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A SURVEY OF CENOZOIC MAMMAL BARAMINS

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ABSTRACT

To expand the sample of statistical baraminology studies, we identified 80 datasets sampled from 29 mammalian orders, from which we performed 82 separate analyses. We analyzed each dataset with standard statistical baraminology techniques: baraminic distance correlation (BDC) and multidimensional scaling (MDS). We evaluated the BDC and MDS results from each character set for potential continuity and discontinuity. We found evidence of holobaramins in 57 of the 82 analyses (69.5%). Of the remaining character sets, three showed evidence of monobaramins and 22 (26.8%) were inconclusive. These results are consistent with previous efforts to test the discontinuity hypothesis, which found that a majority of character sets showed evidence of holobaramins. Tentative holobaramins represent 57 taxonomic groups, many of which have not been previously analyzed by statistical baraminology. Together with previously identified holobaramins, this study increases the number of putative mammal holobaramins to 64.

KEY WORDS

baramin, created kind, baraminology, mammal, Cenozoic, post-Flood, baraminic distance correlation (BDC), multidimensional scaling (MDS)

INTRODUCTION

The concept of the created kind—a group of modern species connected by common ancestry to a population created *ex nihilo* by God—has a long history in creationist thought. Initially suggested before the publication of Darwin's *Origin* by scholars such as Linnaeus and Herbert, creationists throughout the twentieth century have endorsed the created kind as a contrast to species fixity (Wood 2008a). More recently, creationists have produced several large-scale estimates of mammalian created kinds (e.g., Wise 2009; Wood 2011; Lightner 2012).

Although there has been much written about created kinds, statistical baraminology is a comparatively underserved field of research. Few created kinds have been established unequivocally using statistical studies, and even for those taxa which have been investigated, the datasets suitable for baraminological calculations and the reliability of the methods used have rarely been confirmed (Wood 2016a). Some outstanding questions include:

- Are holobaramins marked by discontinuity with all other organisms, defined as “significant, holistic difference” (Wood et al. 2003)?
- Is discontinuity detectable by current statistical baraminology methods?
- What sorts of character or taxon samples are most suitable for statistical baraminology methods?

Previously, Wood (2009) evaluated a collection of 73 baraminology analyses to answer some of these questions. Assuming that discontinuity would distinguish baramins approximately at the taxonomic level of family, he found 62% of analyses revealed discontinuity as predicted, although the fraction was not statistically significant.

This project focused on the mammals of the Cenozoic, more than doubling the number of previous statistical baraminology studies.

Mammals are particularly interesting to creationists because of the numerous stratomorphic series like the famous horse series and because of their affinity to humans. The primary purpose of this work is to provide a database of statistical baraminology studies for future meta-analyses to shed light on such pressing topics as post-Flood diversification, microevolution and speciation, biological imperfection, endemism, and biogeography. We here focus on the question of whether discontinuity distinguishes mammalian families from other similar families and secondarily on the question of what combination of characters and taxa yield successful baraminology studies. These new analyses represent a large-scale first approximation of baramins from at least 79 mammal families.

MATERIALS AND METHODS

We collected character sets from the published literature identified using Google Scholar searches. We selected matrices that primarily focused on a single family or collection of families. We excluded redundant datasets on the same taxa and from the same author(s), and we used the most recent version available. In deference to the baraminological emphasis on holistic similarity and difference, we preferred datasets that had more taxa, characters, or both. In three cases, the kangaroos (Macropodidae), the new world monkey family Cebidae, and the South American notoungulates Interatheriidae, two separate datasets were included for each, since in each case the character and taxon selections were sufficiently different to consider them separately. We also excluded datasets of cetaceans and hominins, in anticipation of forthcoming and more detailed analyses than this simple overview. Six character sets with members from multiple families (Ornithorhynchidae, Felidae, Plesiadapidae, Picrodontidae, Lemuridae, and Omomyidae) were subdivided to include only taxa from an ingroup family and the nearest outgroup clade, and these taxonomic subsets were examined individually rather than *en masse*. For consistency, all selected

datasets were organized and classified by family and order as listed in McKenna and Bell (1997), even when such classification may be out of date.

Prior to any analyses, all hypothetical ancestral taxa were removed from the datasets. Baraminic distance correlation (BDC) and multidimensional scaling (MDS) were calculated using BDISTMDS (<http://coresci.org/bdist.html>). For each baraminology analysis, we recorded the same diagnostic statistics as in Wood's (2008b) previous survey of statistical baraminology studies. Analyses were classified as holobaramin, monobaramin, or inconclusive based on evaluation of the statistical baraminology results. Ideally, a holobaramin can be revealed by BDC showing significant, positive BDC between all members of the holobaramin and significant, negative BDC between the holobaramin members and all outgroup taxa. In practice, this pattern rarely obtains. Alternative patterns include significant, positive BDC between ingroup and outgroup and significant, negative BDC between taxa that also share significant, positive BDC with the same third taxon. MDS aids in the interpretation of the BDC results by confirming clusters of taxa reflected in the BDC results or by revealing complex geometric shapes of taxa (e.g., lines, arcs, or tetrahedra) that cause uninterpretable BDC patterns. Analyses that did not reveal clear clusters of taxa were classified as inconclusive.

RESULTS

We collected 80 character sets from the published literature, from which we performed 82 separate baraminology analyses. The types of characters were most frequently craniodental (43.9%), followed by craniodental with postcranial (28.0%). Only 11% included more types of characters (such as craniodental, postcranial, and soft tissue), and 9.8% consisted only of dental characters. Given the emphasis on holistic analysis in baraminology (Wood et al. 2003), analyses that use more types of characters should be preferred to analyses that use fewer. Especially tentative are any conclusions based solely on dental characters.

We discovered evidence of discontinuity around a holobaramin in 57 (69.5%) of the 82 analyses. Three additional analyses (3.7%) revealed evidence of continuity without discontinuity (monobaramins), and the remaining 22 analyses (26.8%) were classified as inconclusive. A complete account of all results with references is included in the appendix.

Of the 57 analyses that revealed a putative holobaramin, 25 were based on craniodental characters (43.9%), 15 on craniodental and postcranial characters (26.3%), and 6 on dental characters only (10.5%) (see Table 1). These fractions are not substantially different from the overall frequency of character types in the full set of 82 datasets. Of the 22 analyses that were inconclusive, 11 were based on craniodental characters (50.0%), 7 on craniodental and postcranial characters (31.8%), and only two on dental characters only (9.1%). Again, these fractions are not substantially different from the full set of datasets.

The number of taxa and characters used in the analyses that revealed holobaramins were not significantly different from those that resulted in inconclusive results (Table 2). For the MDS analyses, the stress at three dimensions and the dimensions of minimum stress (k_{\min}) also did not differ significantly between analyses that

identified holobaramins when compared to those that did not. In the BDC analyses, the median bootstrap value and the fraction of correlations with bootstrap values >90% (F_{90}) were also not significantly different in BDC analyses that revealed holobaramins than in those that did not.

Based on the successful analyses, we identified 59 putative holobaramins, 49 of which corresponded to families, seven to subfamilies (or portions of families), two to superfamilies (or multiple families), and one (Sirenia) to an infraorder (Table 3). Our results reveal the first five Australian marsupial holobaramins identified by statistical baraminology: the extinct Palorchestidae, the rat kangaroos Hysiprymmodontidae, the kangaroos Macropodidae, ringtail possums Pseudocheirinae, and the koalas Phascolarctidae. Other notable holobaramins first identified here include the Ornithorhynchidae (platypus), the Mephitidae (skunks), and the Sirenia (manatees).

Ten of these groups had already been studied in baraminology analyses (Table 3). Previously identified holobaramins confirmed by this study include Didelphidae (possums), Felidae (cats), Erinaceinae (hedgehogs), and Hippopotamidae (hippos). The Talpidae (moles) were previously inconclusive in Wood's (2009) study, but here they are identified as a holobaramin. The Ursidae (bears), Camelidae (camels), and Brontotheriidae were both previously identified as monobaramins (Wood 2016a), but here are recognized as holobaramins. Our present results also recognize two holobaramins within the Cingulata: Dasypodidae and Glyptodontinae. Previously, Wood (2008b) tentatively proposed that the entire Cingulata might have been a holobaramin, but these results suggest otherwise.

DISCUSSION

Our results substantially expand the available database of baraminology studies. Wood (2016a) previously identified 70 holobaramins identified in studies of 153 taxonomic groups. Our results expand the total number of taxonomic groups studied to 215 with 125 putative holobaramins identified. Within the non-hominin, non-cetacean mammals, we expand the putative holobaramins from 10 to 64. This remains far fewer than the total number of proposed mammal baramins. Assuming that the taxonomic family is roughly equivalent to the holobaramin, McKenna and Bell (1997) list 347 mammal families, of which 64 putative holobaramins is only 18.4%.

Our results agree closely with the mammal ark kind estimations of Lightner (2012). Lightner lists only extant mammals, of which 30 of our holobaramins overlap taxonomically with 30 of her ark kinds. In 28 cases, our holobaramin matches roughly equivalently with her ark kinds, but in three cases, they differ. Our analysis of lemurs identified Indriidae and Palaeopropithecidae as a holobaramin, but Lightner recognized only Indriidae. This could be merely a result of Lightner's exclusion of extinct taxa. In the other two cases, we found holobaramins that appeared to be taxonomically more restricted than Lightner's ark kinds. First, our study of the pigs, Suidae, identified the family as the holobaramin with putative discontinuity observed between the suids and the peccaries (Tayassuidae). In contrast, Lightner included both pigs and peccaries in a single ark kind. Lightner also included

Table 1. Character types in the datasets used in this study. “Holobaramin datasets” refer to datasets in which a putative holobaramin was identified, and “Inconclusive datasets” are datasets which produced inconclusive BDC and MDS results.

| Character type | Frequency in all datasets | Percentage in all datasets | Frequency in holobaramin datasets | Percentage in holobaramin datasets | Frequency in inconclusive datasets | Percentage in inconclusive datasets |
|----------------------------|---------------------------|----------------------------|-----------------------------------|------------------------------------|------------------------------------|-------------------------------------|
| Craniodental | 36 | 43.9% | 25 | 43.9% | 11 | 50.0% |
| Craniodental + postcranial | 23 | 28.0% | 15 | 26.3% | 7 | 31.8% |
| Dental | 8 | 9.8% | 6 | 10.5% | 2 | 9.1% |
| >3 types | 9 | 11.0% | 6 | 10.5% | 1 | 4.5% |
| Cranial + postcranial | 4 | 4.9% | 4 | 7.0% | 0 | 0.0% |
| Dental + postcranial | 1 | 1.2% | 0 | 0.0% | 1 | 4.5% |
| Cranial | 1 | 1.2% | 1 | 1.8% | 0 | 0.0% |

Table 2. Comparison of results between datasets for which a holobaramin was identified and datasets which produced inconclusive results. Welch’s unequal variance *t*-test was used to compare the holobaramin and inconclusive datasets.

| | Holobaramin dataset mean | Inconclusive dataset mean | Welch’s <i>t</i> | P-value |
|------------------------|--------------------------|---------------------------|------------------|---------|
| Taxa in analysis | 21.9 | 26.6 | -1.08 | 0.2915 |
| Characters in analysis | 71.4 | 66.1 | 0.38 | 0.7029 |
| 3D stress | 0.16 | 0.17 | -0.45 | 0.6528 |
| k_{\min} | 6.2 | 6.0 | 0.19 | 0.8473 |
| Median bootstrap | 86.1 | 82.0 | 1.80 | 0.0800 |
| F_{90} | 0.44 | 0.34 | 1.98 | 0.0537 |

three genera (*Cyclopes*, *Myrmecophaga*, and *Tamandua*) from two families in the anteater kind Vermilingua, but our results support only *Myrmecophaga* and *Tamandua* in holobaramin Myrmecophaginae. These results will need further evaluation in future studies to clarify these differences.

Wise’s (2009) estimation of mammalian ark kinds differs substantially from all others in the creationist literature. His estimates are based on the Post-Flood Continuity Criterion (PFCC), which in turn is based on the proposal that an ark kind should leave a continuous fossil record from the Flood to the present. His minimal estimate places a large number of species into as few as 97 ark kinds, and his maximal estimate places fewer species into as many as 234 ark kinds. Not surprisingly, our putative holobaramins do not match his estimated ark kinds (Table 3). Instead, we find tentative evidence of discontinuity within 33 of his minimal ark kinds and 18 of his maximal ark kinds. Sixteen of our putative holobaramins match closely sixteen of Wise’s maximal ark kinds. Resolving the discrepancy between Wise’s results and ours will require assessments of the completeness of the post-Flood fossil record as well as further evaluations of our baraminology results.

Our results should also help creationists evaluate baraminology methods, which have recently come under scrutiny (Wood 2016b; O’Micks 2016; Wood 2017; O’Micks 2017). Previously, Wood’s (2009) survey of 73 statistical baraminology studies examined taxon samples that should have revealed discontinuity as well as those that should not (such as within a single genus). Wood found that 61.6% of the studies were “successful” in either revealing or

not revealing predicted discontinuity. Our survey differs in that we focused exclusively on datasets that should have revealed discontinuity around a taxonomic family, but we found 69.5% of studies revealed evidence of the expected discontinuity. This is an improvement from the previous survey, and the question remains: Why do some datasets reveal evidence of discontinuity while others do not?

We attempted to determine what sort of feature of the datasets might correlate or predict successful discontinuity detection by examining the number of taxa and characters, the stress and k_{\min} of the MDS, and the median bootstrap value and F_{90} of the BDC results, but we found no significant correlations, as in previous studies. Future studies will definitely want to examine these results more closely to determine the type of dataset best suited to these studies.

Future studies should more closely evaluate individual taxonomic groups named herein, especially those where discrepancies between different studies are noted. The current best practice in statistical baraminology is to use multiple holistic character sets compiled from different sources to evaluate the baraminology of a single group, as in hominin baraminology (e.g., see Wood 2016c; 2017). Some of our studies add to baraminology studies of previous groups, while most provide only a first approximation of a group’s baraminology. Nevertheless, the present results provide an important advance in baraminology work and in the study of mammalian created kinds.

Table 3. List of putative holobaramins identified in this study and comparable Ark kinds recognized by Lightner (2012) and Wise (2009). Also listed are previous baraminology studies recorded in Wood (2016a). HB = holobaramin; MB = monobaramin.

| Order | Holobaramin | Rank | Previous Baraminology | Lightner's Ark Kinds | Wise's Minimal Ark Kinds | Wise's Maximal Ark Kinds |
|-----------------|---------------------------------|-------------|-----------------------|----------------------|---|--------------------------|
| Platyptoda | Ornithorhynchidae | Family | | Ornithorhynchidae | Prototheria | Ornithorhynchidae |
| Diprotodontia | Palorchestidae | Family | | | Australidelphia | Australidelphia |
| Diprotodontia | Thylacoleonidae | Family | | | Australidelphia | Australidelphia |
| Diprotodontia | Hypsiprymmodontidae | Family | | Hypsiprymmodontidae | Australidelphia | Australidelphia |
| Diprotodontia | Macropodidae | Family | | Macropodidae | Australidelphia | Australidelphia |
| Diprotodontia | Pseudocheirinae | Subfamily | | Pseudocheiridae | Australidelphia | Australidelphia |
| Diprotodontia | Phascolarctidae | Family | | Phascolarctidae | Australidelphia | Australidelphia |
| Didelphimorphia | Didelphidae | Family | HB Didelphidae | Didelphidae | Didelphimorphia | Didelphidae |
| Sparassodonta | Hathliacynidae | Family | | | Sparassodonta | Sparassodonta |
| Cingulata | Dasypodidae | Family | HB Cingulata | Dasypodidae | Xenarthra | Dasypodidae |
| Cingulata | Glyptodontinae | Subfamily | HB Cingulata | | Xenarthra | |
| Pilosa | Myrmecophaginae | Subfamily | | Vermilingua | Xenarthra | Xenarthra |
| Leptictida | Pseudorhynchocyonidae | Family | | | Leptictida | Leptictidae |
| Rodentia | Castoridae | Family | | Castoridae | Castorimorpha | Castorimorpha |
| Rodentia | Anomaluridae | Family | | Anomaluridae | See Wise (2009) p. 151 for comments on Anomaluromorpha | |
| Rodentia | Caviidae | Family | | Caviidae | See Wise (2009) p. 151 for comments on Hystricognatha | |
| Pholidota | Manidae | Family | | Manidae | Pholidota | Manidae |
| Carnivora | Felidae | Family | HB Felidae | Felidae | Feliformia | Feliformia |
| Carnivora | Felidae | Family | HB Felidae | Felidae | Feliformia | Feliformia |
| Carnivora | Ursidae | Family | MB Ursidae | Ursidae | Caniformia | Caniformia |
| Carnivora | Otariidae | Family | | | Caniformia | Caniformia |
| Carnivora | Odobenidae | Family | | | Caniformia | Caniformia |
| Carnivora | Mustelidae | Family | | Mustelidae | Caniformia | Caniformia |
| Carnivora | Mephitidae | Family | | Mephitidae | Caniformia | Caniformia |
| Chrysochloridea | Chrysochloridae | Family | | Chrysochloridae | See Wise (2009) p. 154 for comments on Chrysochloroidea | |
| Erinaceomorpha | Erinaceinae | Subfamily | HB Erinaceinae | Erinaceinae | Erinaceomorpha | Erinaceidae |
| Erinaceomorpha | Galericinae | Subfamily | | Galericinae | Erinaceomorpha | Erinaceidae |
| Erinaceomorpha | Talpidae | Family | ? Talpidae | Talpidae | Erinaceomorpha | Erinaceomorpha |
| Soricomorpha | Nyctitheriidae | Family | | | Soricomorpha | Nyctitheriidae |
| Soricomorpha | Soricidae | Family | | Soricidae | Soricomorpha | Soriocoidea |
| Chiroptera | Rhinolophidae | Family | | Rhinolophidae | Chiroptera | Yinochiroptera |
| Primates | Picrodontidae | Family | | | Picrodontidae | Picrodontidae |
| Primates | Plesiadapidae | Family | | | Plesiadapidae | Plesiadapidae |
| Primates | Lemuridae | Family | | Lemuridae | Strepsirrhini | Lemuroidea |
| Primates | Indriidae + Palaeopropithecidae | Superfamily | | Indriidae | Strepsirrhini | Lemuroidea |
| Primates | Lepilemuridae | Family | | Lepilemuridae | Strepsirrhini | Lemuroidea |
| Primates | Cheirogaleidae | Family | | Cheirogaleidae | Strepsirrhini | Lemuroidea |
| Primates | Carpolestidae | Family | | | Carpolestoidea | Carpolestidae |
| Primates | Omomyidae | Family | | | Tarsioidea | Omomyidae |
| Primates | Callitrichidae | Family | | | Anthropoidea | Anthropoidea |
| Condylarthra | Didolodontidae | Family | | | Didolodontidae | Didolodontidae |
| Artiodactyla | Suidae | Family | | Suoidea | See Wise (2009) p. 157 for comments on Suoidea | |
| Artiodactyla | Hippopotamidae | Family | HB Hippopotamidae | | See Wise (2009) p. 157 for comments on Suoidea | |
| Artiodactyla | Camelidae | Family | MB Camelidae | Camelidae | Cameloidea | Camelidae |
| Notoungulata | Leontiniidae | Family | | | Toxodontia | Toxodontia |
| Notoungulata | Toxodontidae | Family | | | Toxodontia | Toxodontia |
| Notoungulata | Interatheriidae | Family | | | Tyotheria | Interatheriidae |
| Notoungulata | Hegetotheriidae | Family | | | Hegetotheria | Hegetotheria |
| Astrapotheria | Astrapotheriidae | Family | | | Astrapotheria | Astrapotheriidae |
| Xenoungulata | Carodniidae | Family | | | Xenoungulata | Carodniidae |
| Perissodactyla | Palaeotheriinae s. l. | Subfamily | | | Hippomorpha | Palaeotheriidae |
| Perissodactyla | Brontotheriidae | Family | MB Brontotheriidae | | Brontotherioidea | Brontotheriidae |
| Perissodactyla | Chalicotherioidea | Superfamily | | | Chalicotherioidea | Chalicotherioidea |
| Perissodactyla | Lophodontinae | Subfamily | | | Tapiroidea | Lophodontidae |
| Uranotheria | Sirenia | Infraorder | | | Sirenia | Sirenia |
| Uranotheria | Desmostylidae | Family | | | Behemota | Behemota |
| Hyracoidea | Procaviidae | Family | | Procaviidae | Hyracoidea | Hyracoidea |
| Proboscidea | Gomphotheriidae | Family | | | Behemota | Behemota |
| Proboscidea | Elephantidae | Family | | Elephantidae | Behemota | Behemota |

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REFERENCES

- Lightner, J.K. 2012. Mammalian ark kinds. *Answers Research Journal* 5:151-204.
- McKenna, M.C. and S.K. Bell. 1997. *Classification of Mammals above the Species Level*. New York: Columbia University Press.
- O'Micks, J. 2016. Reply to "Taxon sample in hominin baraminology: A response to O'Micks." *Answers Research Journal* 9:373-375.
- O'Micks, J. 2017. Rebuttal to "Reply to O'Micks concerning the geology and taphonomy of the *Homo naledi* site" and "Identifying humans in the fossil record: A further response to O'Micks." *Answers Research Journal* 10:63-70.
- Wise, K.P. 2009. Mammal kinds: how many were on the ark? *CORE Issues in Creation* 5:129-161.
- Wood, T.C. 2008a. Species variability and creationism. *Origins* 62:6-25.
- Wood, T.C. 2008b. Animal and plant baramins. *CORE Issues in Creation* 3:1-258.
- Wood, T.C. 2009. Natura facit saltum: The case for discontinuity. *CORE Issues in Creation* 5:113-127.
- Wood, T.C. 2011. Terrestrial mammal families and creationist perspectives on speciation. *Journal of Creation Theology and Science Series B: Life Sciences* 1:2-5.
- Wood, T.C. 2016a. A list and bibliography of identified baramins. *Journal of Creation Theology and Science Series B: Life Sciences* 6:91-101.
- Wood, T.C. 2016b. Taxon sample size in hominin baraminology: A response to O'Micks. *Answers Research Journal* 9:369-372.
- Wood, T.C. 2016c. An evaluation of *Homo naledi* and "early" *Homo* from a young-age creationist perspective. *Journal of Creation Theology and Science Series B: Life Sciences* 6:14-30.
- Wood, T.C. 2017. Identifying humans in the fossil record: A further response to O'Micks. *Answers Research Journal* 10:57-62.
- Wood, T.C., K.P. Wise, R. Sanders, and N. Doran. 2003. A refined baramin concept. *Occasional Papers of the BSG* 3:1-14.

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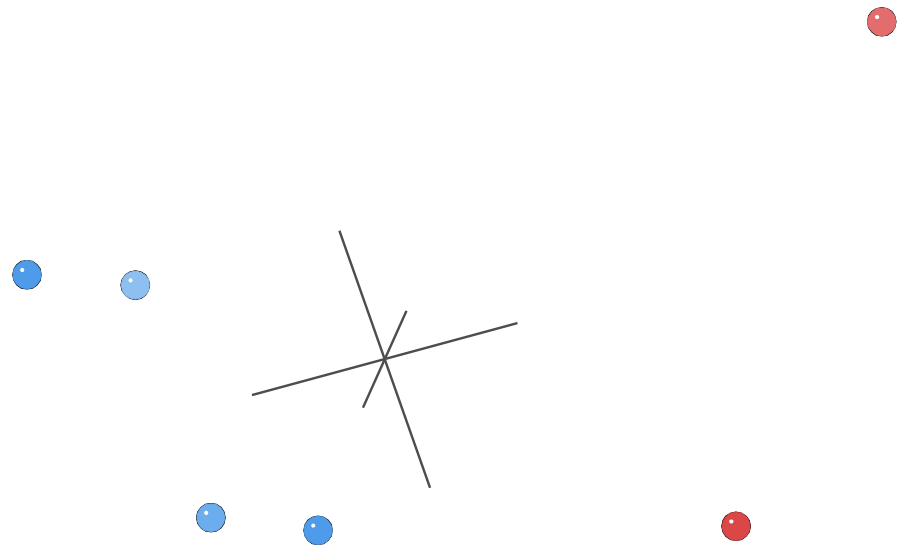
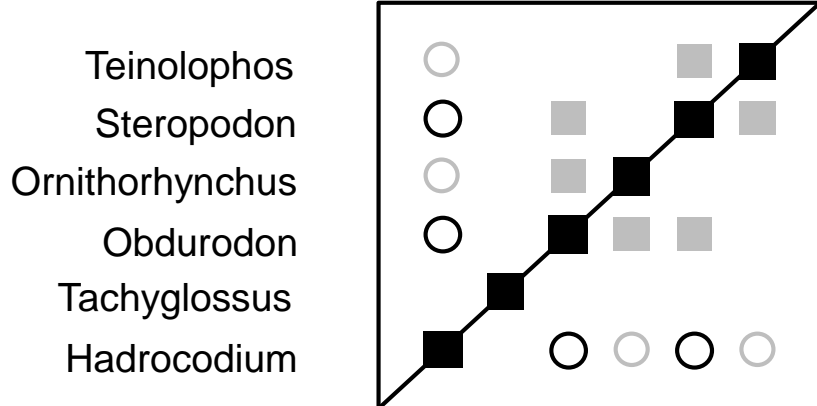
Appendix. BDC and MDS analyses of Cenozoic Mammalia

This appendix contains the results of all 82 BDC and MDS analyses. All baraminology analyses were conducted in BDISTMDS (coresci.org/bdistmds.html). For all Multidimensional Scaling (MDS) illustrations, outgroup taxa are shown in red and ingroup taxa in blue, unless otherwise indicated. Statistics for Baraminic Distance Correlation (BDC) and MDS are summarized in a table, as in Wood's (2008) "Animal and Plant Baramins." As in Wood (2008), F_{90} is the fraction of taxon pairs having bootstrap values >90%, and k_{min} is the MDS dimensionality at which the minimum stress is observed. A conclusion is also included in the table (HB = holobaramin, HB? = provisional holobaramin, MB = monobaramin, and Inc = inconclusive). The conclusion is briefly explained for each data set. The taxa are classified and ordered as in McKenna and Bell (1997). An index of the mammal taxa is provided below, page numbers are indicated at the bottom right of each page.

| | | | | | |
|-------------------|----|---------------------|----|-----------------------|----|
| Anomaluridae | 23 | Hegetotheriidae | 71 | Otariidae | 34 |
| Anthracotheriidae | 62 | Hippopotamidae | 61 | Palaeoryctidae | 27 |
| Aplodontidae | 20 | Hyaenodontidae | 29 | Palaeotheriidae | 74 |
| Astrapotheriidae | 72 | Hyopsodontidae | 58 | Palorchestidae | 4 |
| Barbourofelinae | 32 | Hypsiprymnodontidae | 6 | Peramelidae | 3 |
| Brontotheriidae | 75 | Interatheriidae | 69 | Phascolarctidae | 10 |
| Caenolestidae | 12 | Interatheriidae | 70 | Phyllostomidae | 46 |
| Camelidae | 63 | Lemuridae | 49 | Picrodontidae | 47 |
| Carodniidae | 73 | Leontiniidae | 67 | Plesiadapidae | 48 |
| Carpolestidae | 52 | Lepilemuridae | 50 | Procaviidae | 81 |
| Castoridae | 21 | Leporidae | 19 | Procyonidae | 38 |
| Caviidae | 24 | Lophiodontidae | 78 | Pseudocheirinae | 9 |
| Cebidae | 54 | Loridae | 51 | Pseudorhynchocyonidae | 17 |
| Cebidae | 55 | Louisinidae | 57 | Rhinocerotidae | 77 |
| Cervidae | 65 | Macropodidae | 7 | Rhinolophidae | 44 |
| Chalicotheriidae | 76 | Macropodidae | 8 | Sirenia | 79 |
| Chrysochloridae | 39 | Manidae | 28 | Soricidae | 43 |
| Cricetidae | 22 | Mephitidae | 37 | Suidae | 60 |
| Dasypodidae | 14 | Mormoopidae | 45 | Talpidae | 41 |
| Desmostylidae | 80 | Moschidae | 64 | Thylacoleonidae | 5 |
| Didelphidae | 11 | Mustelidae | 36 | Toxodontidae | 68 |
| Didolodontidae | 59 | Myrmecophagidae | 16 | Ursidae | 33 |
| Echimyidae | 26 | Notohippidae | 66 | | |
| Elephantidae | 83 | Nyctitheriidae | 42 | | |
| Erinaceidae | 40 | Ochotonidae | 18 | | |
| Felidae | 30 | Octodontidae | 25 | | |
| Felidae | 31 | Odobenidae | 35 | | |
| Glyptodontidae | 15 | Omomyidae | 53 | | |
| Gomphotheriidae | 82 | Ornithorhynchidae | 2 | | |
| Hathliacynidae | 13 | Orycteropodidae | 56 | | |

Rowe, T., T.H. Rich, P. Vickers-Rich, M. Springer, and M.O. Woodburne. 2008. The oldest platypus and its bearing on divergence timing of the platypus and echidna clades. *Proceedings of the National Academy of Sciences USA* 105:1238-1242.

Characters: Craniodental and postcranial



Order Platypoda
Family Ornithorhynchidae

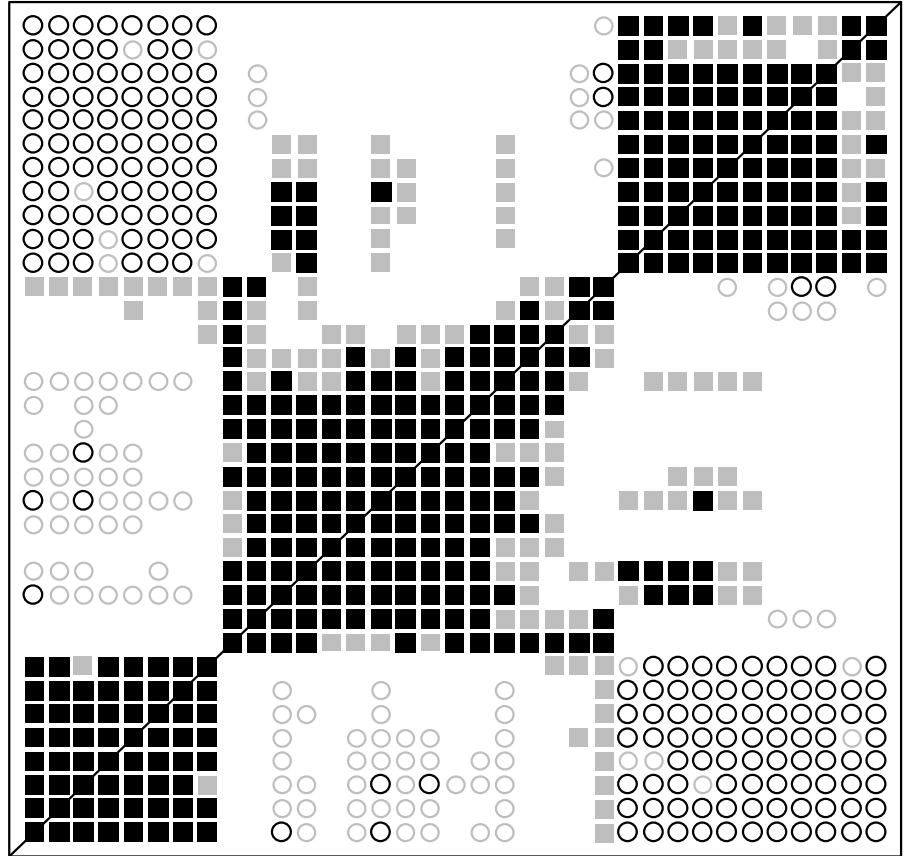
| | |
|----------------------------------|------|
| Published taxa | 6 |
| Published characters | 390 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 6 |
| Characters used for calculations | 50 |
| Median bootstrap value | 91 |
| F ₉₀ | 0.53 |
| Stress of 3D MDS | 0.15 |
| k _{min} | 2 |
| Conclusion | HB? |

Notes: Character set has been reduced to just Monotremata from the published dataset. BDC results have poor bootstrap values, and MDS reveals a disperse cloud of taxa. Nevertheless, ornithorhynchids share significant, positive BDC, and are noticeably separated from outgroup taxa in MDS. We may provisionally accept holobaramin Ornithorhynchidae.

Travouillon, K.J., S.J. Hand, M. Archer, and K.H. Black. 2014. Earliest modern bandicoot and bilby (Marsupialia, Peramelidae, and Thylacomyidae) from the Miocene of the Riversleigh World Heritage Area, Northwestern Queensland, Australia. *Journal of Vertebrate Paleontology* 34:375-382.

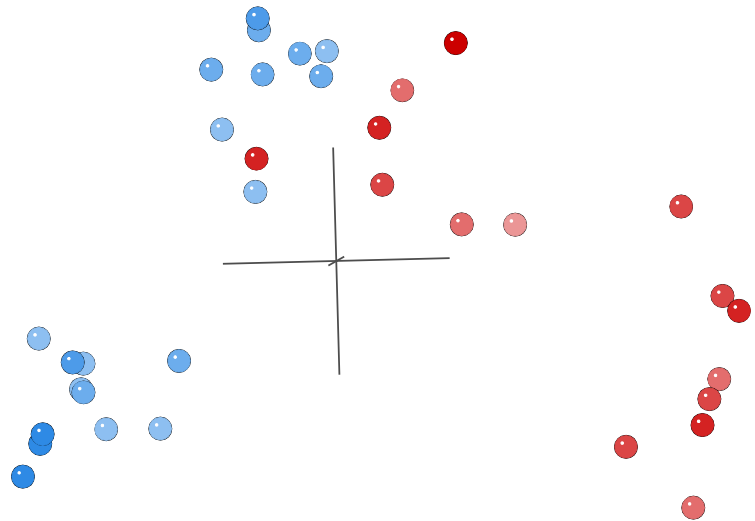
Characters: Craniodental

- Macrotus leucura*
- Macrotus lagotis*
- Isoodon macrourus*
- Isoodon obesulus*
- Isoodon auratus*
- Perameles gunnii*
- Perameles nasuta*
- Perameles sobbei*
- Perameles eremiana*
- Perameles bougainville*
- Chaeropus ecaudatus*
- Yarala kida*
- Yarala burchfieldi*
- Galadi grandis*
- Galadi speciosus*
- Galadi amplus*
- Peroryctes broadbenti*
- Peroryctes raffrayana*
- Rhynchomeles prattorum*
- Echymipera rufescens*
- Echymipera kalubu*
- Echymipera davidi*
- Echymipera clara*
- Microperoryctes papuensis*
- Microperoryctes longicauda*
- cf. Peroryctes tedfordi*
- Bulungu palara*
- Djarthia murgonensis*
- Mutpuracinus archibaldi*
- Barinya wangala*
- Dasyurus hallucatus*
- Sminthopsis macroura*
- Dasyuroides byrnei*
- Phascogale tapoatafa*
- Antechinus stuartii*



Order Peramelia
Family Peramelidae

| | |
|----------------------------------|------|
| Published taxa | 42 |
| Published characters | 156 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 35 |
| Characters used for calculations | 140 |
| Median bootstrap value | 91 |
| F ₉₀ | 0.52 |
| Stress of 3D MDS | 0.26 |
| k _{min} | 12 |
| Conclusion | Inc |

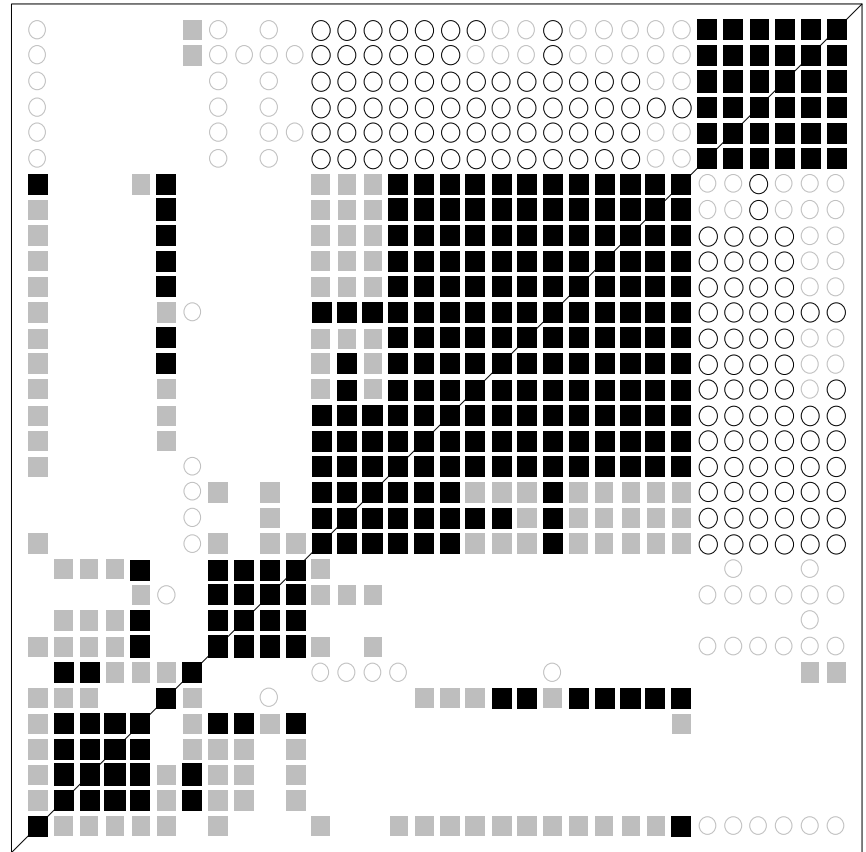


Notes: Three groups appear in both BDC and MDS: one with only peramelids, one with only outgroup taxa, and one with both. BDC and MDS do not support a clear holobaramin, even provisionally.

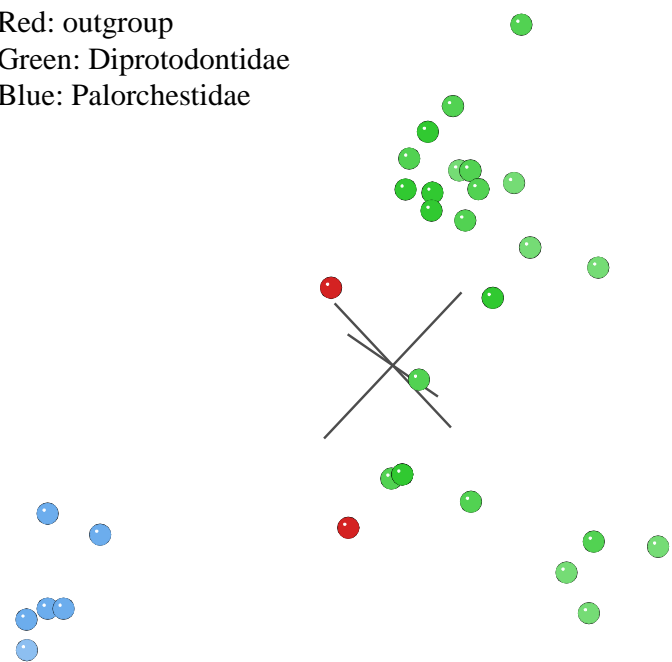
Black, K. 2008. Diversity, Phylogeny and biostratigraphy of Diprotodontoids (Marsupialia: Diprotodontidae, Palorchestidae) from the Riversleigh World Heritage Area. Ph.D. Dissertation. Sydney, Australia: University of New South Wales.

Characters: Craniodental

- Pr. ponticulus*
- Pr. novaculacephalus*
- Palorchestes painei*
- Palorchestes pickeringi*
- Palorchestes parvus*
- Palorchestes azael*
- Silvabestius michaelbirti*
- Silvabestius johnnilandi*
- Nimbadoron lavarackorum*
- Plaisiodon centralis*
- Neohelos stirtoni*
- Neohelos gingari*
- Neohelos tirarensis*
- Neohelos gadiyanga*
- Kolopsis yperus*
- Kolopsis torus*
- Kolopsoides cultridens*
- Kolopsis rotundus*
- Maokopia ronaldi*
- Zygomaturus trilobus*
- Hulitherium tomasetti*
- Euryzygoma dunense*
- Meniscolophus mawsoni*
- Euowenia grata*
- Diprotodon optatum*
- Vombatus ursinus*
- Namilamadeta albivenator*
- Pyramios alcootensis*
- Pitikantia dailyi*
- Ngapakaldia tedfordi*
- Ngapakaldia bonythoni*
- Alkwertatherium webbi*



Red: outgroup
 Green: Diprotodontidae
 Blue: Palorchestidae

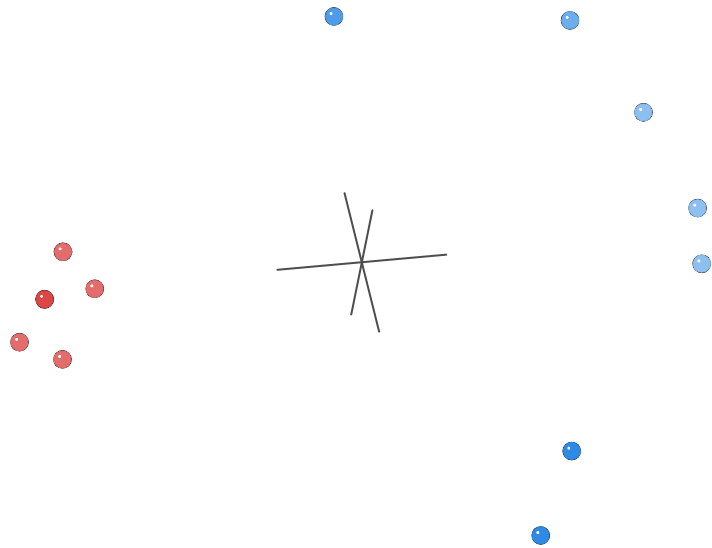
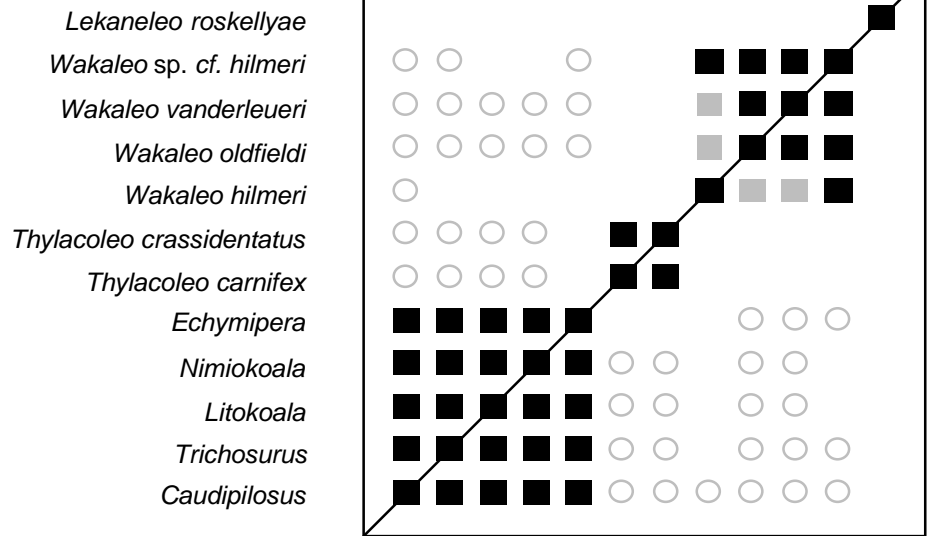


Order Diprotodontia
 Family Palorchestidae

| | |
|----------------------------------|------|
| Published taxa | 35 |
| Published characters | 77 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 32 |
| Characters used for calculations | 57 |
| Median bootstrap value | 79 |
| F ₉₀ | 0.39 |
| Stress of 3D MDS | 0.1 |
| k _{min} | 4 |
| Conclusion | HB |

Notes: This character set is taken from chapter ten of Black's dissertation. BDC appears to support clear separation of palorchestids from outgroup taxa, except for *Vombatus ursinus*, which is correlated positively with two palorchestids (but with low bootstrap values). The MDS definitely supports an inference of discontinuity between Palorchestidae and the outgroup taxa. Palorchestidae is probably a holobaramin.

Gillespie, A.K. 2007. Diversity and systematics of marsupial lions from the Riversleigh World Heritage Area and the evolution of the Thylacoleonidae. Ph.D. Dissertation. Sydney, Australia: University of New South Wales.
 Characters: Craniodental



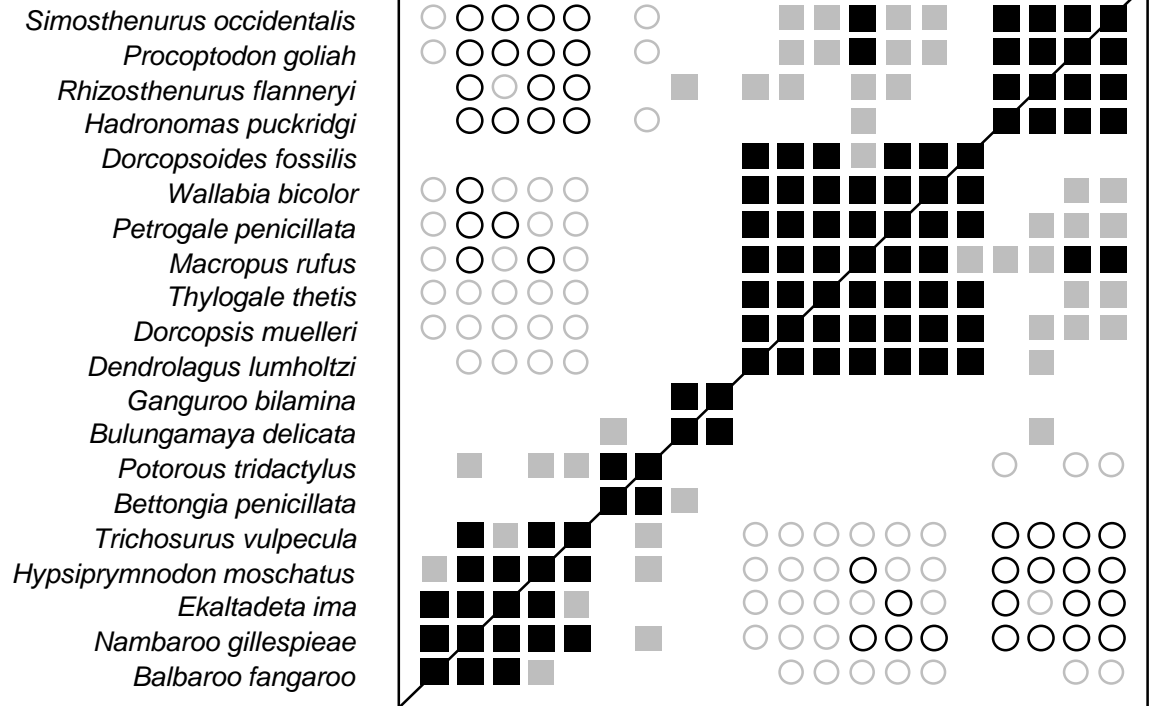
Order Diprotodontia
 Family Thylacoleonidae

| | |
|----------------------------------|------|
| Published taxa | 17 |
| Published characters | 72 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 12 |
| Characters used for calculations | 51 |
| Median bootstrap value | 89 |
| F ₉₀ | 0.47 |
| Stress of 3D MDS | 0.12 |
| k _{min} | 6 |
| Conclusion | HB? |

Notes: This dataset comes from chapter 11 of Gillespie's dissertation. The BDC shows what appear to be four groups of thylacoleonids all negatively correlated with the outgroup taxa but with little evidence of positive correlation with each other. The MDS reveals an arc of thylacoleonid taxa surrounding a tight cluster of outgroup taxa. The curvilinear distribution of thylacoleonid taxa might account for the poor intrafamilial BDC results. Hence we may provisionally accept Thylacoleonidae as a holobaramin.

Bates, H., K.J. Travouillon, B. Cooke, R.M.D. Beck, S.J. Hand, and M. Archer. 2014. Three new Miocene species of musky rat-kangaroos (Hypsiprymnodontidae, Macropodoidea): description, phylogenetics, and paleoecology. *Journal of Vertebrate Paleontology* 34:383-396.

Characters: Craniodental



Bulungamaya delicata ●

Ganguroo bilamina ●

Hadronomas puckridgi ●

Order Diprotodontia
Family Hypsiprymnodontidae

| | |
|----------------------------------|------|
| Published taxa | 33 |
| Published characters | 107 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 20 |
| Characters used for calculations | 93 |
| Median bootstrap value | 84 |
| F ₉₀ | 0.36 |
| Stress of 3D MDS | 0.12 |
| k _{min} | 5 |
| Conclusion | HB |

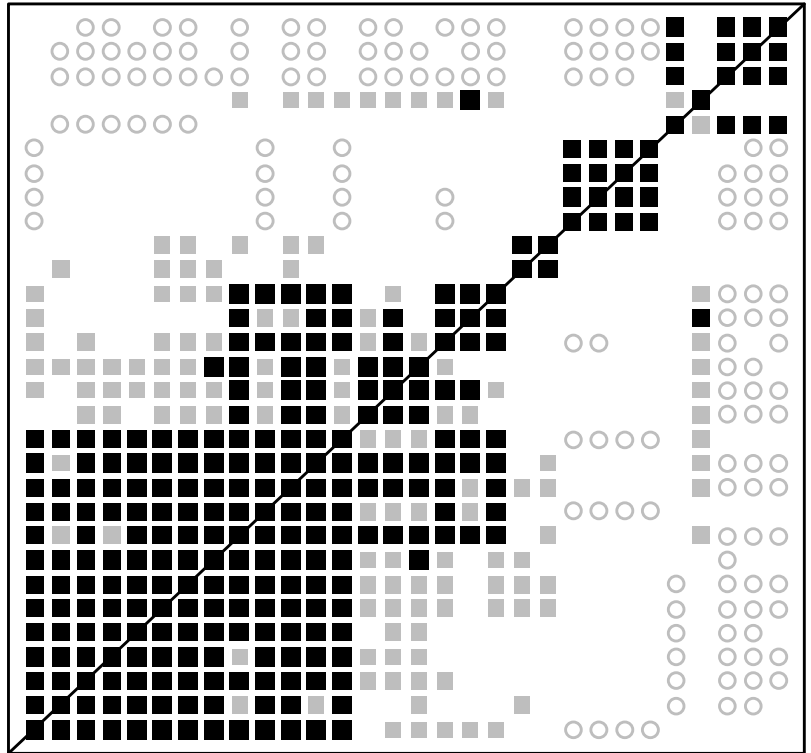
Red: outgroup
Green: Macropodidae
Blue: Hypsiprymnodontidae

Notes: MDS reveals a disperse cluster of ingroup taxa, but BDC supports recognizing a discontinuity between ingroup and outgroup. There is only one taxon pair with significant, positive BDC between the Macropodidae and Hypsiprymnodontidae: *Rhizosthenurus flanneryi* and *Bulungamaya delicata* respectively. Since the bootstrap value for that correlation is only 79% and the MDS reveals that *Bulungamaya* is an outlier from the hypsiprymnodontids and not adjacent to the macropodids, we may consider that correlation spurious. Hypsiprymnodontidae is probably a holobaramin.

Prideaux, G.J. and R.H. Tedford. 2012. *Tjukuru wellsi*, gen. et sp. nov., a lagostrophine kangaroo (Diprotodontia, Macropodidae) from the Pliocene (Tirarian) of northern South Australia. *Journal of Vertebrate Paleontology* 32:717-721.

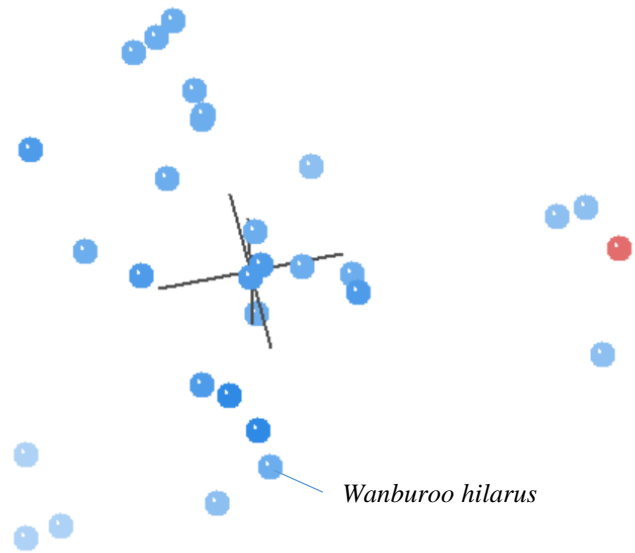
Characters: Craniodental and postcranial

Sthenurus_andersoni
 Simosthenurus_occidentalis
 Procoptodon_goliah
 Wanburoo_hilarus
 Hadronomas_puckridgi
 Ngamaroo_archeri
 Hypsiprymnodon_moschatus
 Potorous_tridactylus
 Bettongia_penicillata
 Troposodon_minor
 Lagostrophus_fasciatus
 Dendrolagus_matschiei
 Dendrolagus_bennettianus
 Bohra_illuminata
 Dorcopsoides_fossilis
 Dorcopsulus_vanheurni
 Dorcopsis_veterum
 Protemnodon_anak
 Setonix_brachyurus
 Thylogale_billardieri
 Prionotemnus_palankarinnicus
 Petrogale_brachyotis
 Wallabia_bicolor
 Lagorchestes_hirsutus
 Lagorchestes_conspicillatus
 Macropus_obustus
 Macropus_fuliginosus
 Macropus_eugenii
 Onychogalea_unguifera
 Baringa_nelsonensis



Order Diprotodontia
 Family Macropodidae

| | |
|----------------------------------|------|
| Published taxa | 36 |
| Published characters | 83 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 30 |
| Characters used for calculations | 80 |
| Median bootstrap value | 81 |
| F ₉₀ | 0.34 |
| Stress of 3D MDS | 0.18 |
| k _{min} | 6 |
| Conclusion | HB? |

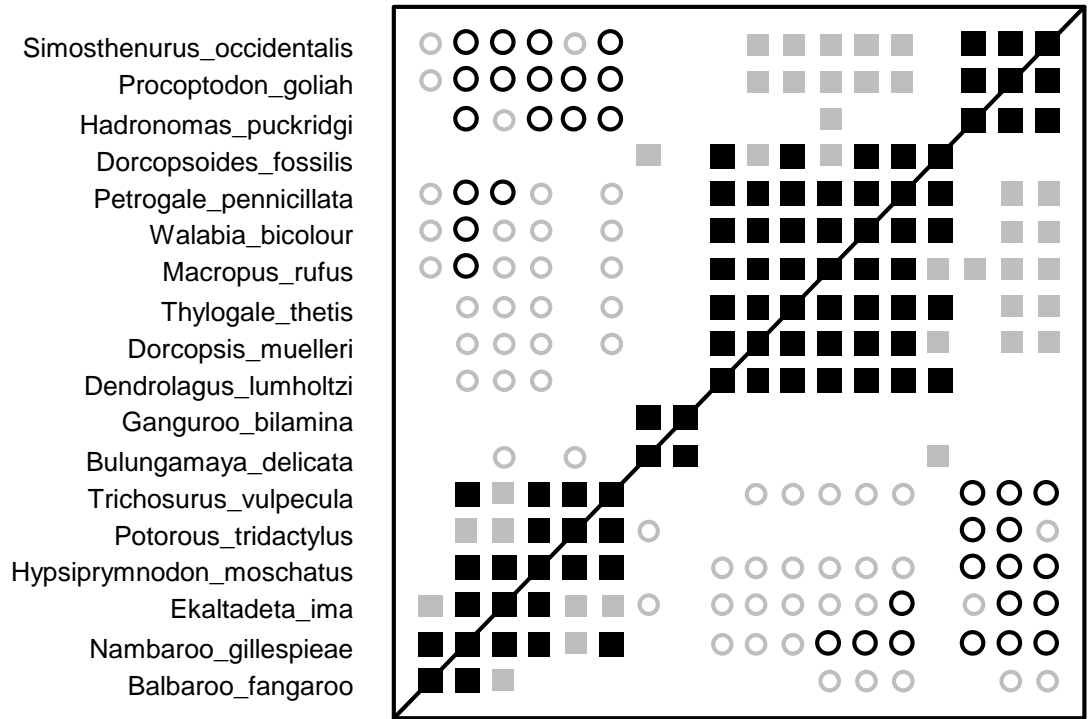


Notes: The outgroup indicated in the MDS results is only the hypsiprymnodontid *Hypsiprymnodon moschatus*. Other non-macropodid taxa include the potoroids *Bettongia* and *Potorous*.

The BDC results indicate a clear separation between macropodids and the three outgroup taxa. Also, BDC shows positive BDC between stem macropodoid *Ngamaroo* and the outgroup taxa and negative BDC between *Ngamaroo* and the ingroup. Macropodoid *Wanburoo* appears to correlate with the outgroup and ingroup taxa, but the MDS results reveal that *Wanburoo* is not actually part of the outgroup cluster. Instead, it appears to be part of the Macropodidae holobaramin.

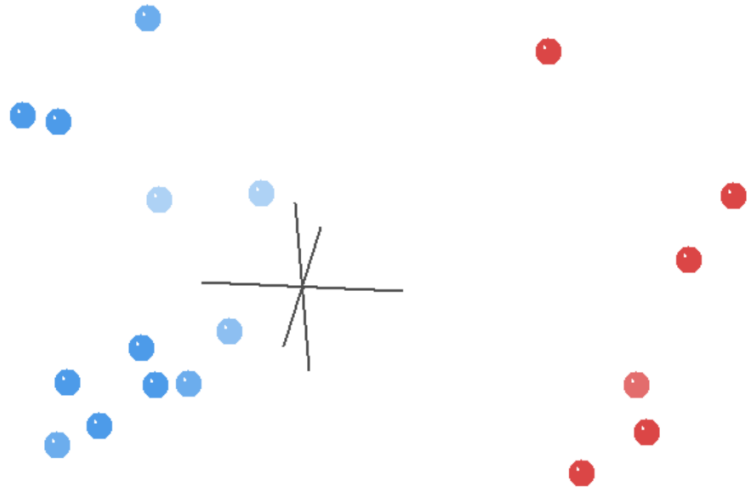
Kear, B.P., B.N. Cooke, M. Archer, and T.F. Flannery. 2007. Implications of a new species of the Oligo-Miocene kangaroo (Marsupialia: Macropodoidea) *Nambaroo*, from the Riversleigh World Heritage Area, Queensland, Australia. *Journal of Paleontology* 81:1147-1167.

Characters: Craniodental and postcranial



Order Diprotodontia
Family Macropodidae

| | |
|----------------------------------|------|
| Published taxa | 25 |
| Published characters | 104 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 18 |
| Characters used for calculations | 85 |
| Median bootstrap value | 83 |
| F ₉₀ | 0.34 |
| Stress of 3D MDS | 0.15 |
| k _{min} | 6 |
| Conclusion | HB |

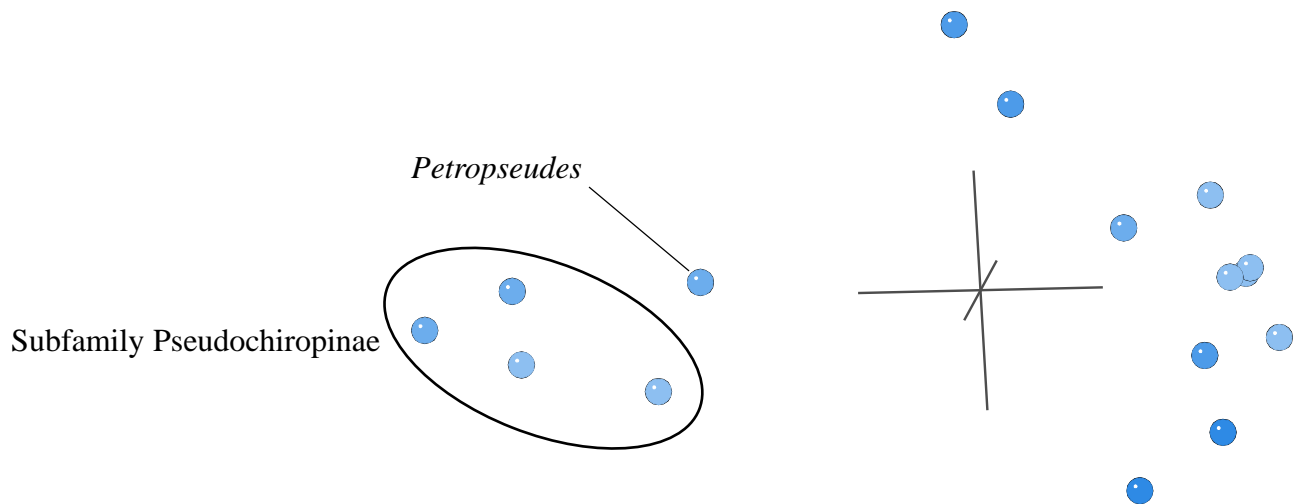
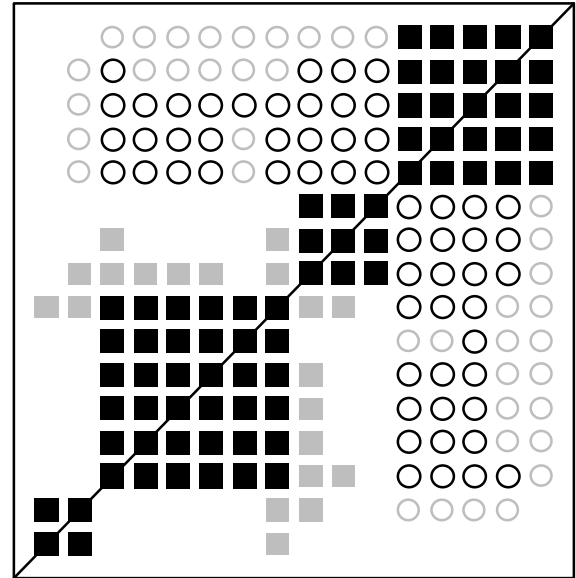


Notes: This second macropodid dataset is included because it contains fewer taxa than the previous Prideaux and Tedford dataset but more characters. In this case, the outgroups and macropodids are definitely separated in both BDC and MDS results, supporting the inference that Macropodidae is a holobaramin.

Springer, M.S. 1993. Phylogeny and rates of character evolution among ringtail possums (Pseudocheiridae, Marsupialia). *Australian Journal of Zoology* 41:273-291.

Characters: Craniodental

Petropseudes dahli
Pseudochirops corinnae
Pseudochirops cupreus
Pseudochirops archeri
Pseudochirops albertisii
Pildra
Parjara
Marlu
Pseudocheirus peregrinus
Pseudochirulus herbertensis
Pseudochirulus mayeri
Pseudochirulus forbesi
Pseudochirulus caroli
Pseudochirulus canescens
Petauroides volans
Hemibelideus lemuroides



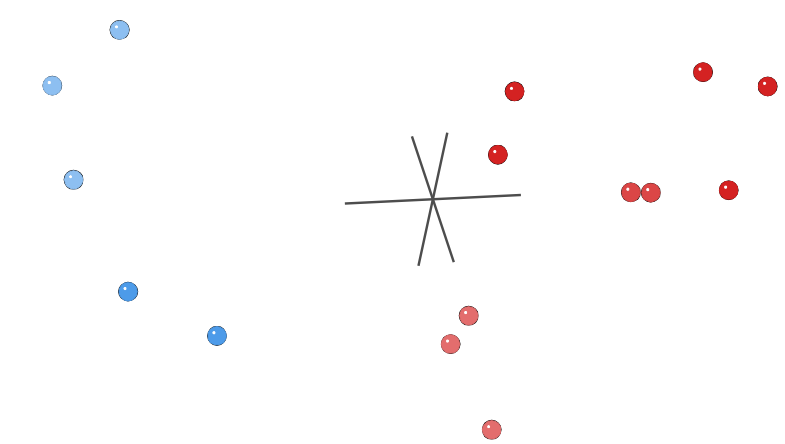
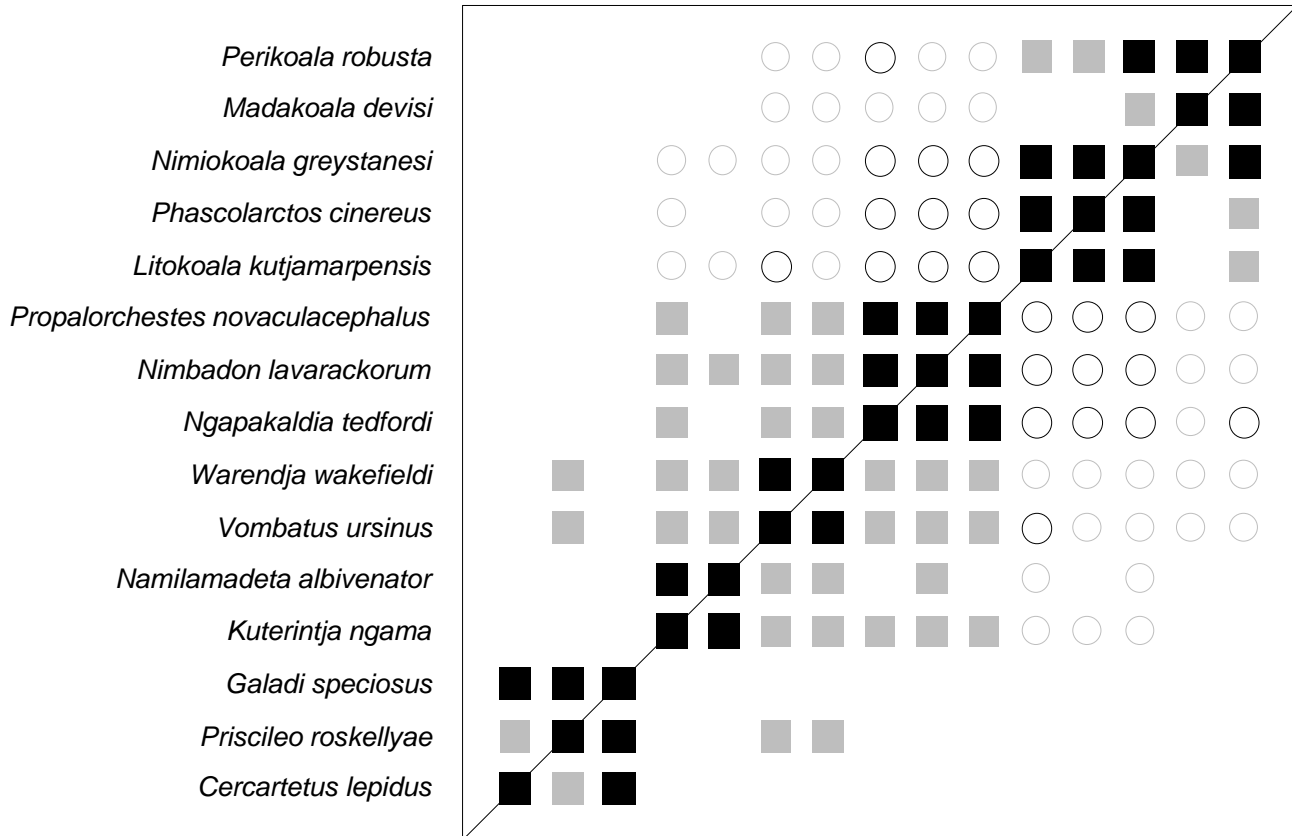
Order Diprotodontia
 Family Pseudocheiridae

| | |
|----------------------------------|------|
| Published taxa | 16 |
| Published characters | 56 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 16 |
| Characters used for calculations | 56 |
| Median bootstrap value | 87.5 |
| F ₉₀ | 0.46 |
| Stress of 3D MDS | 0.04 |
| k _{min} | 4 |
| Conclusion | HB |

Notes: Two holobaramins: Subfamily Pseudochiropinae + *Petropseudes*, and all other taxa.

Black, K.H., M. Archer, and S.J. Hand. 2012. New Tertiary koala (Marsupialia, Phascolarctidae) from Riversleigh, Australia, with a revision of phascolarctid phylogenetics, paleoecology, and paleobiodiversity. *Journal of Vertebrate Paleontology* 32:125-138.

Characters: Craniodental

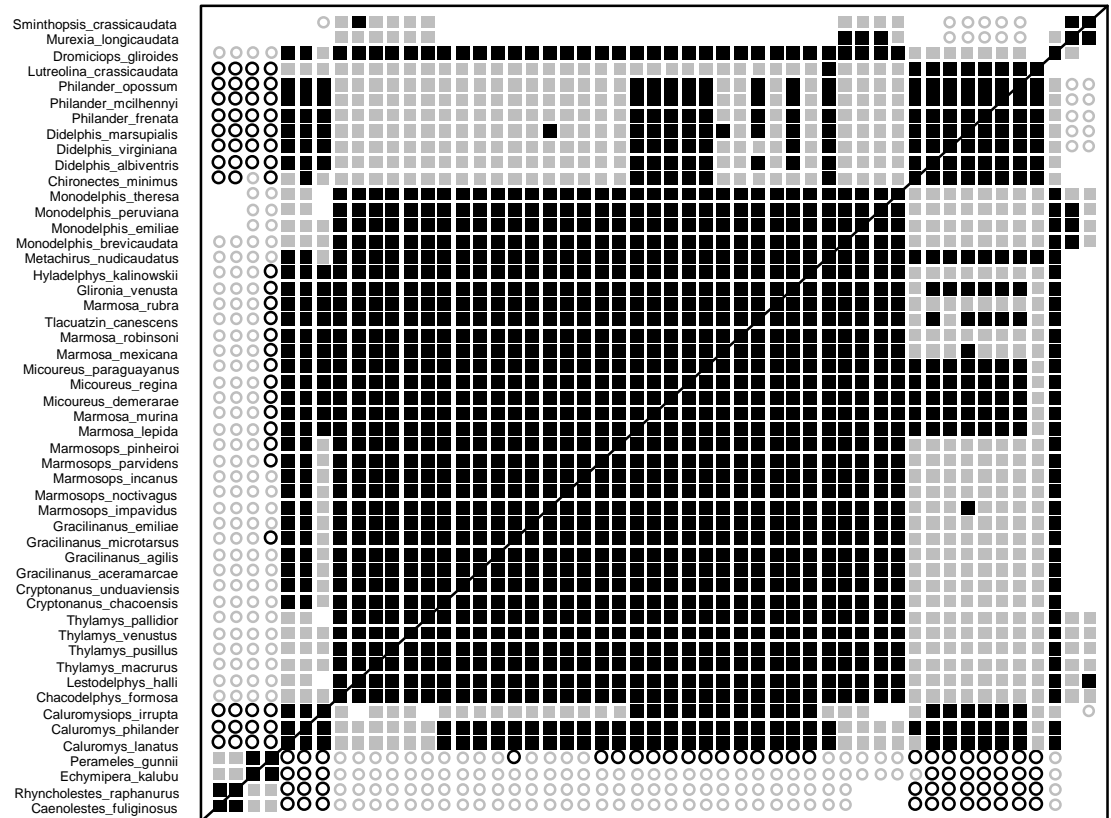


Order Diprotodontia
Family Phascolarctidae

| | |
|----------------------------------|------|
| Published taxa | 19 |
| Published characters | 71 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 15 |
| Characters used for calculations | 50 |
| Median bootstrap value | 86 |
| F ₉₀ | 0.38 |
| Stress of 3D MDS | 0.1 |
| k _{min} | 6 |
| Conclusion | HB |

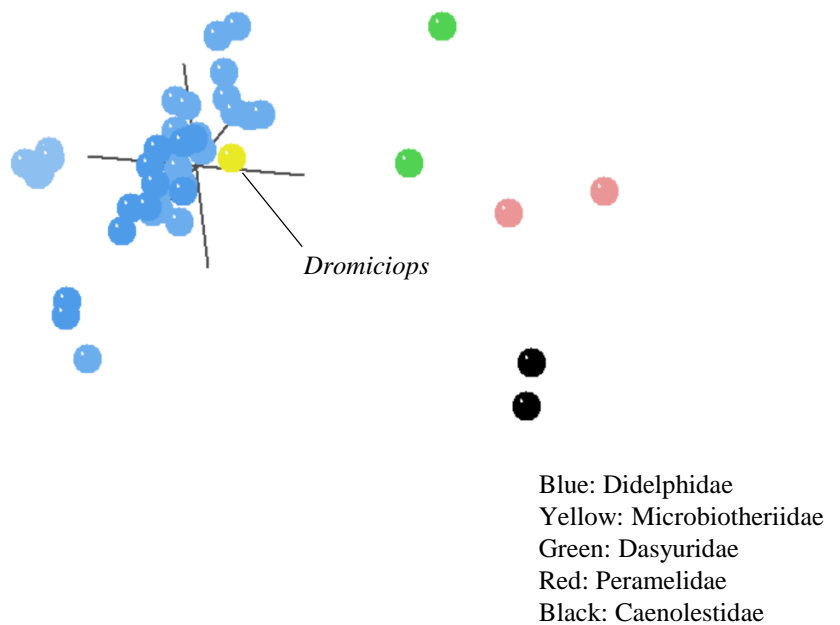
Notes: Phascolarctidae is a holobaramin.

Voss, R.S. and S.A. Jansa. 2009. Phylogenetic relationships and classification of didelphid marsupials, an extant radiation of New World metatherian mammals. *Bulletin of the American Museum of Natural History* 322:1-177. Characters: External morphology, craniodental, karyotype



Order Didelphimorphia
Family Didelphidae

| | |
|----------------------------------|------|
| Published taxa | 51 |
| Published characters | 129 |
| Character relevance | 0.95 |
| Taxic relevance | 0 |
| Taxa used for calculations | 51 |
| Characters used for calculations | 115 |
| Median bootstrap value | 97 |
| F ₉₀ | 0.61 |
| Stress of 3D MDS | 0.21 |
| k _{min} | 8 |
| Conclusion | HB? |

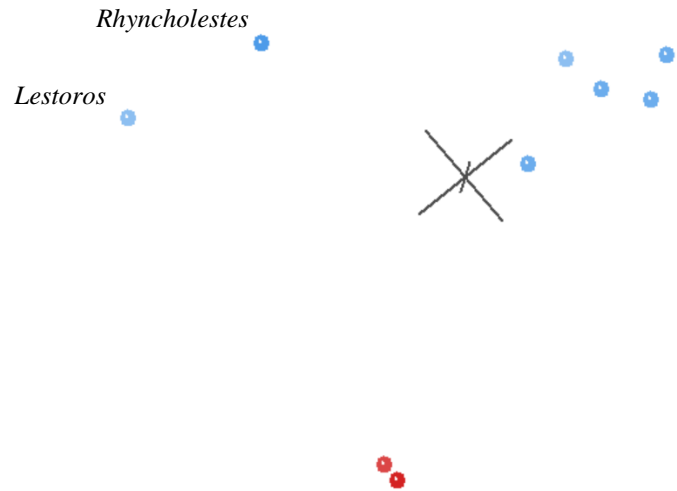
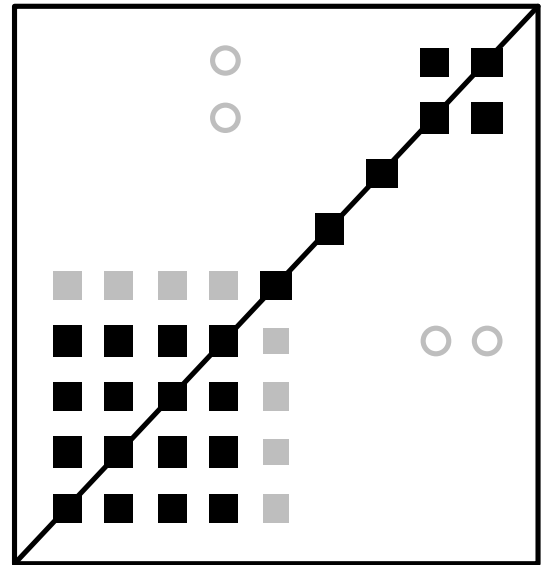


Notes: BDC supports inference of discontinuity between two groups: 1. Peramelidae + Caenolestidae and 2. Didelphidae + Microbiotheriidae + Dasyuridae. MDS supports including microbiotheriid *Dromiciops* in Didelphidae, but dasyurids are more distant. Either Didelphidae + *Dromiciops* or Didelphidae + *Dromiciops* + Dasyuridae is a holobaramin.

Ojala-Barbour, R., C.M. Pinto, M.J. Brito V.L. Albuja T.E. Lee Jr, and B.D. Patterson. 2013. A new species of shrew-opossum (Paucituberculata: Caenolestidae) with a phylogeny of extant caenolestids. *Journal of Mammalogy* 94:967-982.

Characters: External morphology, craniodental

Monodelphis_domestica
 Metachirus_nudicaudatus
 Rhyncholestes_raphanurus
 Lestoros_inca
 Caenolestes_convelatus
 Caenolestes_fuliginosus
 Caenolestes_condorensis
 Caenolestes_sangay
 Caenolestes_caniventer



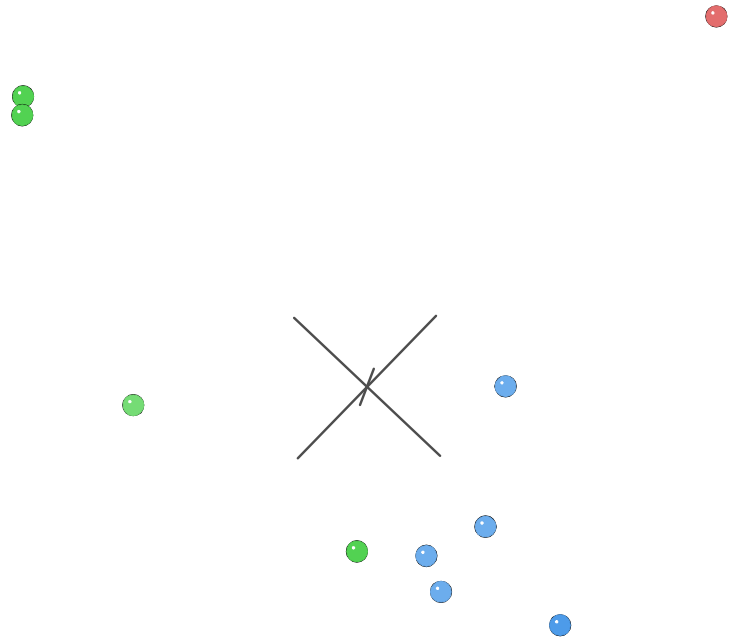
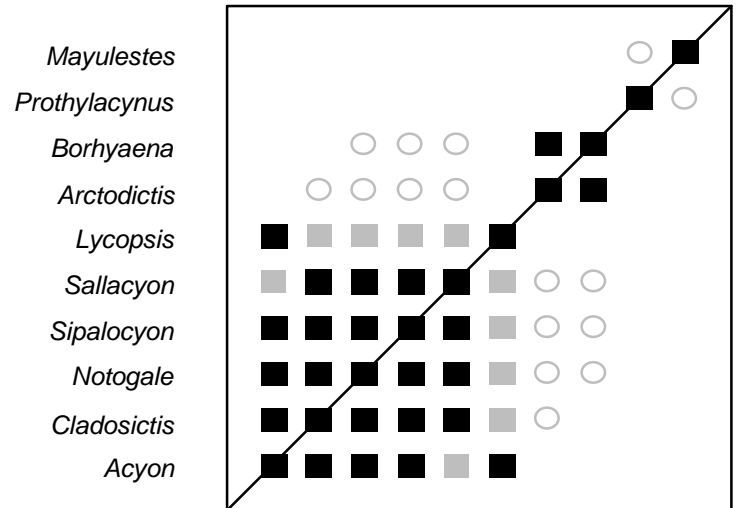
Order Paucituberculata
 Family Caenolestidae

| | |
|----------------------------------|------|
| Published taxa | 9 |
| Published characters | 33 |
| Character relevance | 0.95 |
| Taxic relevance | 0 |
| Taxa used for calculations | 9 |
| Characters used for calculations | 33 |
| Median bootstrap value | 94 |
| F ₉₀ | 0.56 |
| Stress of 3D MDS | 0.08 |
| k _{min} | 5 |
| Conclusion | MB |

Notes: BDC correlations between *Caenolestes* species suggests that at minimum, *Caenolestes* is a monobaramin. There is no evidence that would suggest continuity between the three caenolestid genera or discontinuity of any kind.

Forasiepi, A.M., M.R. Sánchez-Villagra, F.J. Goin, M. Takai, N. Shigehara, and R.F. Kay. 2006. A new species of Hathliacynidae (Metatheria, Sparassodonta) from the middle Miocene of Quebrada Honda, Bolivia. *Journal of Vertebrate Paleontology* 26:670-684.

Characters: Craniodental

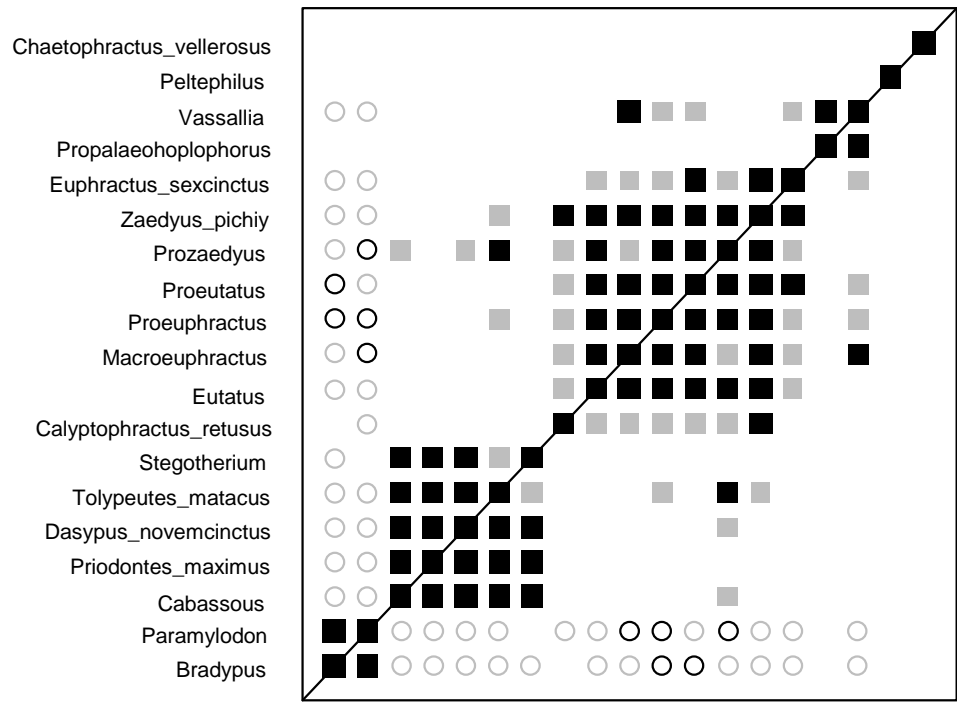


Order Sparassodonta
Family Hathliacynidae

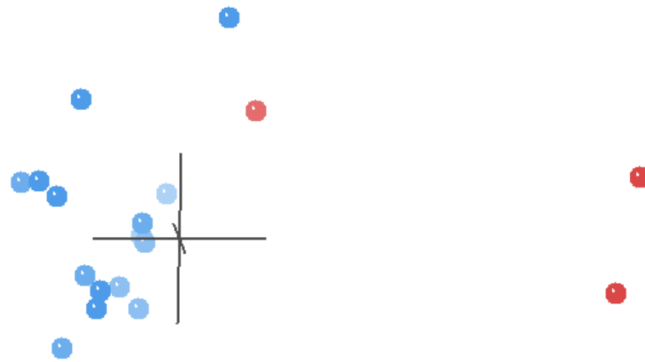
| | |
|----------------------------------|------|
| Published taxa | 10 |
| Published characters | 47 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 10 |
| Characters used for calculations | 39 |
| Median bootstrap value | 89 |
| F ₉₀ | 0.4 |
| Stress of 3D MDS | 0.08 |
| k _{min} | 3 |
| Conclusion | HB? |

Red: outgroup
Green: Borhyaenidae
Blue: Hathliacynidae

Notes: Hathliacynidae + *Lycopsis* could be a holobaramin; evidence of discontinuity has poor bootstrap values.



Order Cingulata
Family Dasypodidae

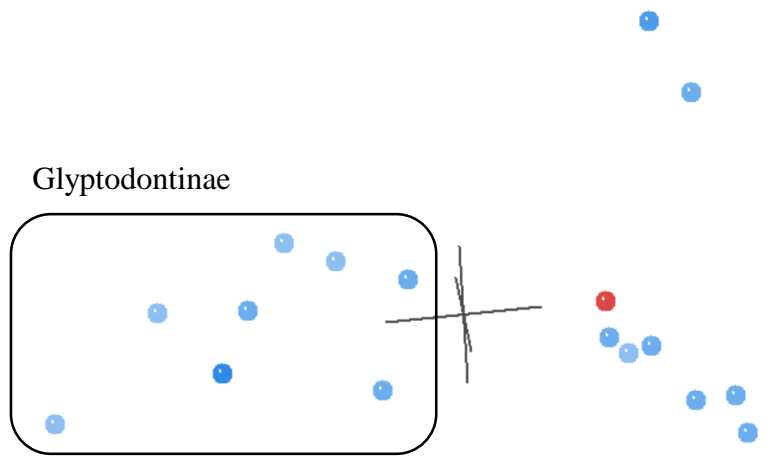
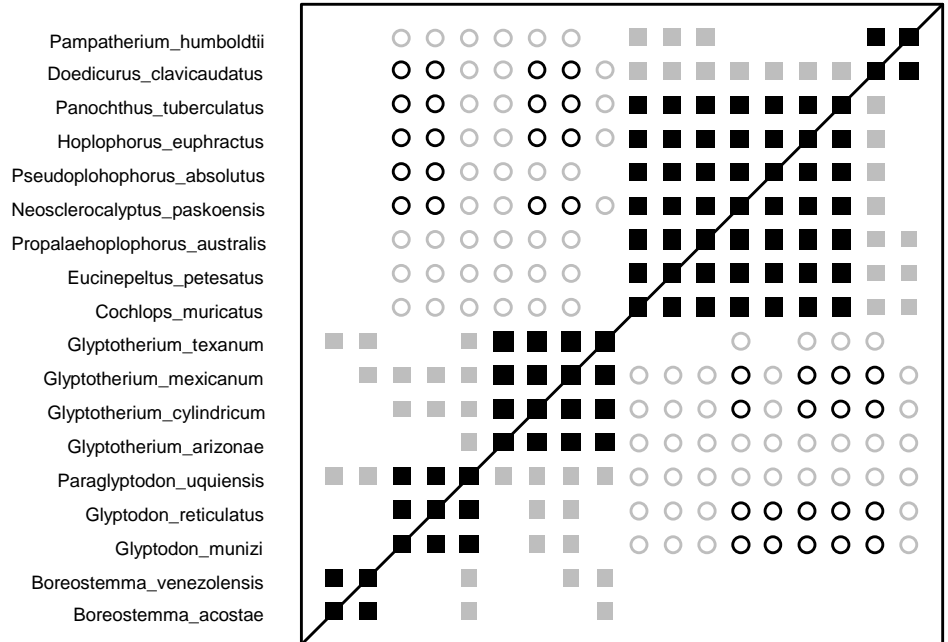


| | |
|----------------------------------|------|
| Published taxa | 22 |
| Published characters | 144 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 19 |
| Characters used for calculations | 106 |
| Median bootstrap value | 94 |
| F ₉₀ | 0.58 |
| Stress of 3D MDS | 0.29 |
| k _{min} | 11 |
| Conclusion | HB |

Notes: Significant, negative BDC between two outgroup taxa and the ingroup taxa is widespread but has poor bootstrap values. Chlamyphorid *Peltephilus* clusters with the dasypodids in both BDC and MDS results. Dasypodidae + *Peltephilus* are a holobaramin. Dasypodid *Chaetophractis* appears to be separate from the holobaramin.

Zurita, A.E., L.R.G. Ruiz, A.J. Gómez-Cruz, and J.E. Arenas-Mosquera. 2013. The most complete known Neogene Glyptodontidae (Mammalia, Xenarthra, Cingulata) from northern South America: taxonomic, paleobiogeographic, and phylogenetic implications. *Journal of Vertebrate Paleontology* 33:696-708.

Characters: Cranial and postcranial



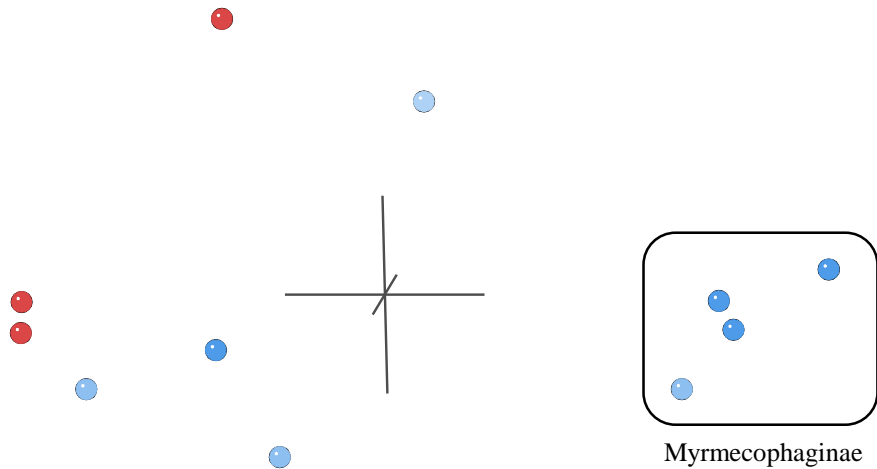
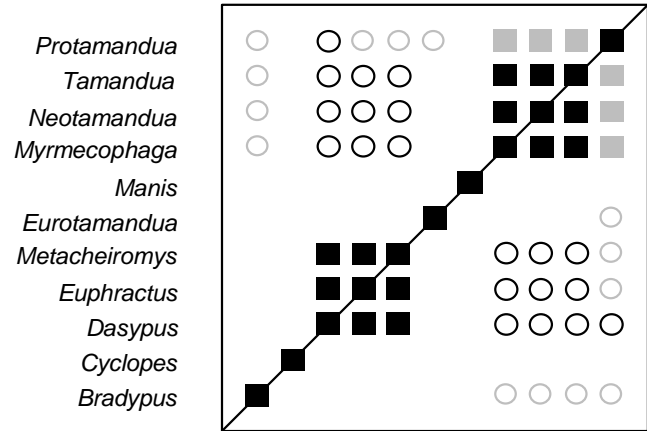
Order Cingulata
Family Glyptodontidae

| | |
|----------------------------------|------|
| Published taxa | 18 |
| Published characters | 26 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 18 |
| Characters used for calculations | 23 |
| Median bootstrap value | 75 |
| F ₉₀ | 0.31 |
| Stress of 3D MDS | 0.13 |
| k _{min} | 4 |
| Conclusion | HB |

Notes: Subfamily Glyptodontinae is well-separated in both BDC and MDS analyses. Glyptodontinae is probably a holobaramin.

Gaudin, T.J. and D.G. Branham. 1998. The phylogeny of the Myrmecophagidae (Mammalia, Xenarthra, Vermilingua) and the relationship of *Eurotamandua* to the Vermilingua. *Journal of Mammalian Evolution* 5:237-265.

Characters: Cranial and postcranial



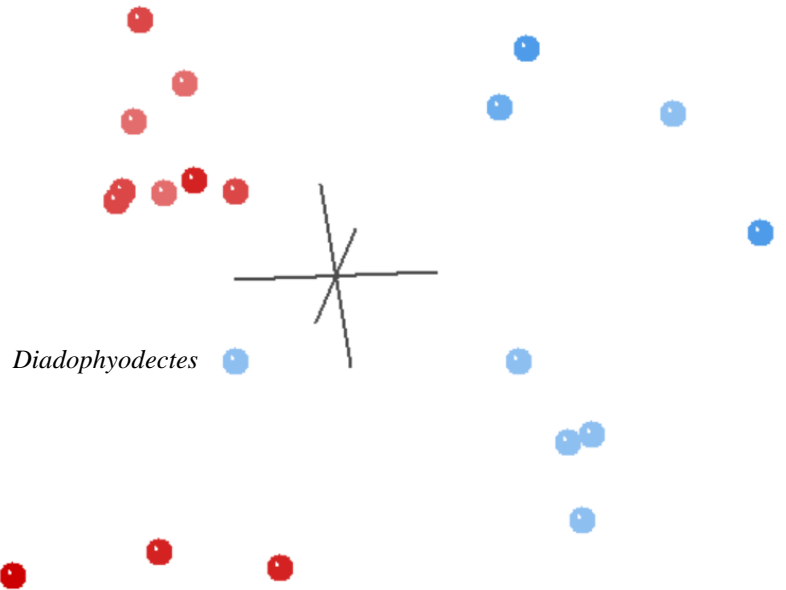
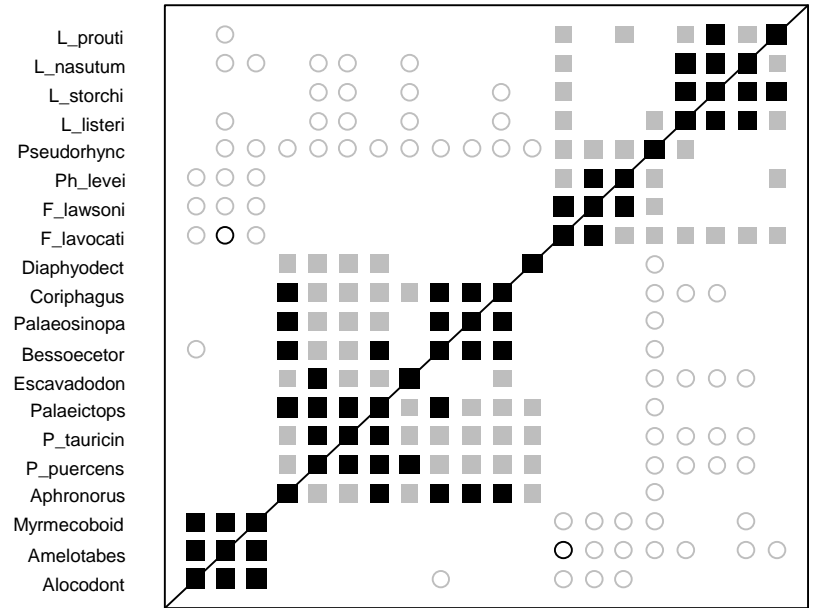
Order Pilosa
Family Myrmecophagidae

| | |
|----------------------------------|------|
| Published taxa | 12 |
| Published characters | 107 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 11 |
| Characters used for calculations | 88 |
| Median bootstrap value | 92 |
| F ₉₀ | 0.58 |
| Stress of 3D MDS | 0.13 |
| k _{min} | 6 |
| Conclusion | HB? |

Notes: Both the MDS and BDC results appear to show a separation between the subfamily Myrmecophaginae (*Myrmecophaga*, *Neotamandua*, *Tamandua*, *Protamandua*) and all other taxa. Myrmecophaginae appears to be a holobaramin.

Hooker, J.J. 2013. Origin and evolution of the Pseudorhynchocyoniidae, a European Paleogene family of insectivorous placental mammals. *Palaeontology* 56:807-835.

Characters: Craniodental



Order Leptictida
Family Pseudorhynchocyoniidae

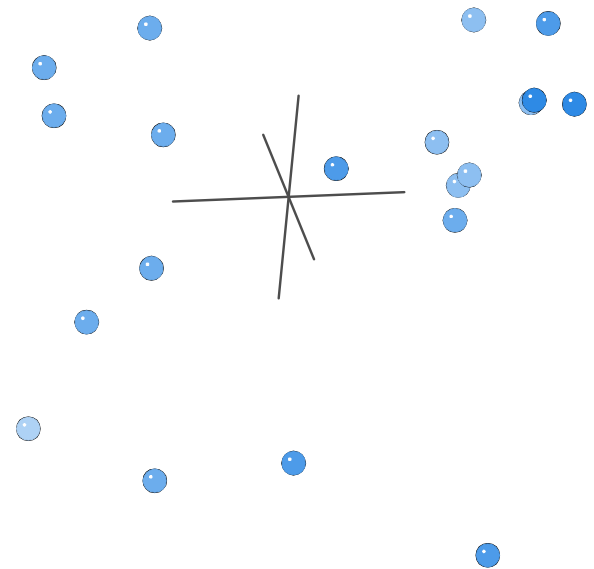
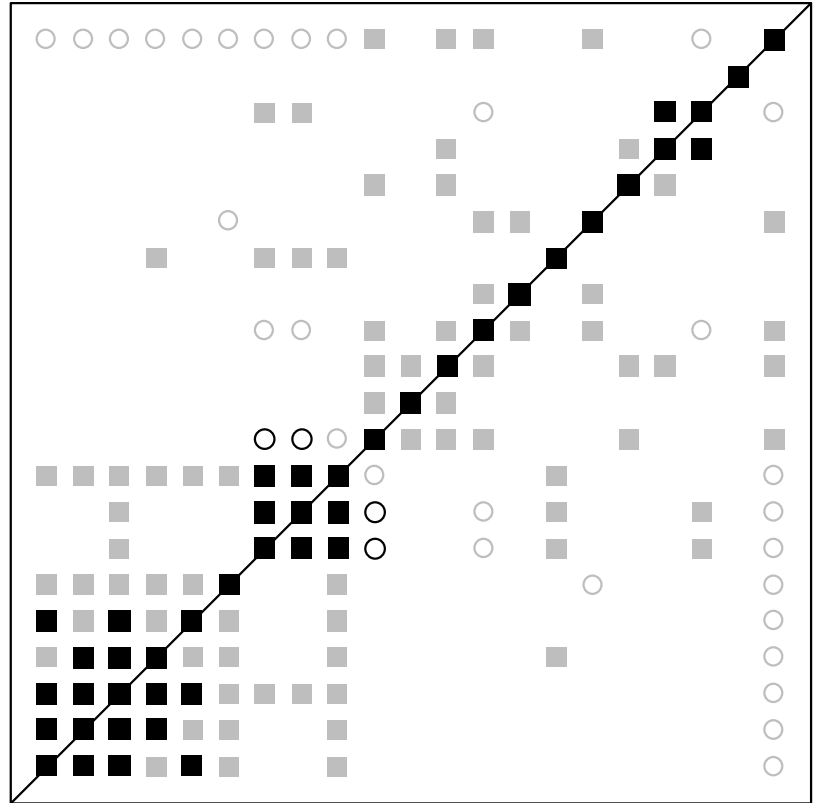
| | |
|----------------------------------|------|
| Published taxa | 21 |
| Published characters | 56 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 21 |
| Characters used for calculations | 38 |
| Median bootstrap value | 78 |
| F ₉₀ | 0.12 |
| Stress of 3D MDS | 0.19 |
| k _{min} | 4 |
| Conclusion | HB? |

Notes: BDC reveals significant, negative BDC with poor bootstrap values, and the MDS shows a diffuse cluster of taxa. Nevertheless, the data support recognizing Pseudorhynchocyoniidae as a provisional holobaramin. The position of *Diadophyodectes*, which clusters with the outgroup in MDS and BDC results, remains uncertain.

Fostowicz-Frelik, Ł., G.J. Frelik, and M. Gasparik. 2010. Morphological phylogeny of pikas (Lagomorpha: Ochotona), with a description of a new species from the Pliocene/Pleistocene transition of Hungary. *Proceedings of the Academy of Natural sciences of Philadelphia* 159:97-118.

Characters: Craniodental

Prolagus sardus
O. pusilla
O. thibetana
O. nihewanica
O. dauurica
O. chowmincheni
O. roylei
O. transcaucasica
O. gudrunae
O. tedfordi
O. lagreli
O. antiqua
O. zabiensis
O. polonica
O. kormosi
O. rufescens
O. dodogolica
O. rutila
O. princeps
O. pallasii
O. alpina



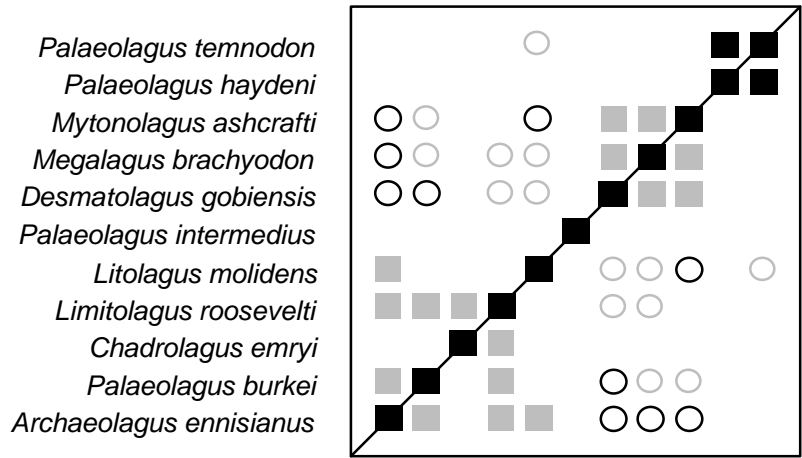
Order Lagomorpha
 Family Ochotonidae

| | |
|----------------------------------|------|
| Published taxa | 21 |
| Published characters | 38 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 21 |
| Characters used for calculations | 24 |
| Median bootstrap value | 69 |
| F ₉₀ | 0.06 |
| Stress of 3D MDS | 0.22 |
| k _{min} | 6 |
| Conclusion | Inc |

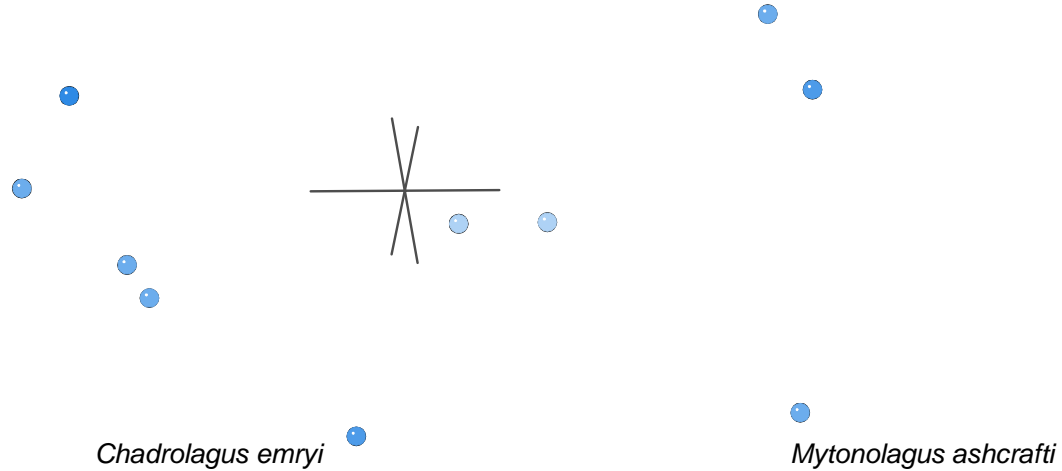
Notes: In BDC results, taxic groups are not evident and bootstrap values are very poor. MDS suggests Ochotonidae forms a cluster distinct from the one outgroup taxon.

Fostowicz-Frelik, Ł. 2013. Reassessment of *Chadrolagus* and *Litolagus* (Mammalia: Lagomorpha) and a new genus of North American Eocene lagomorph from Wyoming. *American Museum Novitates* 3773:1-76.

Characters: Craniodental



Palaeolagus intermedius ●

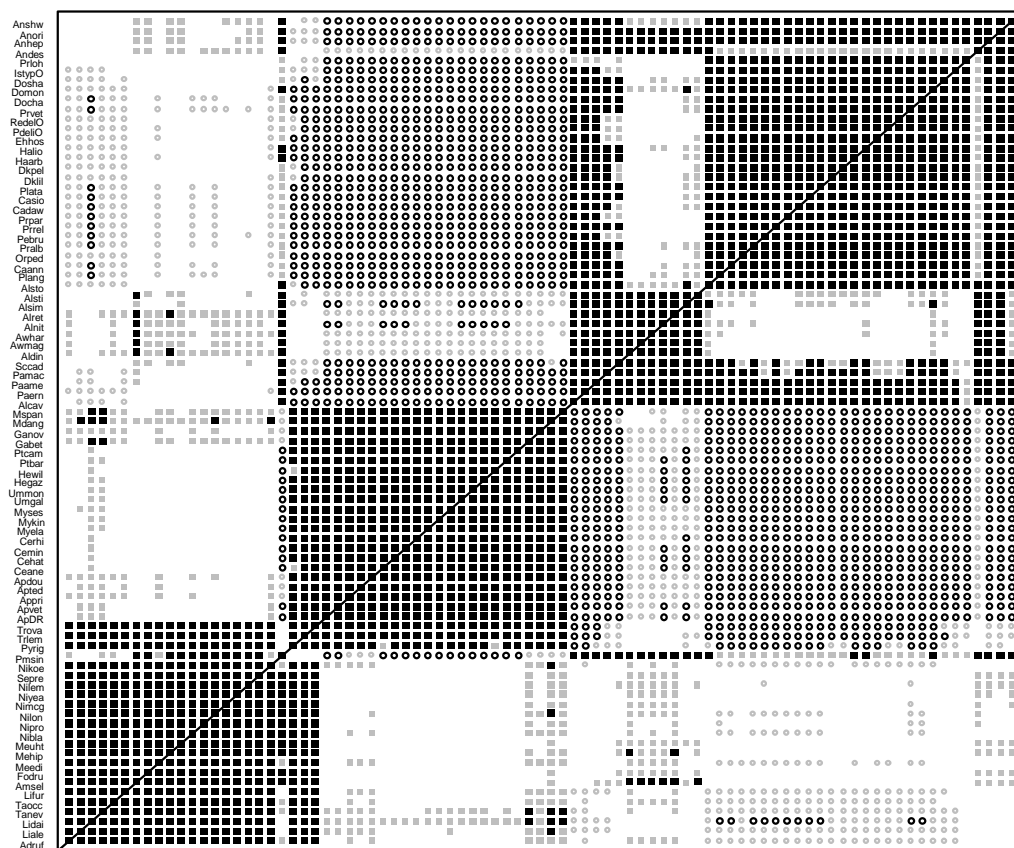


Order Lagomorpha
Family Leporidae

| | |
|----------------------------------|------|
| Published taxa | 11 |
| Published characters | 44 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 11 |
| Characters used for calculations | 34 |
| Median bootstrap value | 89 |
| F ₉₀ | 0.49 |
| Stress of 3D MDS | 0.11 |
| k _{min} | 6 |
| Conclusion | Inc |

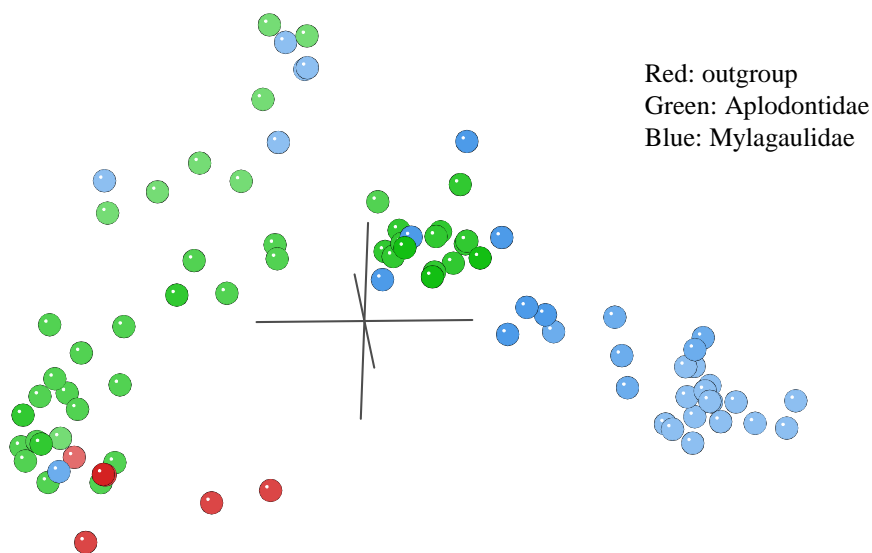
Notes: BDC results have poor bootstrap values, and MDS results indicate a highly diffuse cloud of taxa.

Characters: Craniodental



Order Rodentia
Family Aplodontidae

| | |
|----------------------------------|------|
| Published taxa | 107 |
| Published characters | 249 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 85 |
| Characters used for calculations | 145 |
| Median bootstrap value | 95 |
| F ₉₀ | 0.54 |
| Stress of 3D MDS | 0.15 |
| k _{min} | 7 |
| Conclusion | Inc |

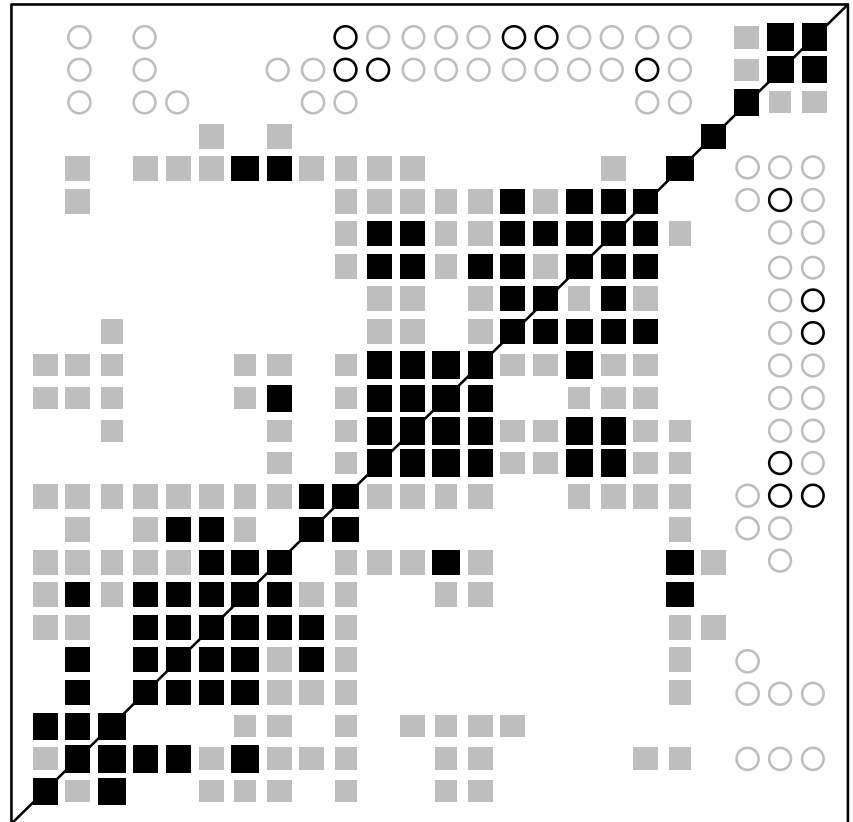


Notes: No evidence of discontinuity within superfamily Aplodontioidea. MDS reveals a cluster of taxa with families blending into one another.

Rybczynski, N. 2007. Castorid phylogenetics: implications for the evolution of swimming and tree-exploitation in beavers. *Journal of Mammalian Evolution* 14:1-35.

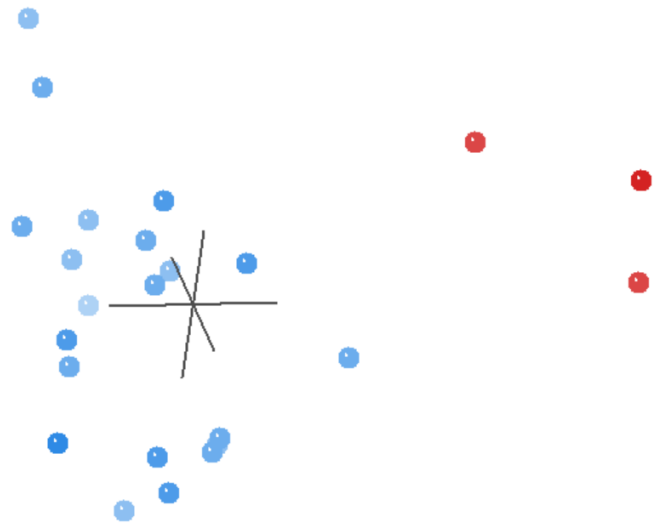
Characters: Cranial and postcranial

Paramys_delicatus
 Paramys_copei
 Eutypomys_thompsoni
 Anchitheriomys
 Castor_fiber
 Pseudopaleocastor_barbouri
 Palaeocastor_magnus
 Palaeocastor_fossor
 Euhapsis_ellicotae
 Euhapsis_breugerom
 Palaeocastor_sp2
 Palaeocastor_cf_simplicidens
 Palaeocastor_cf_nebrascensis
 Capacikala_sp1
 Trogontherium_cuvieri
 Castoroides_ohioensis
 Steneofiber_eseri
 Steneofiber_depereti
 Dipoides_tanneri
 Dipoides_stirtoni
 Dipoides_smithi
 Migmacastor_procumbodens
 Eucastor_tortus
 Agnotocastor_praetereadens



Order Rodentia
 Family Castoridae

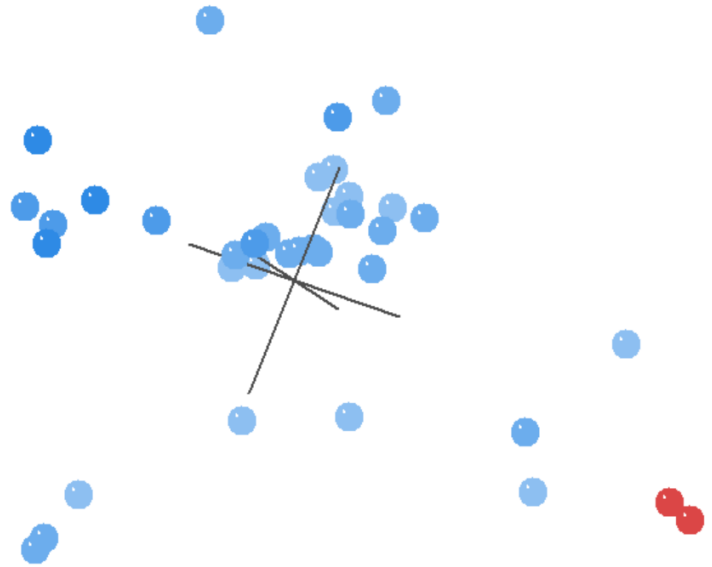
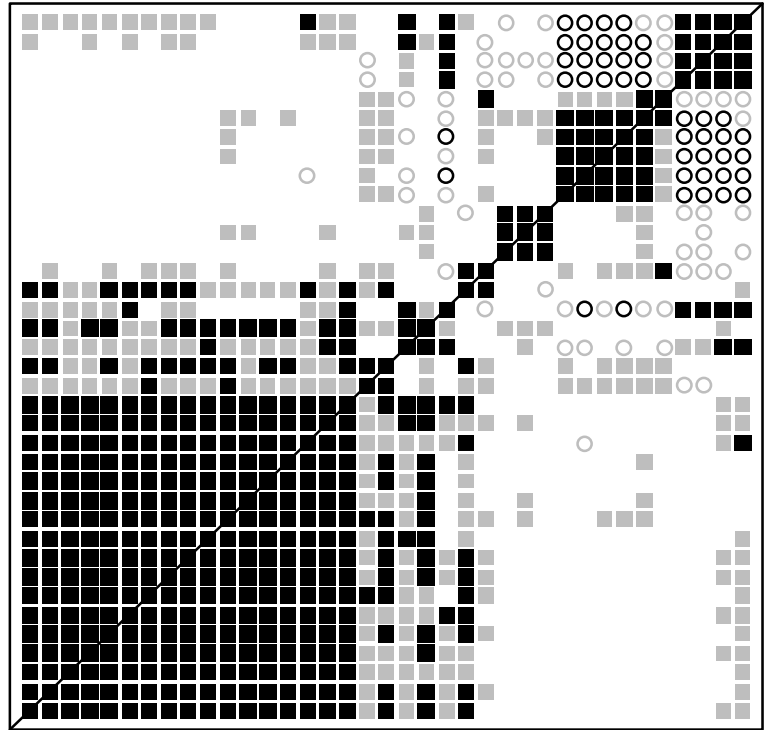
| | |
|----------------------------------|------|
| Published taxa | 39 |
| Published characters | 88 |
| Character relevance | 0.75 |
| Taxic relevance | 0.5 |
| Taxa used for calculations | 24 |
| Characters used for calculations | 41 |
| Median bootstrap value | 78 |
| F ₉₀ | 0.16 |
| Stress of 3D MDS | 0.21 |
| k _{min} | 6 |
| Conclusion | HB |



Notes: Castorid taxa are well separated from the outgroup in both BDC and MDS results. Castoridae appears to be a holobaramin.

Characters: Craniodental

- Pappocricetodon_antiquus
- Palasiomys_conulus
- Primisminthus_yuenus
- Banyuesminthus_uniconjugatus
- Selenomys_mimicus
- Mirabella_tuberosa
- Meteamys_alpani
- Deperetomys_intermedius
- PARACRICETOPS_VIRGATOINCISUS
- Cricetops_dormitor
- Paracricetodon_spectabilis
- Trakymys_saratji
- Paracricetodon_dehmi
- Melissiodon_quercyi
- Edimella_kempeni
- Raricricetodon_minor
- Pappocricetodon_schaubi
- Pappocricetodon_rencunensis
- Muhsinia_steffensi
- Enginia_gertcheki
- Raricricetodon_zhongtiaensis
- Witenia_fusca
- Ulaancricetodon_badamae
- Eucricetodon_longidens
- Eucricetodon_dubius
- Eucricetodon_caducus
- Eucricetodon_asiatricus
- Atavocricetodon_atavus
- Eocricetodon_meridionalis
- Eocricetodon_borealis
- Aralocricetodon_schokensis
- Oxynocricetodon_leptaleos
- Pseudocricetodon_thaleri
- Pseudocricetodon_montalbanensis
- Kerosinia_variabilis
- Heterocricetodon_helbingi
- Adelomyarion_vireti



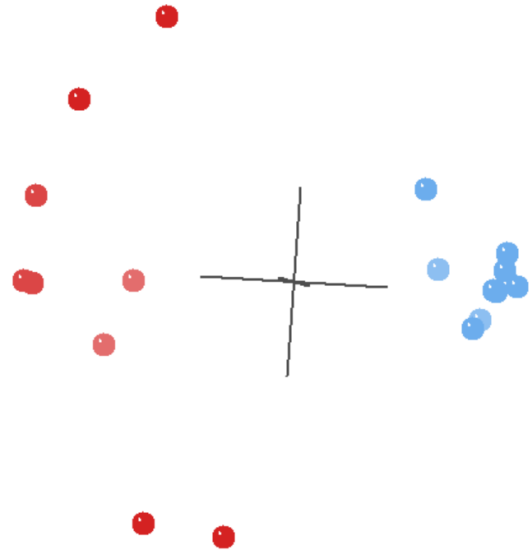
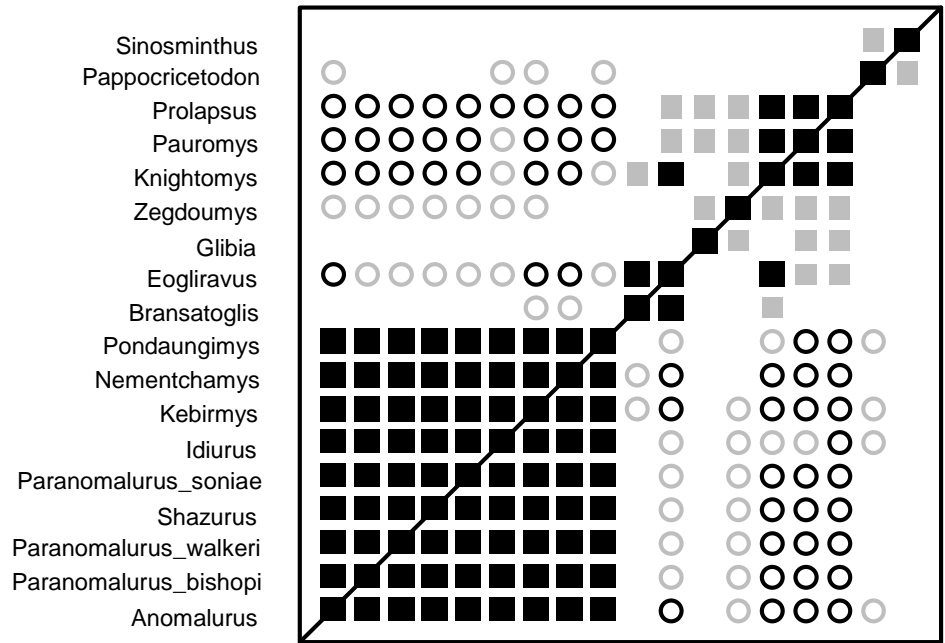
Order Rodentia
Family Cricetidae

| | |
|----------------------------------|------|
| Published taxa | 37 |
| Published characters | 67 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 37 |
| Characters used for calculations | 50 |
| Median bootstrap value | 73 |
| F ₉₀ | 0.34 |
| Stress of 3D MDS | 0.18 |
| k _{min} | 5 |
| Conclusion | Inc |

Notes: No evidence of discontinuity in BDC or MDS results.

Sallam, H.M., E.R. Seiffert, E.L. Simons, and C. Brindley. 2010. A large-bodied anomaluroid rodent from the earliest late Eocene of Egypt: phylogenetic and biogeographic implications. *Journal of Vertebrate Paleontology* 30:1579-1593.

Characters: Craniodental



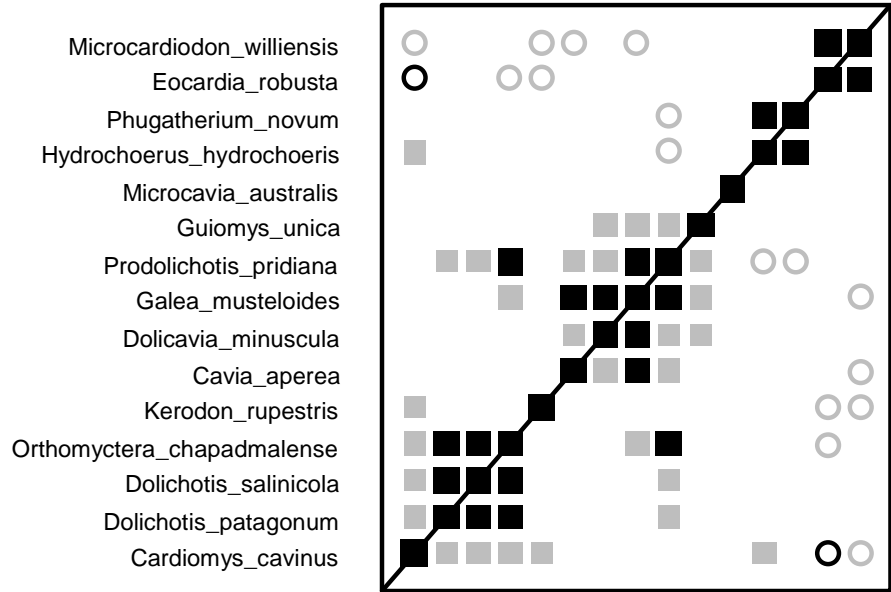
Order Rodentia
Family Anomaluridae

| | |
|----------------------------------|------|
| Published taxa | 29 |
| Published characters | 99 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 18 |
| Characters used for calculations | 72 |
| Median bootstrap value | 89 |
| F ₉₀ | 0.48 |
| Stress of 3D MDS | 0.18 |
| k _{min} | 8 |
| Conclusion | HB |

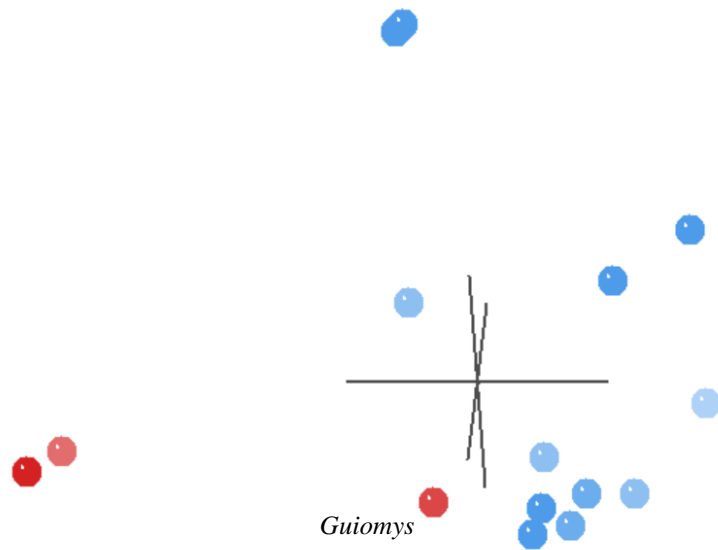
Notes: Anomaluridae form a recognizable cluster in both BDC and MDS results. Anomaluridae is probably a holobaramin.

Pérez, M.E. and M.G. Vucetich. 2011. A new extinct genus of Caviioidea (Rodentia, Hystricognathi) from the Miocene of Patagonia (Argentina) and the evolution of cavioid mandibular morphology. *Journal of Mammalian Evolution* 18:163-183.

Characters: Craniodental



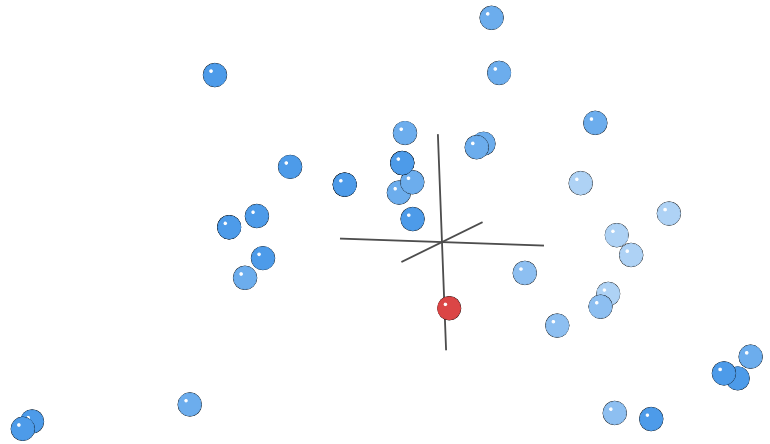
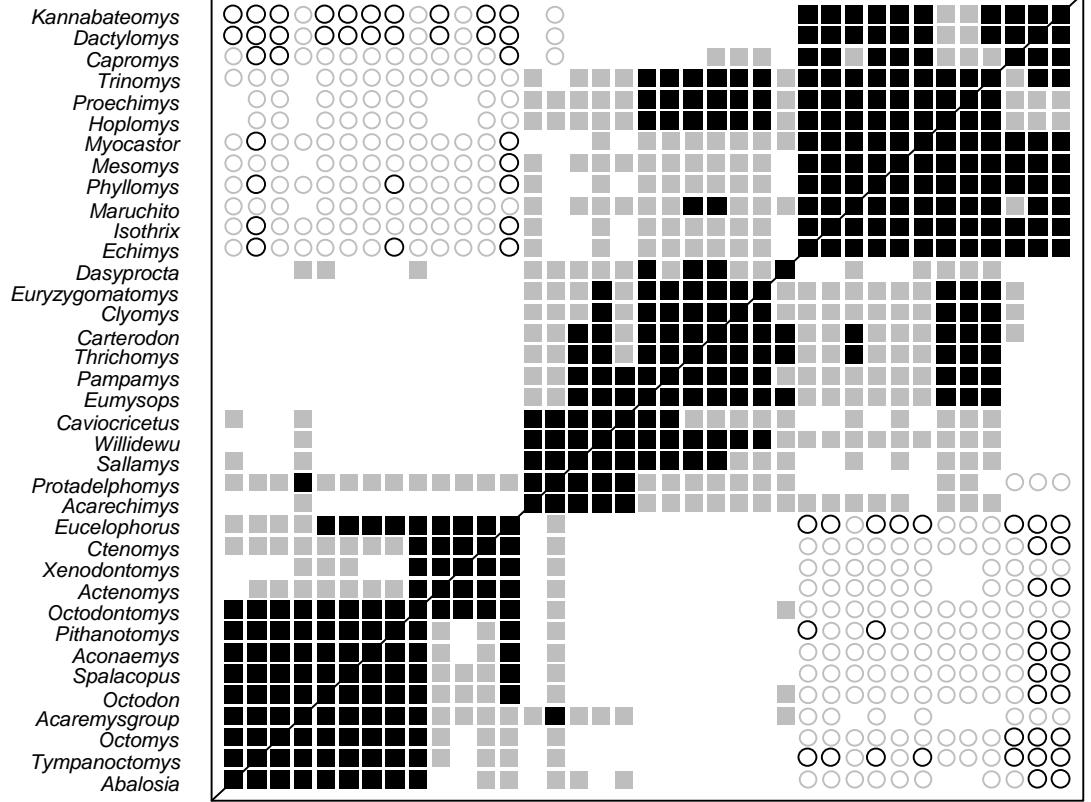
Order Rodentia
 Family Caviidae



| | |
|----------------------------------|------|
| Published taxa | 35 |
| Published characters | 89 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 15 |
| Characters used for calculations | 44 |
| Median bootstrap value | 74 |
| F ₉₀ | 0.1 |
| Stress of 3D MDS | 0.13 |
| k _{min} | 5 |
| Conclusion | HB? |

Notes: Caviidae + *Guiomys* might be a holobaramin. BDC evidence is weak but MDS seems to support a discontinuity.

Characters: Craniodental

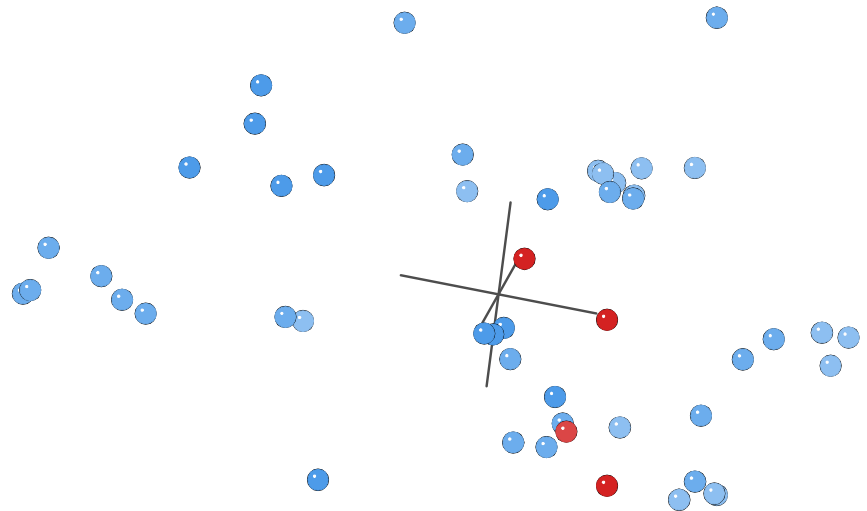
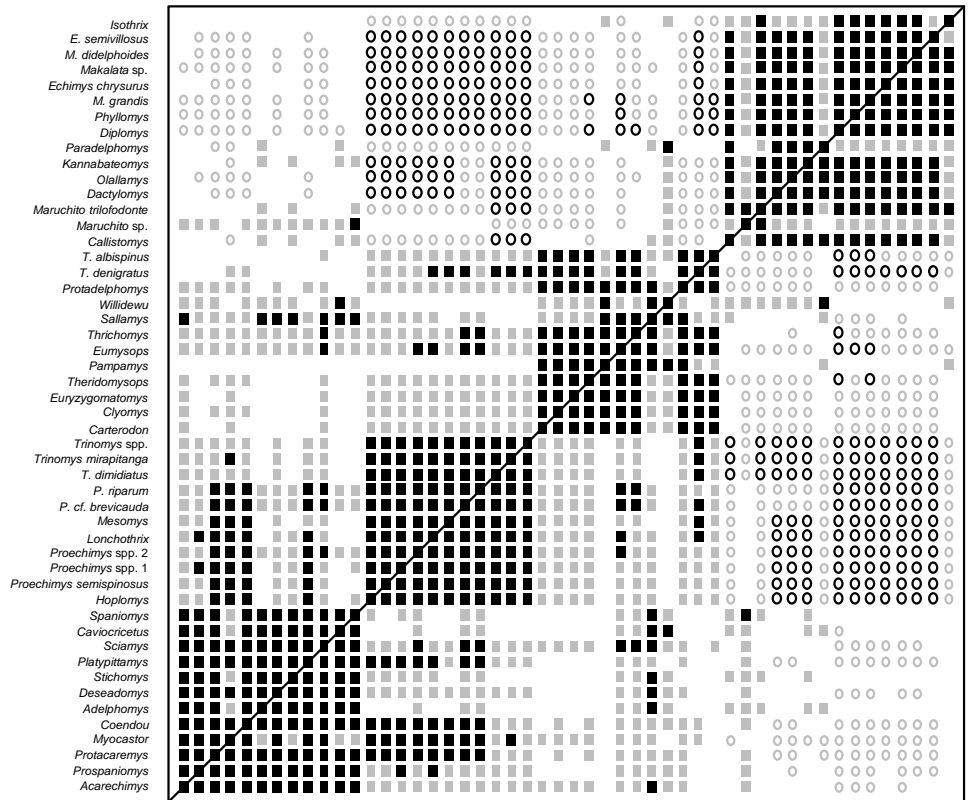


Order Rodentia
Family Octodontidae

| | |
|----------------------------------|------|
| Published taxa | 54 |
| Published characters | 73 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 37 |
| Characters used for calculations | 44 |
| Median bootstrap value | 76 |
| F ₉₀ | 0.29 |
| Stress of 3D MDS | 0.15 |
| k _{min} | 4 |
| Conclusion | Inc |

Notes: In BDC, there appear to be two groups, but *Protadelphomys* shares positive BDC with members of both groups. There is no obvious clustering in the MDS results. There is no clear evidence of discontinuity.

Characters: Craniodental



Order Rodentia
Family Echimyidae

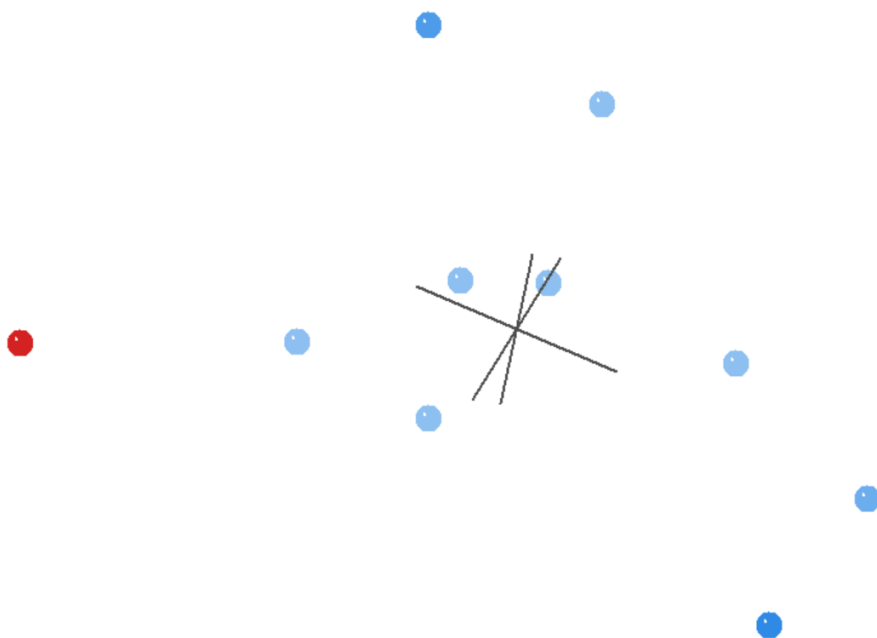
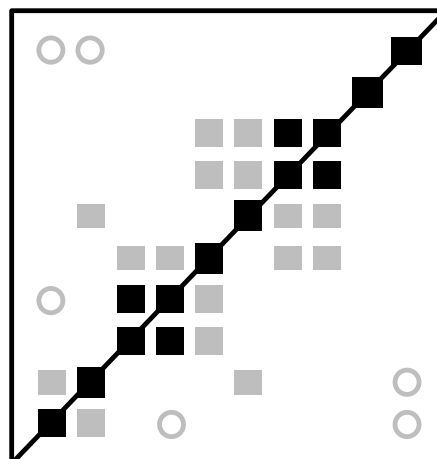
| | |
|----------------------------------|------|
| Published taxa | 54 |
| Published characters | 50 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 50 |
| Characters used for calculations | 37 |
| Median bootstrap value | 78 |
| F ₉₀ | 0.34 |
| Stress of 3D MDS | 0.13 |
| k _{min} | 3 |
| Conclusion | Inc |

Notes: Inconclusive; no strong evidence of discontinuity.

Rankin, B.D. and P.A. Holroyd. 2014. *Aceroryctes dulcis*, a new palaeoryctid (Mammalia, Eutheria) from the early Eocene of the Wasatch Formation of southwestern Wyoming, USA. *Canadian Journal of Earth Sciences* 51:919-926.

Characters: Dental

Asioryctes_nemegetensis
 Palaeoryctes_puercensis
 Palaeoryctes_punctatus
 Palaeoryctes_jepsoni
 Lainoryctes_youzwysnyi
 Eoryctes_melanus
 Ottoryctes_winkleri
 Aceroryctes_dulcis
 Palaeoryctes_cruoris
 Aaptoryctes_ivyi



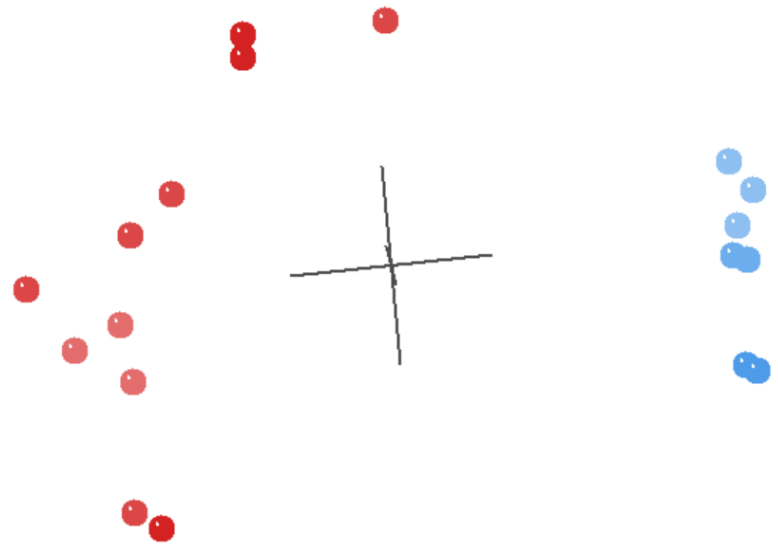
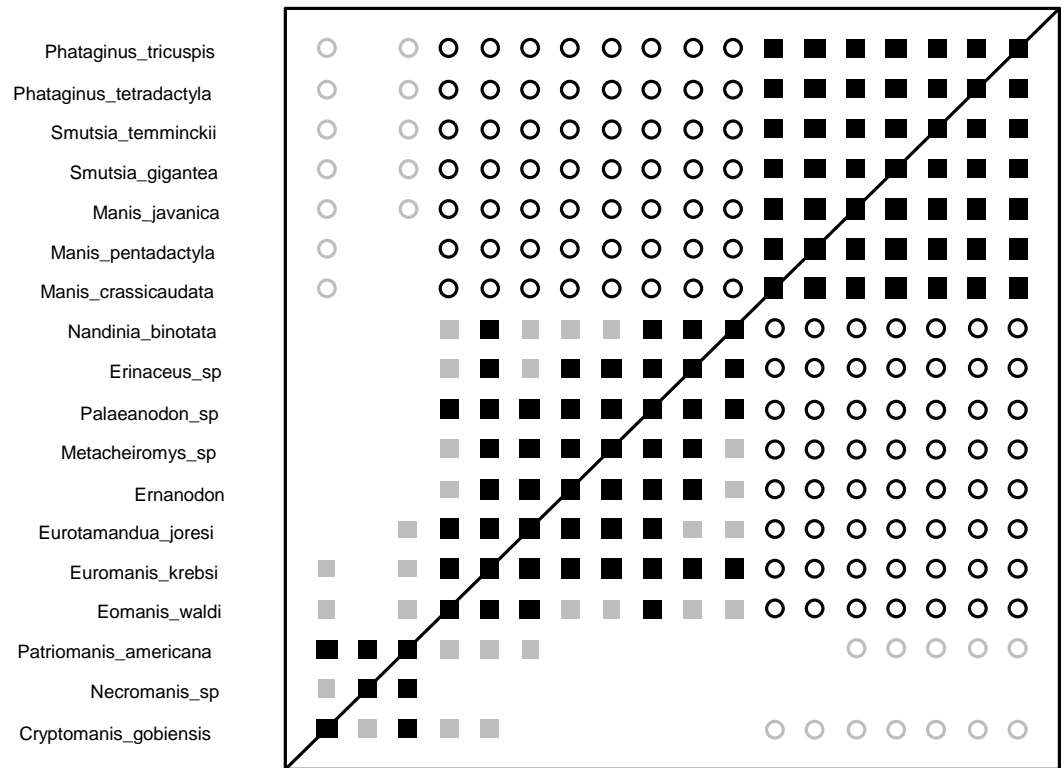
Order Insectivora
 Family Palaeoryctidae

| | |
|----------------------------------|------|
| Published taxa | 10 |
| Published characters | 32 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 10 |
| Characters used for calculations | 18 |
| Median bootstrap value | 78 |
| F ₉₀ | 0.09 |
| Stress of 3D MDS | 0.17 |
| k _{min} | 4 |
| Conclusion | Inc |

Notes: The outgroup taxon *Asioryctes* is moderately separated from the ingroup taxa, but not sufficiently to warrant inferring discontinuity.

Kondrashov, P. and A.K. Agadjanian. 2012. A nearly complete skeleton of *Ernanodon* (Mammalia, Palaeanodonta) from Mongolia: morphofunctional analysis. *Journal of Vertebrate Paleontology* 32:983-1001.

Characters: Cranial and postcranial



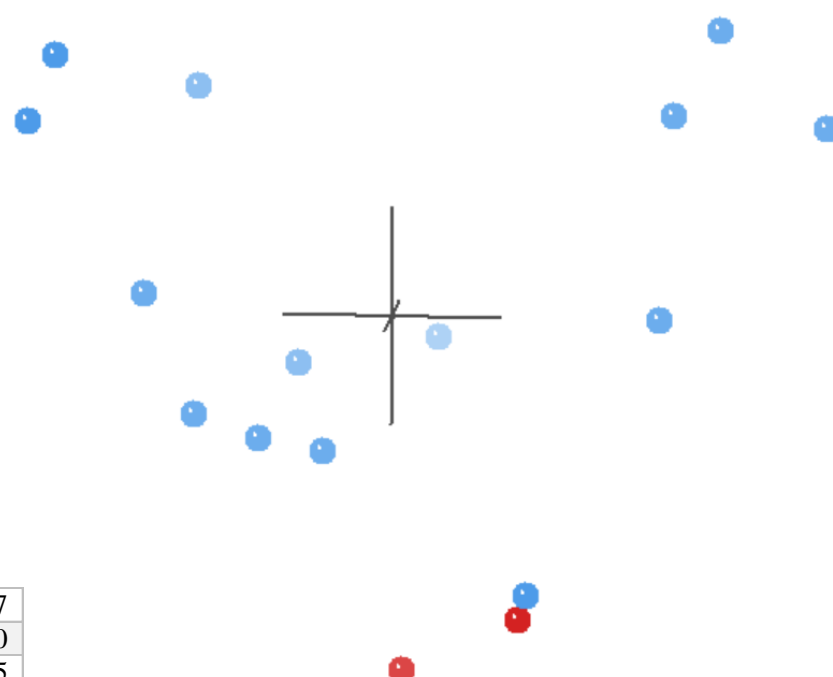
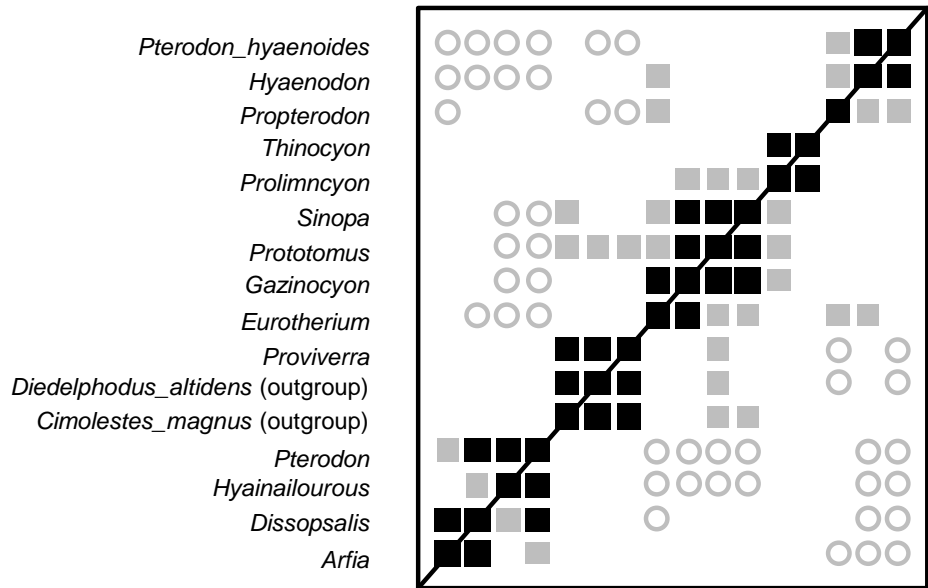
Order Pholidota
Family Manidae

| | |
|----------------------------------|------|
| Published taxa | 18 |
| Published characters | 395 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 18 |
| Characters used for calculations | 209 |
| Median bootstrap value | 100 |
| F ₉₀ | 0.77 |
| Stress of 3D MDS | 0.16 |
| k _{min} | 9 |
| Conclusion | HB |

Notes: Manidae is well-separated in both BDC and MDS results. Manidae is probably a holobaramin.

Polly, P.D. 1996. The skeleton of *Gazinocyon vulpeculus* gen. et comb. nov. and the cladistic relationships of Hyaenodontidae (Eutheria, Mammalia). *Journal of Vertebrate Paleontology* 16:303-319.

Characters: Craniodental and postcranial



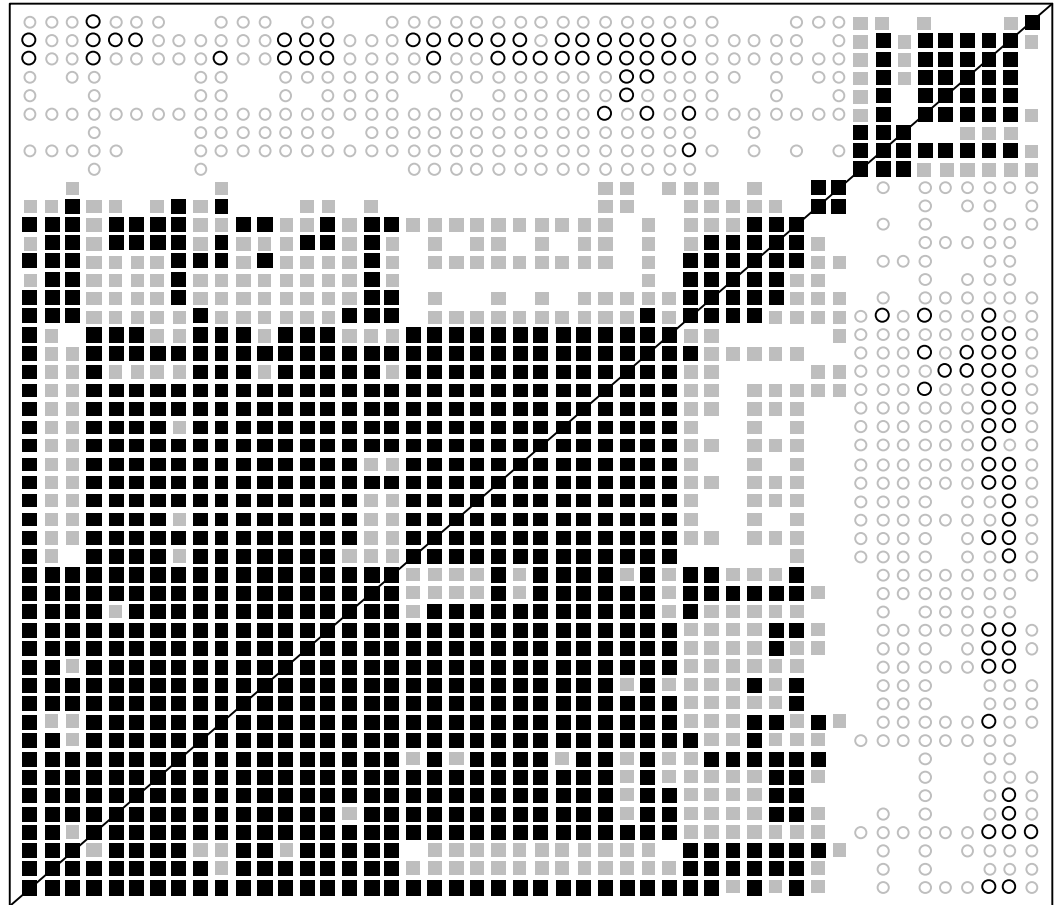
Order Carnivora
Family Hyaenodontidae

| | |
|----------------------------------|------|
| Published taxa | 17 |
| Published characters | 60 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 16 |
| Characters used for calculations | 32 |
| Median bootstrap value | 75 |
| F ₉₀ | 0.12 |
| Stress of 3D MDS | 0.12 |
| k _{min} | 5 |
| Conclusion | Inc |

Notes: The BDC results mostly have poor bootstrap values, and there is no discontinuity evident in the MDS.

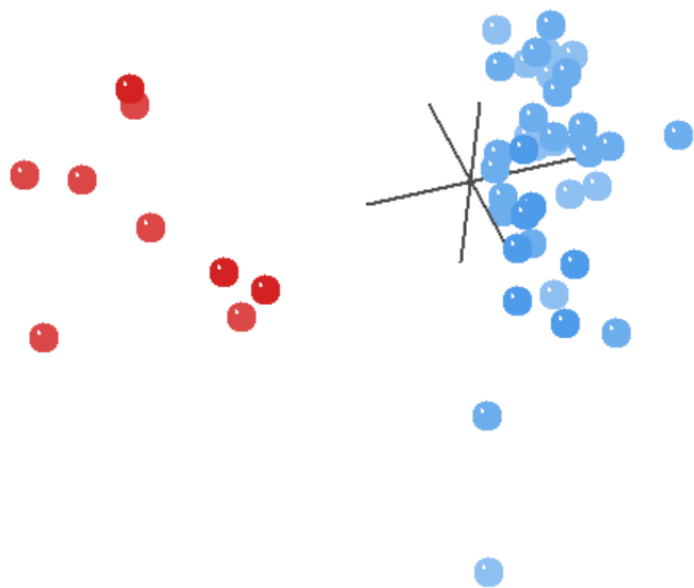
Characters: Craniodental

- Proteles_cristatus
- Palinhyena_reperta_SMNHM37M38
- Palinhyena_reperta_AMNHCHL46
- Lycyaena_chaeretic_AMNHCH11L12
- Palinhyena_reperta_AMNHCh42L344
- Hyaenotherium_wongii_AMNHcomposite
- Hyaena_brunnea
- Hyaena_hyena
- Crocuta_crocuta
- Smilodon_composite
- Panthera_atrox_bebbi
- Pseudaelurus_validusAMNH61803
- Metailurus_sp131854
- Homotherium_nestianum_AMNH104641
- Machairodontinae_genus_indet
- Dinofelis_CB20
- Dinofelis_BC120SMNH
- Felis_yagouaroundi
- Lynx_canadensis
- Lynx_rufus_X9489
- Felis_nigripes
- Felis_geoffroyi
- Felis_colocolo
- Felis_temmincki
- Felis_silvestris
- Felis_viverrina
- Lynx_rufus
- Felis_serval
- Felis_pardalis
- Felis_bengalensis
- Metailurus_parvulusM3895
- Metailurus_major_SMNHM3841M3842
- Metailurus_major
- Panthera_uncia
- Felis_concolor
- Felis_caracal
- Felis_concolor_X8627
- Felis_bituminosa
- Felis_shansius
- Felis_attica
- Panthera_onca
- Panthera_tigris
- Panthera_pardus
- Panthera_leo
- Neofelis_nebulosa
- Felis_atrox_LACM579
- Felis_atrox_LACM290019
- Acinonyx_jubatus



Order Carnivora
Family Felidae

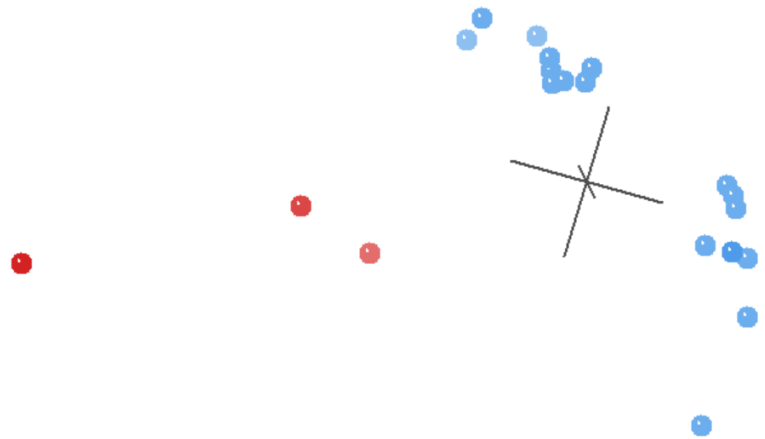
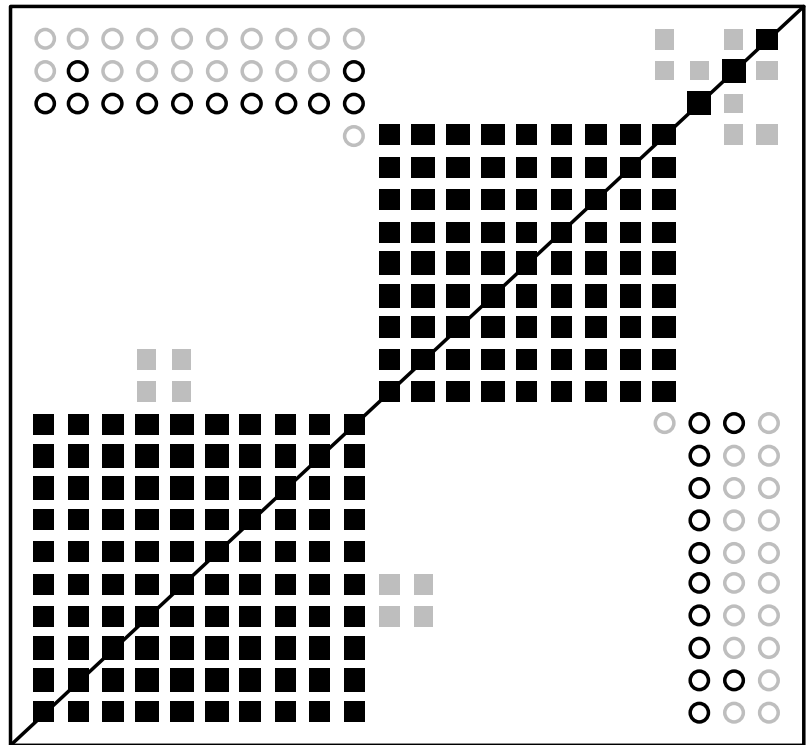
| | |
|----------------------------------|------|
| Published taxa | 168 |
| Published characters | 154 |
| Character relevance | 0.75 |
| Taxic relevance | 0.5 |
| Taxa used for calculations | 48 |
| Characters used for calculations | 103 |
| Median bootstrap value | 87 |
| F ₉₀ | 0.46 |
| Stress of 3D MDS | 0.32 |
| k _{min} | 11 |
| Conclusion | HB |



Notes: The published dataset was trimmed to include just the Felidae and Hyaenidae as in Robinson and Cavanaugh's original analysis (1998. Evidence for a holobaraminic origin of the cats. *CRSQ* 35:2-14). BDC and MDS support a discontinuity between the two families. Felidae is likely a holobaramin.

Characters: Craniodental

- Dinictis_felina
- Cryptoprocta_ferox
- Canis_lupus
- Pseudaelurus_spp.
- Metailurus_parvulus
- Metailurus_major
- Nimravides_ssp.
- Promegantereon_ogygia
- Machairodus_aphanistus
- Dinofelis_petteri
- Dinofelis_cristata
- Dinofelis_barlowi
- Xenosmilus_hodsonae
- Homotherium_serum
- Homotherium_latidens
- Homotherium_crenatidens
- Megantereon_whitei
- Megantereon_nihowanensis
- Megantereon_cultridens
- Smilodon_populator
- Smilodon_fatalis
- Amphimachairodus_giganteus



Order Carnivora
Family Felidae

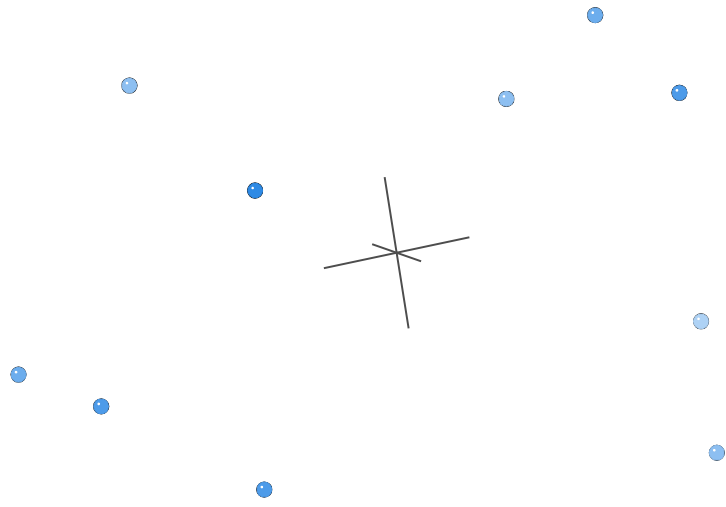
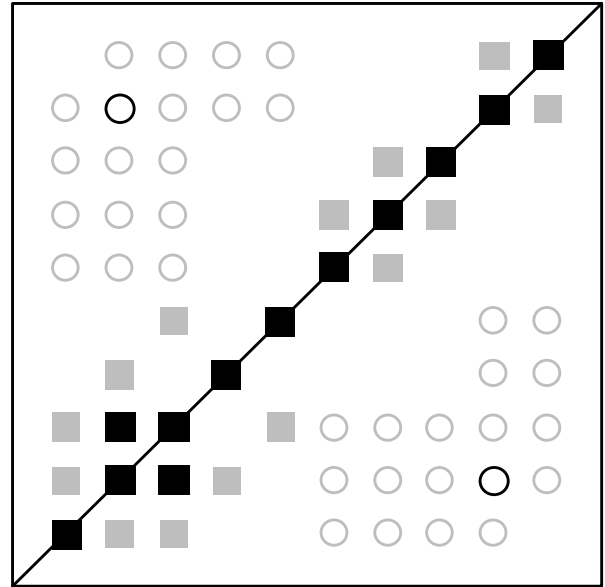
| | |
|----------------------------------|------|
| Published taxa | 22 |
| Published characters | 50 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 22 |
| Characters used for calculations | 50 |
| Median bootstrap value | 85 |
| F ₉₀ | 0.41 |
| Stress of 3D MDS | 0.12 |
| k _{min} | 6 |
| Conclusion | HB |

Notes: Machairodontinae is the focus of this character set, but there are four extinct genera recognized as Felinae by McKenna and Bell (1997): *Pseudaelurus*, *Metailurus*, *Nimravides*, and *Dinofelis*. Results show the felids well-separated from outgroup taxa in MDS and BDC results. BDC indicates two groups of felids with positive BDC between them (but with poor bootstrap values between them). Felidae is likely a holobaramin.

Morlo, M., S. Peigne, and D. Nagel. 2004. A new species of *Prosansanosmilus*: implications for the systematic relationships of the family Barbouroufelidae new rank (Carnivora, Mammalia). *Zoological Journal of the Linnean Society* 140:43-61.

Characters: Dental

Nimravus intermedius
Eofelis edwardsii
Ginsburgsmilus
Proailurus lemanensis
Early Pseudaelurus
Prosansanosmilus eggeri
Afrosmilus turkanae
P. peregrinus
A. hispanicus
A. africanus

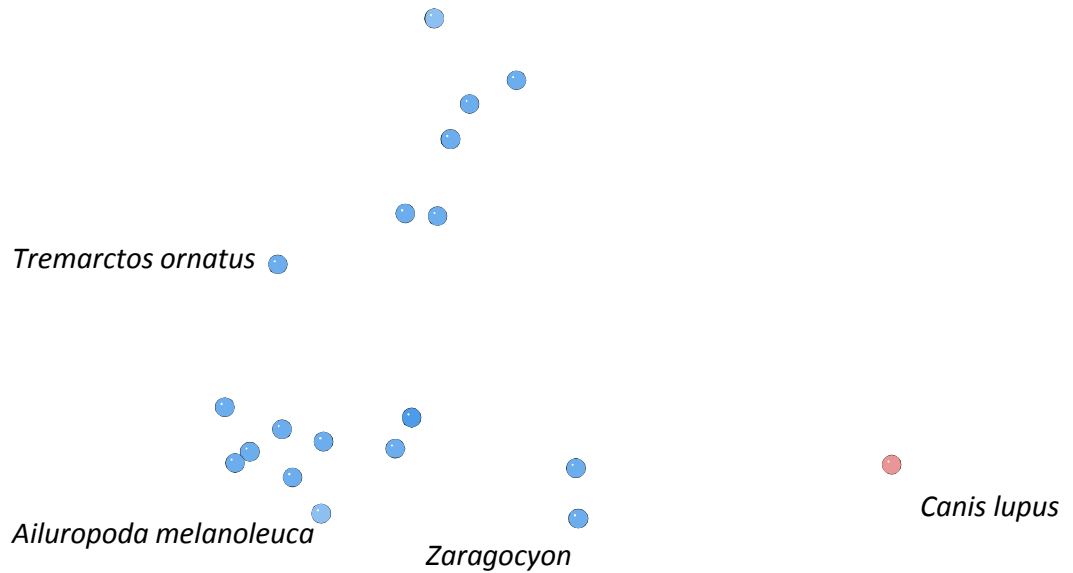
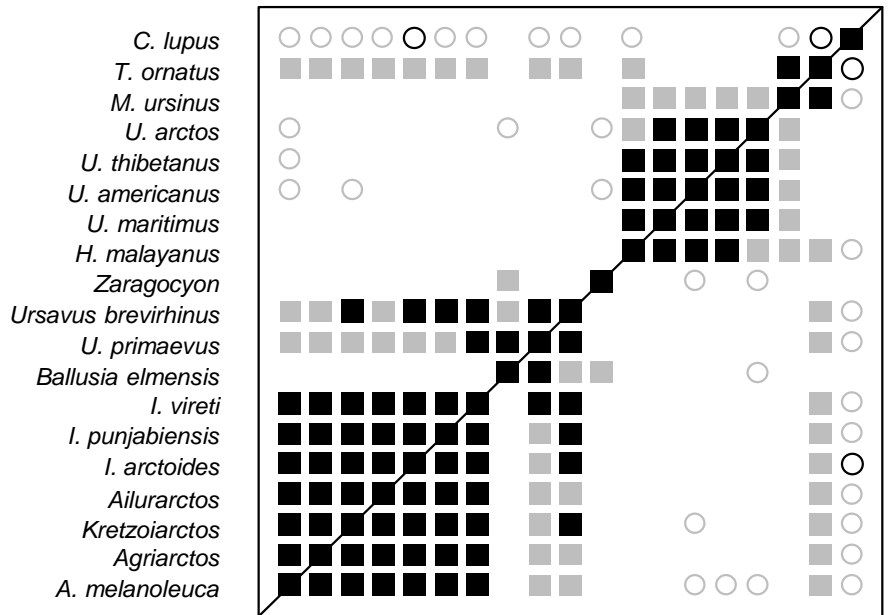


Order Carnivora
 Subfamily Barbouroufelinae

| | |
|----------------------------------|------|
| Published taxa | 11 |
| Published characters | 22 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 10 |
| Characters used for calculations | 16 |
| Median bootstrap value | 71 |
| F ₉₀ | 0.11 |
| Stress of 3D MDS | 0.1 |
| k _{min} | 3 |
| Conclusion | Inc |

Notes: The BDC results have poor bootstrap values, and the MDS reveals a diffuse distribution of taxa. There is no evidence of discontinuity.

Characters: Craniodental



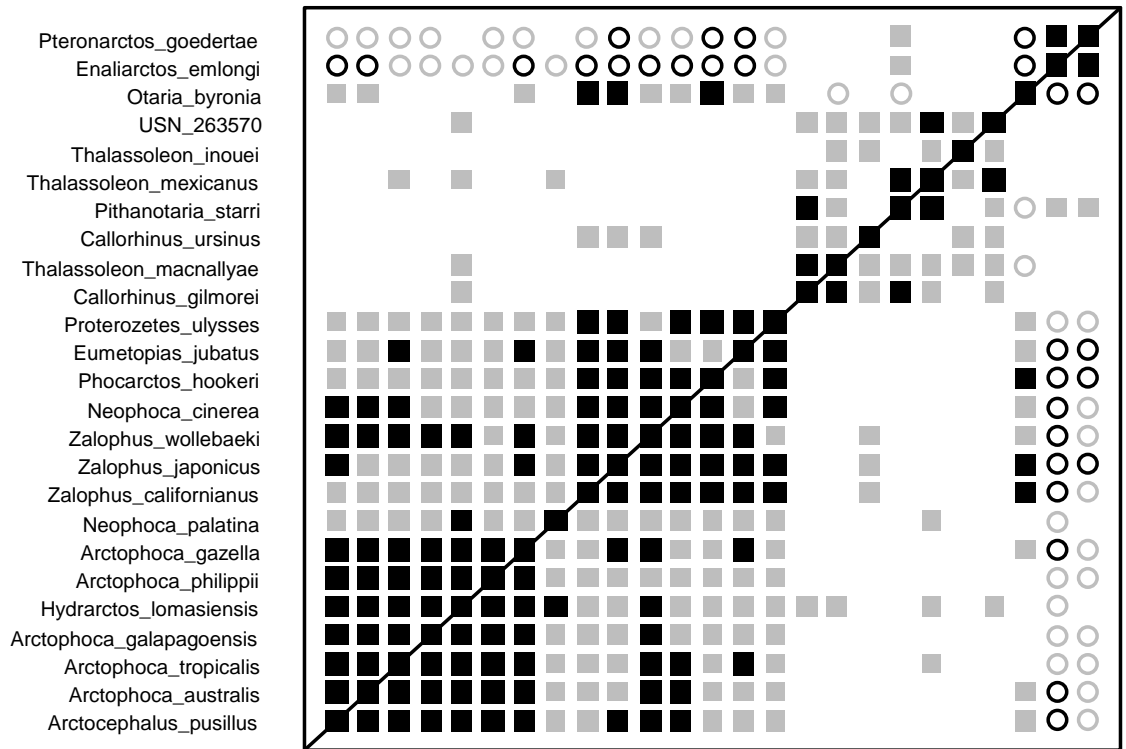
Order Carnivora
Family Ursidae

| | |
|----------------------------------|------|
| Published taxa | 19 |
| Published characters | 82 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 19 |
| Characters used for calculations | 53 |
| Median bootstrap value | 82 |
| F ₉₀ | 0.23 |
| Stress of 3D MDS | 0.14 |
| k _{min} | 5 |
| Conclusion | HB? |

Notes: Ursidae is well-separated from the outgroup taxon *Canis lupus* in both BDC and MDS results. Within the group, ursids form two groups that are connected only by the spectacled bear *Tremarctos ornatus*, which also appears in an intermediate position in the MDS results. Ursidae appears to be a holobaramin.

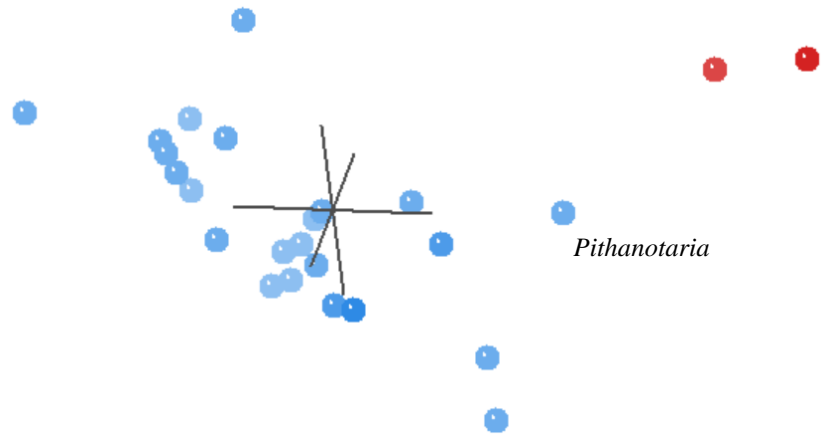
Churchill, M., R.W. Boessenecker, and M.T. Clementz. 2014. Colonization of the southern hemisphere by fur seals and sea lions (Carnivora: Otariidae) revealed by combined evidence phylogenetic and Bayesian biogeographical analysis. *Zoological Journal of the Linnean Society* 172:200-225.

Characters: Craniodental, postcranial, behavioral



Order Carnivora
Family Otariidae

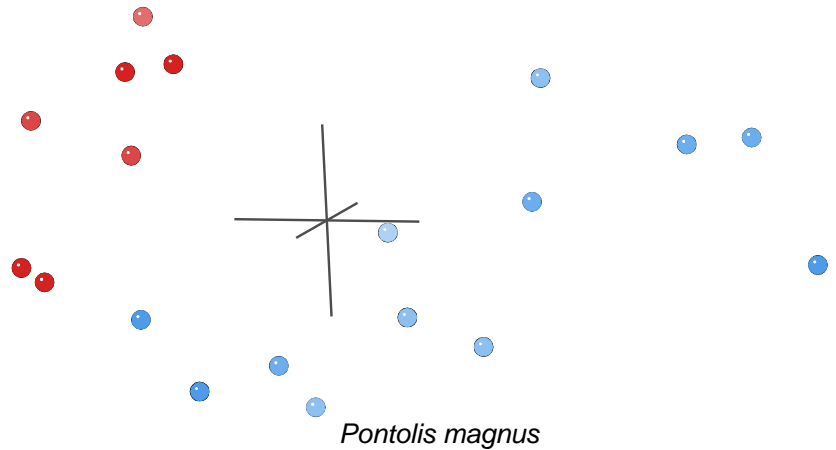
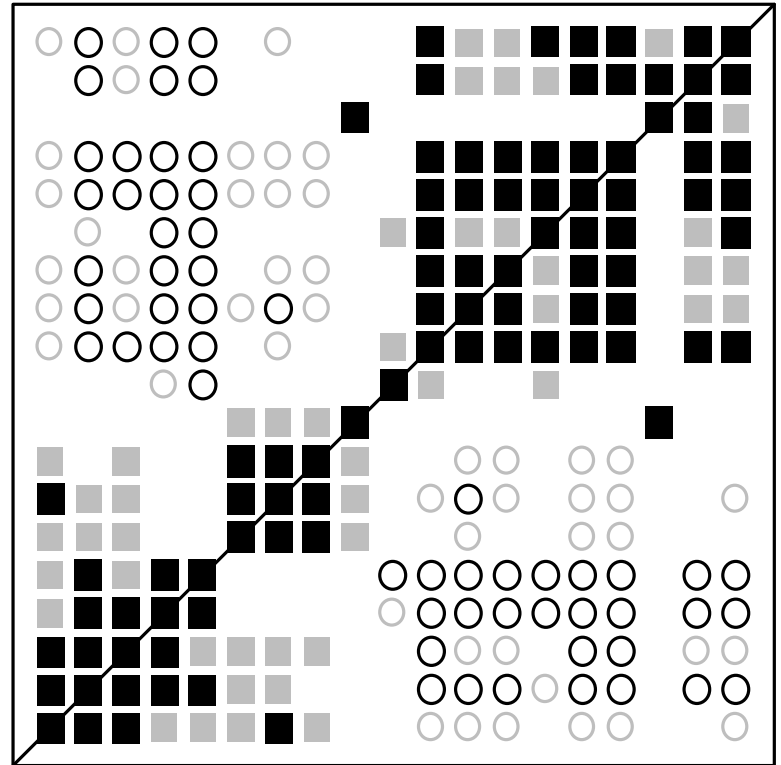
| | |
|----------------------------------|------|
| Published taxa | 25 |
| Published characters | 107 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 25 |
| Characters used for calculations | 84 |
| Median bootstrap value | 80 |
| F ₉₀ | 0.24 |
| Stress of 3D MDS | 0.26 |
| k _{min} | 10 |
| Conclusion | HB |



Notes: Otariidae is well-separated from the outgroup taxa in both BDC and MDS results. BDC reveals positive correlation between the outgroup taxa and *Pithanotaria*, but their proximity is not evident in the MDS results. Otariidae is likely a holobaramin.

Characters: Craniodental and postcranial

- Proneotherium repenningi*
- Neotherium mirum*
- Imagotaria downsi*
- Pteronarctos goedertae*
- Enaliarctos emlongi*
- Callorhinus ursinus*
- Monachus monachus*
- Erignathus barbatus*
- Desmatophoca oregonensis*
- Allodesmus gracilis*
- Pontolis magnus*
- Gomphotaria pugnax*
- Dusignathus seftoni*
- Dusignathus santacruzensis*
- Valenictus chulavistensis*
- Odobenus rosmarus*
- Protodobenus japonicus*
- Ontocetus emmonsi*
- Aivukus cedrosensis*



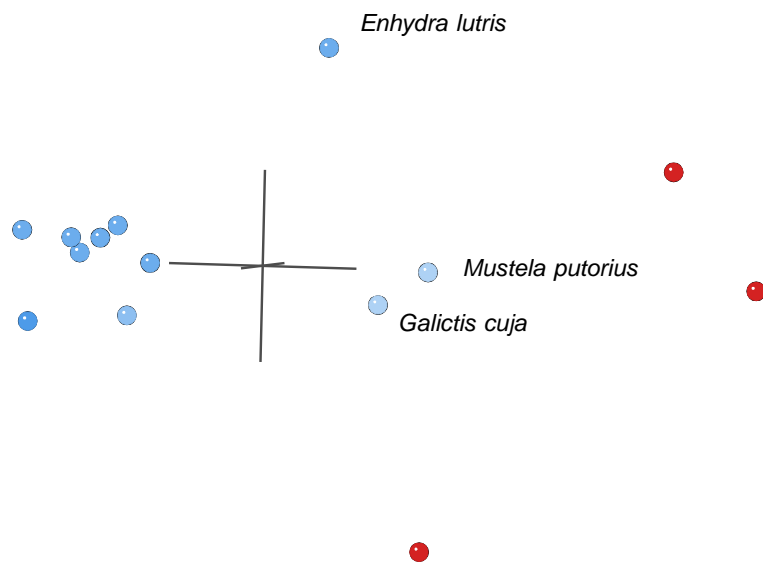
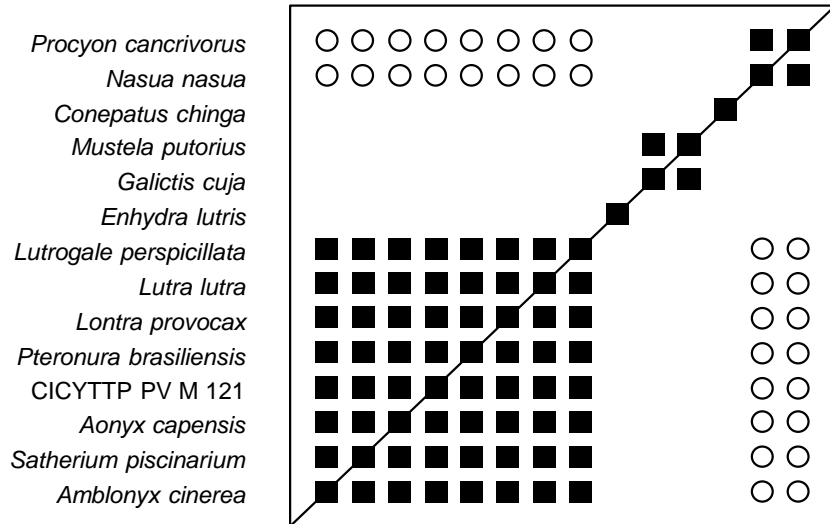
Order Carnivora
Family Odobenidae

| | |
|----------------------------------|------|
| Published taxa | 23 |
| Published characters | 90 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 19 |
| Characters used for calculations | 82 |
| Median bootstrap value | 89 |
| F ₉₀ | 0.48 |
| Stress of 3D MDS | 0.14 |
| k _{min} | 7 |
| Conclusion | HB? |

Notes: Odobenidae is well separated from the outgroup taxa in the BDC results, with the exception of *Pontolis*, which is positively correlated with members of the ingroup and outgroup. The MDS results do not reveal a clear distinction between the two groups, although *Pontolis* does not appear to be closely associated with the outgroup taxa. Odobenidae might be a holobaramin.

Prevosti, F.J. and B.S. Ferrero. 2008. A Pleistocene giant river otter from Argentina: remarks on the fossil record and phylogenetic analysis. *Journal of Vertebrate Paleontology* 28:1171-1181.

Characters: Craniodental



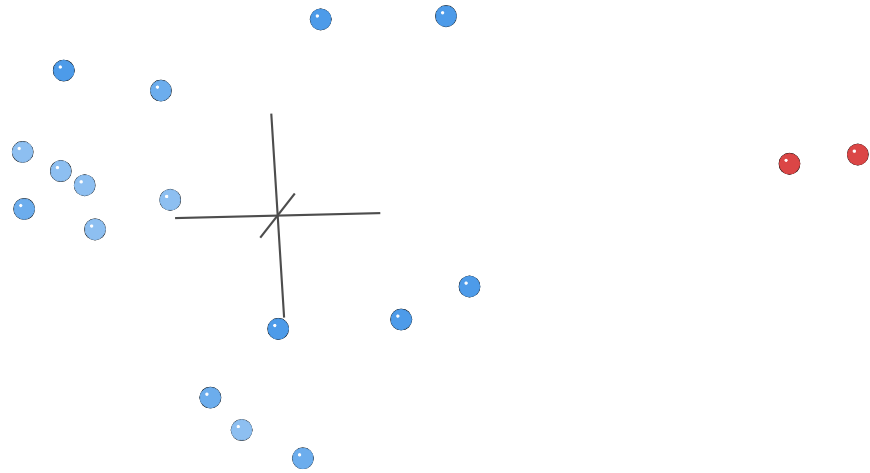
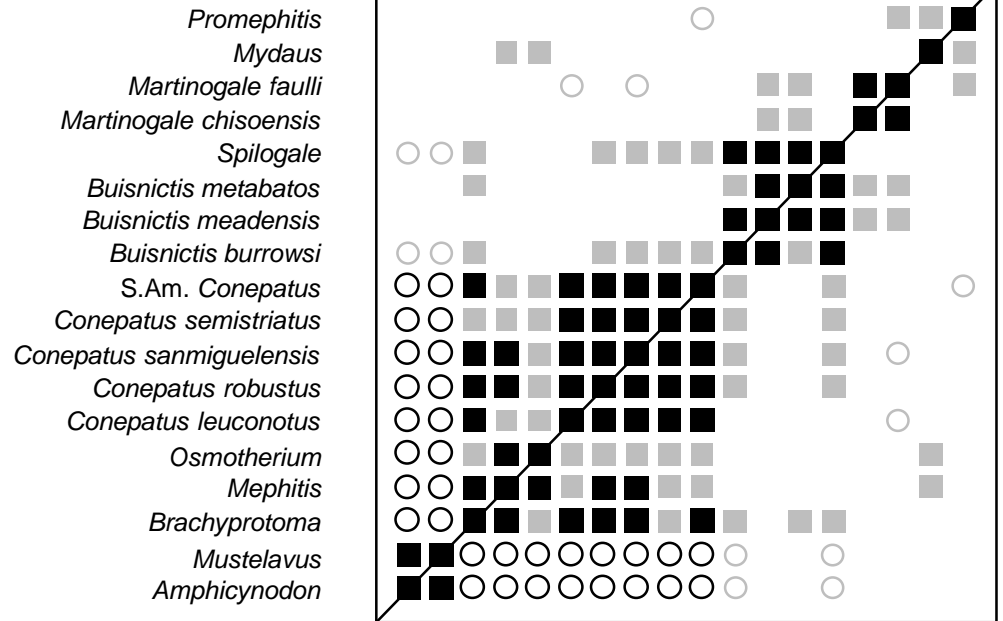
Order Carnivora
Family Mustelidae

| | |
|----------------------------------|------|
| Published taxa | 41 |
| Published characters | 69 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 14 |
| Characters used for calculations | 64 |
| Median bootstrap value | 99 |
| F ₉₀ | 0.78 |
| Stress of 3D MDS | 0.11 |
| k _{min} | 6 |
| Conclusion | HB? |

Notes: A cluster of mustelid taxa appear to be a holobaramin, but *Mustela*, *Galictis* and *Enhydra* are not connected to the main cluster of mustelid taxa in either the MDS or BDC results. This result should be examined further.

Wang, X., Ó. Carranza-Castañeda, and J.J. Aranda-Gómez. 2014. A transitional skunk, *Buisnictis metabatos* sp. nov. (Mephitidae, Carnivora), from Baja California Sur and the role of southern refugia in skunk evolution. *Journal of Systematic Palaeontology* 12:291-302.

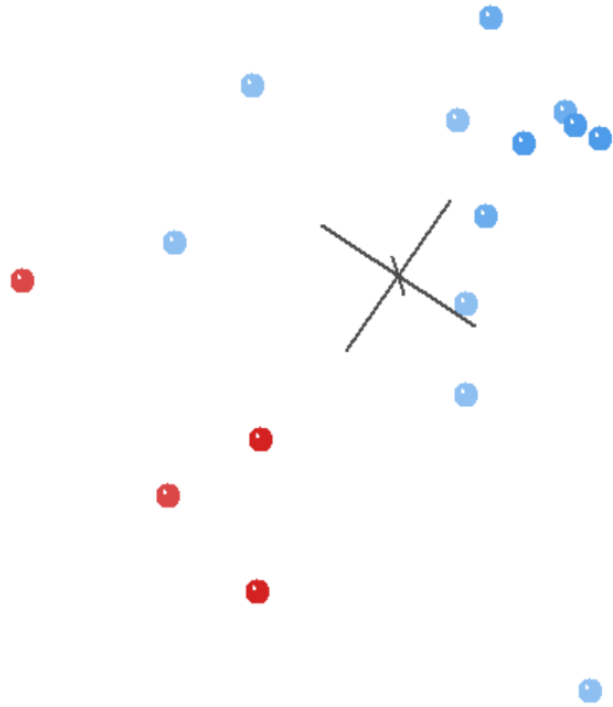
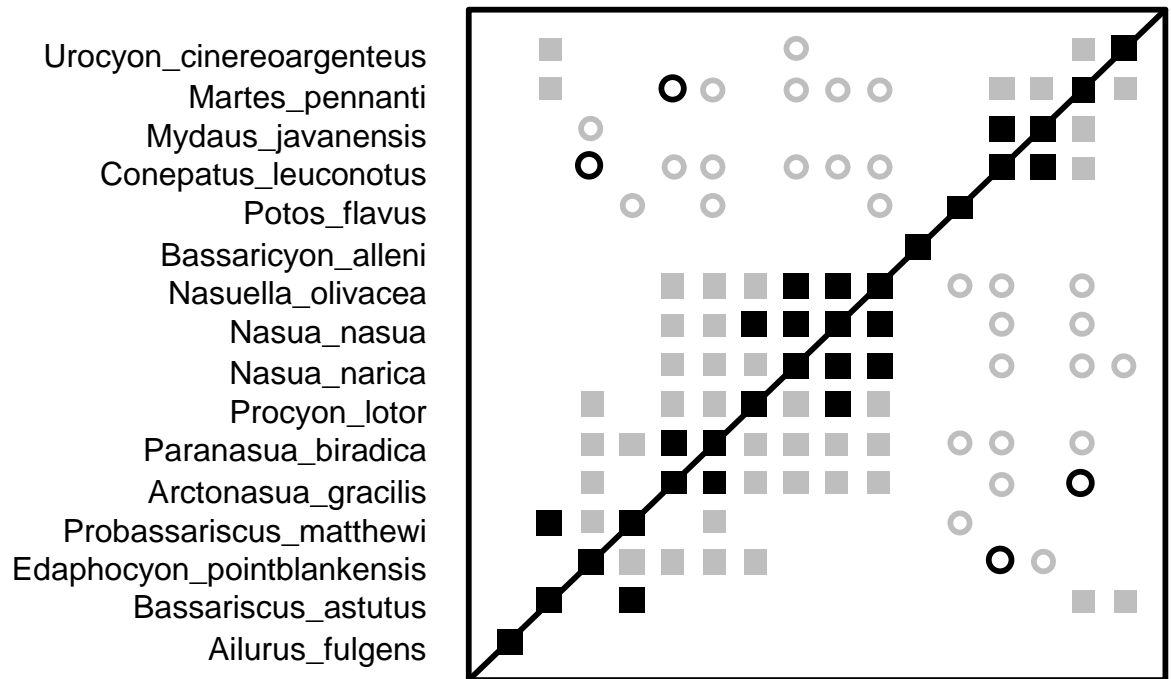
Characters: Craniodental



Order Carnivora
 Family Mephitidae

| | |
|----------------------------------|------|
| Published taxa | 21 |
| Published characters | 39 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 18 |
| Characters used for calculations | 31 |
| Median bootstrap value | 74 |
| F ₉₀ | 0.26 |
| Stress of 3D MDS | 0.09 |
| k _{min} | 4 |
| Conclusion | HB |

Notes: Mephitidae is clearly separated from the outgroup taxa in both BDC and MDS results. Mephitidae is likely a holobaramin.



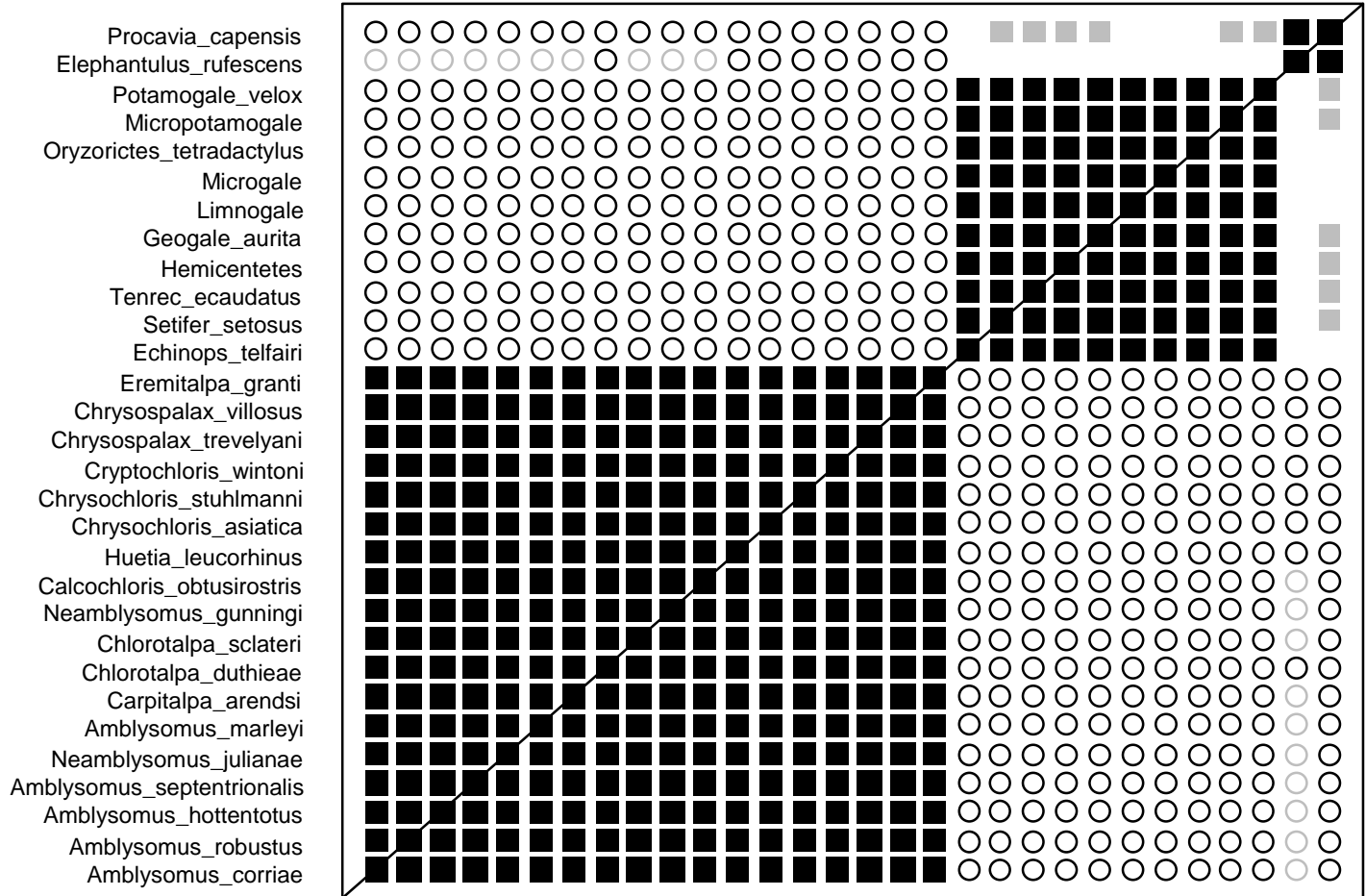
Order Carnivora
Family Procyonidae

| | |
|----------------------------------|------|
| Published taxa | 20 |
| Published characters | 78 |
| Character relevance | 0.75 |
| Taxic relevance | 0.25 |
| Taxa used for calculations | 16 |
| Characters used for calculations | 69 |
| Median bootstrap value | 79 |
| F ₉₀ | 0.32 |
| Stress of 3D MDS | 0.15 |
| k _{min} | 6 |
| Conclusion | Inc |

Notes: BDC results shows potential discontinuity around some procyonid taxa, but the MDS results show a diffuse cloud of all taxa. There is no clear evidence of discontinuity.

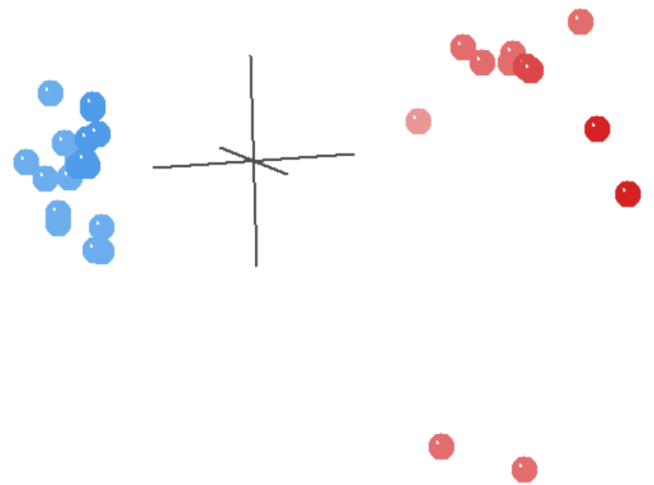
Asher, R.J., S. Maree, G. Bronner, N.C. Bennett, P. Bloomer, P. Czechowski, M. Meyer, and M. Hofreiter. 2010. A phylogenetic estimate for golden moles (Mammalia, Afrotheria, Chrysochloridae). *BMC Evolutionary Biology* 10:69.

Characters: Craniodental and postcranial



Order Chrysochloridea
 Family Chrysochloridae

| | |
|----------------------------------|------|
| Published taxa | 30 |
| Published characters | 144 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 30 |
| Characters used for calculations | 134 |
| Median bootstrap value | 100 |
| F ₉₀ | 0.93 |
| Stress of 3D MDS | 0.11 |
| k _{min} | 6 |
| Conclusion | HB |

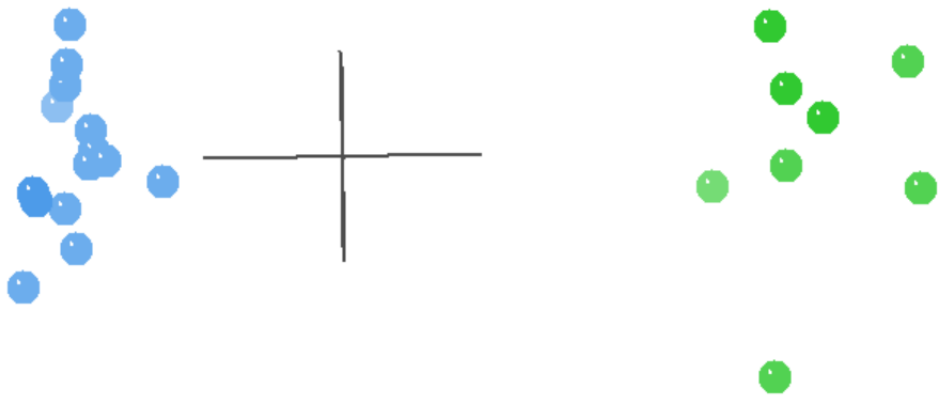
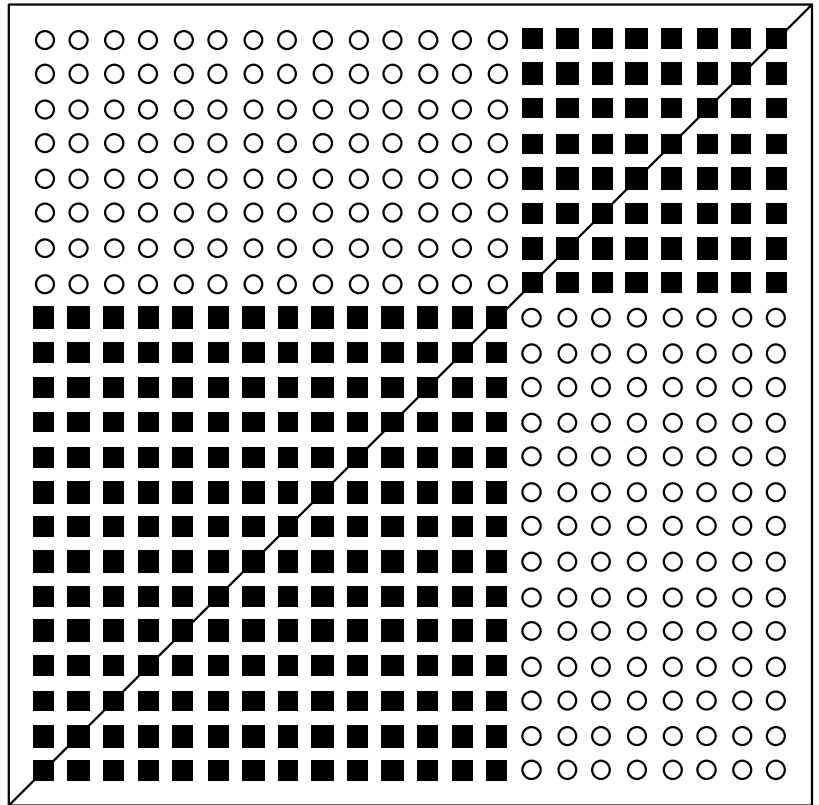


Notes: Chrysochloridae is well-separated from the outgroup taxa in MDS and BDC results. Chrysochloridae is a holobaramin.

He, K., J.-H. Chen, G.C. Gould, N. Yamaguchi, H.-S. Ai, Y.-X. Wang, Y.-P. Zhang, and X.-L. Jiang. 2012. An estimation of Erinaceidae phylogeny: a combined analysis approach. *PLoS ONE* 7(6):e39304.

Characters: Craniodental, postcranial, external morphology

Neotetracus_sinensis
 Neohylomys_hainanensis
 Hylomys_suillus
 Hylomys_parvus
 Hylomys_megalotis
 Podogymnura_truei
 Podogymnura_aureospinula
 Echinorex_gymnura
 Paraechinus_hypomelas
 Paraechinus_micropus
 Paraechinus_aethiopicus
 Hemiechinus_collaris
 Hemiechinus_auritus
 Mesechinus_hughi
 Mesechinus_dauricus
 Erinaceus_europaeus
 Erinaceus_concolor
 Erinaceus_amurensis
 Atelerix_sclateri
 Atelerix_frontalis
 Atelerix_algirus
 Atelerix_albiventris

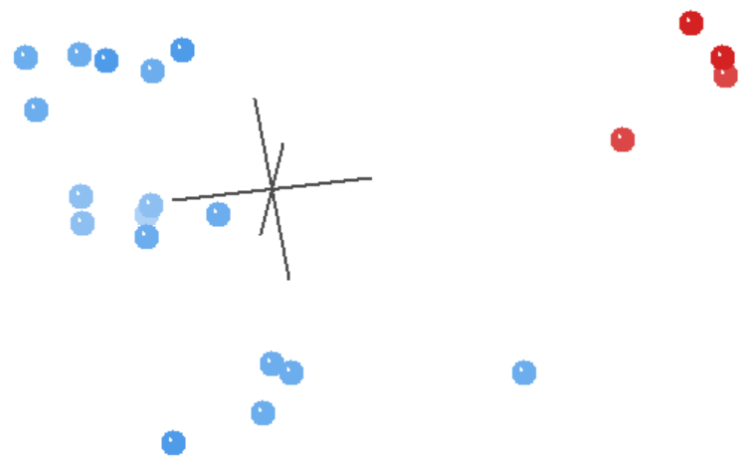
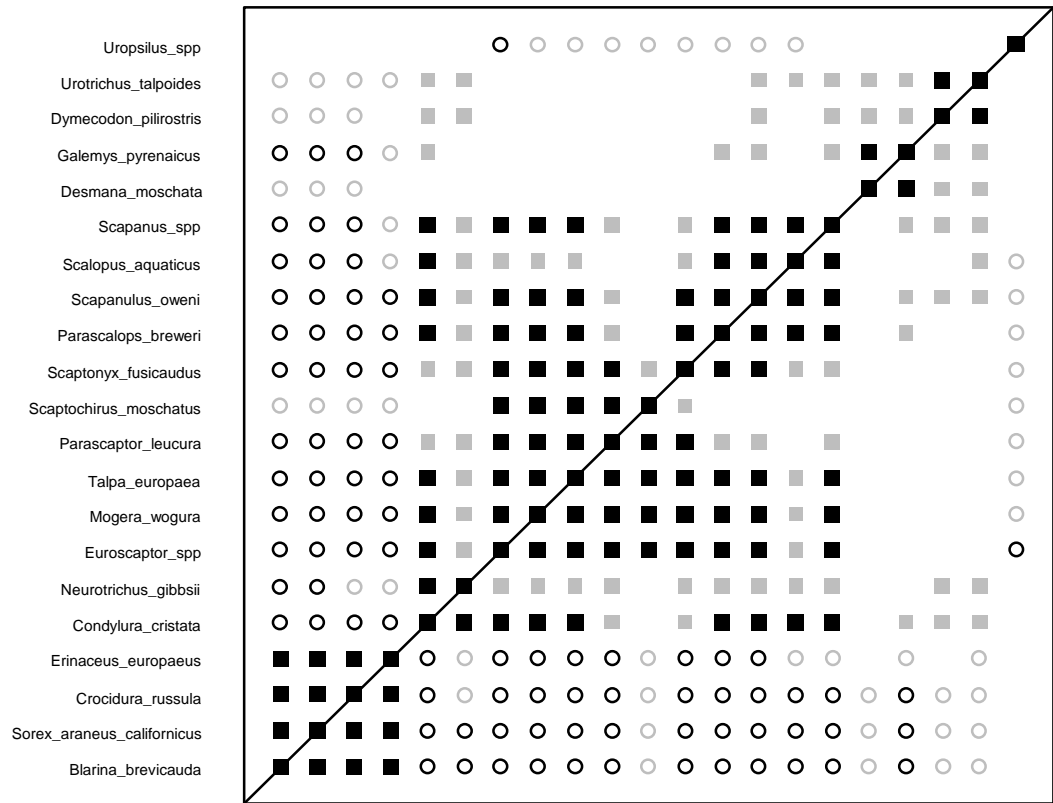


Order Erinaceomorpha
 Family Erinaceidae

| | |
|----------------------------------|------|
| Published taxa | 23 |
| Published characters | 135 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 23 |
| Characters used for calculations | 97 |
| Median bootstrap value | 100 |
| F ₉₀ | 1 |
| Stress of 3D MDS | 0.07 |
| k _{min} | 4 |
| Conclusion | HB |

Notes: Subfamilies Erinaceinae and Galericinae are both well-separated in the BDC and MDS results. Both are likely holobaramins.

Characters: Craniodental and postcranial



Order Erinaceomorpha
 Family Talpidae

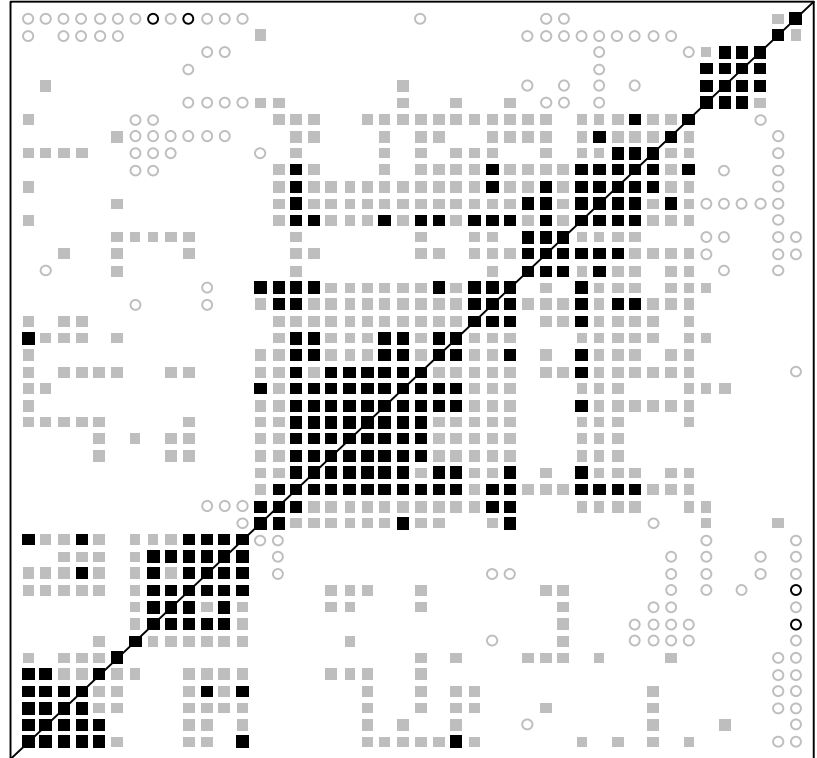
| | |
|----------------------------------|------|
| Published taxa | 21 |
| Published characters | 157 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 21 |
| Characters used for calculations | 135 |
| Median bootstrap value | 90 |
| F ₉₀ | 0.49 |
| Stress of 3D MDS | 0.21 |
| k _{min} | 9 |
| Conclusion | HB |

Notes: Talpidae is well-separated from the outgroup taxa in both the BDC and MDS results. Talpidae is likely a holobaramin.

Manz, C.L. and J.I. Bloch. 2015. Systematics and phylogeny of Paleocene-Eocene Nyctitheriidae (Mammalia, Eulipotyphla?) with description of a new species from the Late Paleocene of the Clarks Fork Basin, Wyoming, USA. *Journal of Mammalian Evolution* 22:307-342.

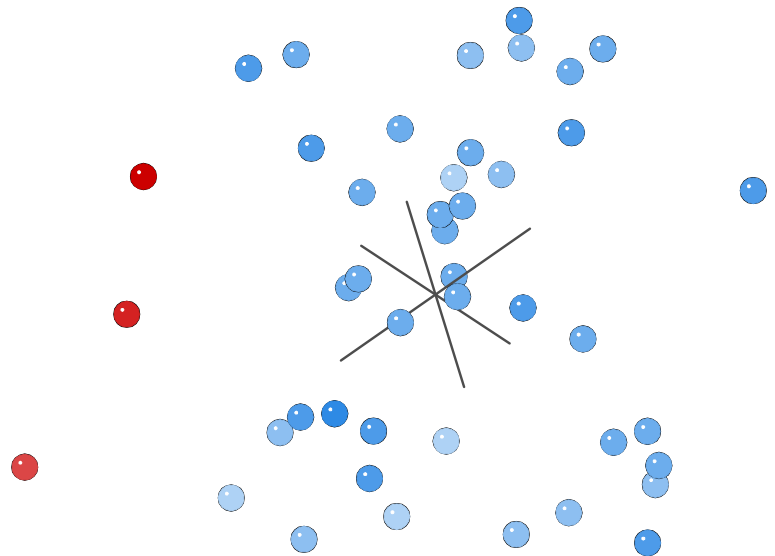
Characters: Dental

Maelestes gobiensis
Macrocranium junnei
Oedolius perexiguus
Eosoricodon terrigena
Bumbanius rarus
Asionyctia guoi
Saturninia carbonum
Scraeva hatherwoodensis
Cryptotopos woodi
Cryptotopos hartenbergi
Cryptotopos beatus
Saturninia intermedia
Saturninia gracilis
Euronyctia montana
Euronyctia grisollensis
Amphidozotherium cayluxi
Leptacodon munusculum
Saturninia grandis
Leptacodon catulus
Leptacodon packi
Leptacodon choristus
Saturninia mamertensis
Leptacodon nascimentoi
Leptacodon tener
Plagioctenodon krausae
Plagioctenodon thewisseni
Plagioctenodon rosei
Plagioctenodon dormaalensis
Leptacodon acherontus
Leptacodon donkroni
Adunator minutus
Wyonycteris galensis
Wyonycteris richardi
Wyonycteris chalik
Wyonycteris primitivus
Plagioctenoides tombowni
Plagioctenoides microlestes
Piacentidens lotus P3
Limacomyssus habrus
Ceutholestes dolosus
Nyctitherium serotinum
Nyctitherium velox
Nyctitherium krishtalkai
Acrodentis rosenorum



Order Soricomorpha
 Family Nyctitheriidae

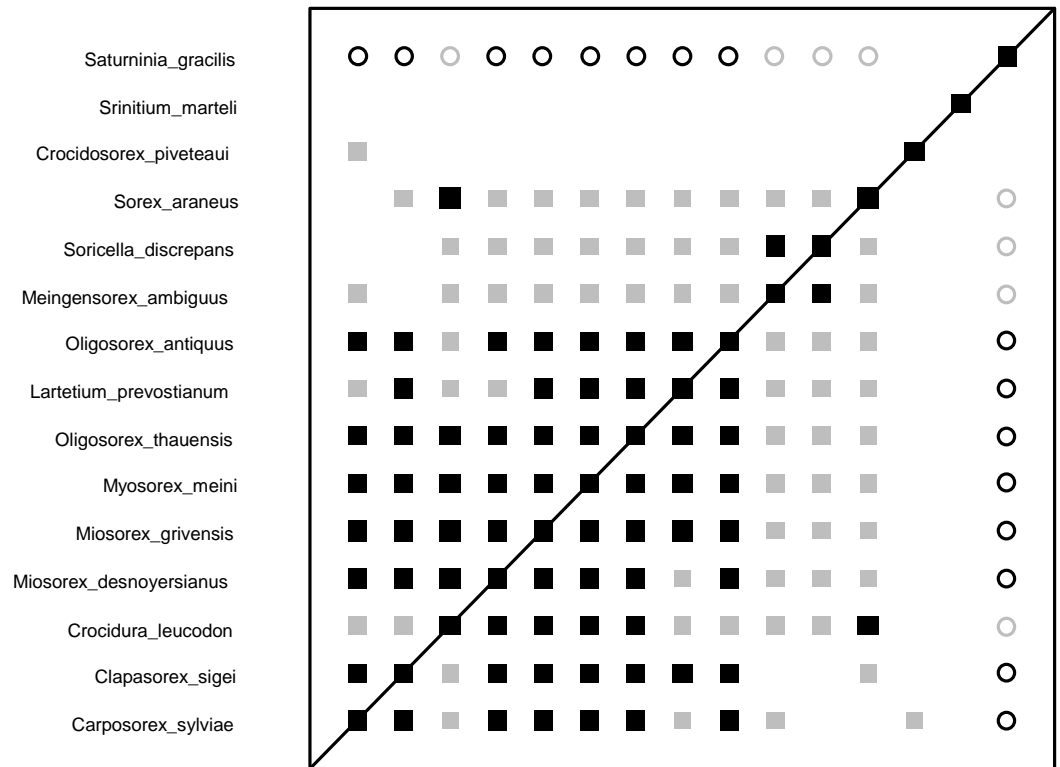
| | |
|----------------------------------|------|
| Published taxa | 54 |
| Published characters | 66 |
| Character relevance | 0.75 |
| Taxic relevance | 0.5 |
| Taxa used for calculations | 44 |
| Characters used for calculations | 34 |
| Median bootstrap value | 65 |
| F ₉₀ | 0.1 |
| Stress of 3D MDS | 0.3 |
| k _{min} | 7 |
| Conclusion | HB? |



Notes: BDC seems to support separation between the outgroup taxa and ingroup taxa, but the MDS results are far less clear. Given that the 3D stress is high (0.3), we may very tentatively accept Nyctitheriidae as a holobaramin.

Hugueney, M. and O. Maridet. 2011. Early Miocene soricids (Insectivora, Mammalia) from Limagne (Central France): new systematic comparisons, updated biostratigraphic data and evolutionary implications. *Geobios* 44:225-236.

Characters: Craniodental



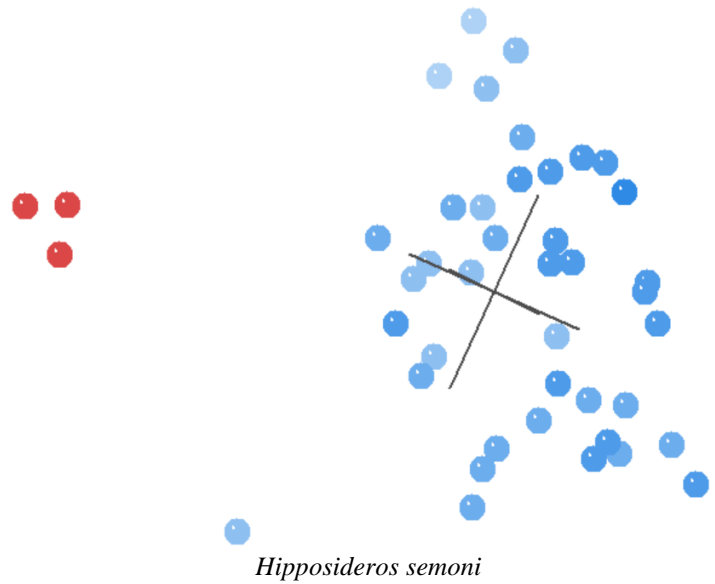
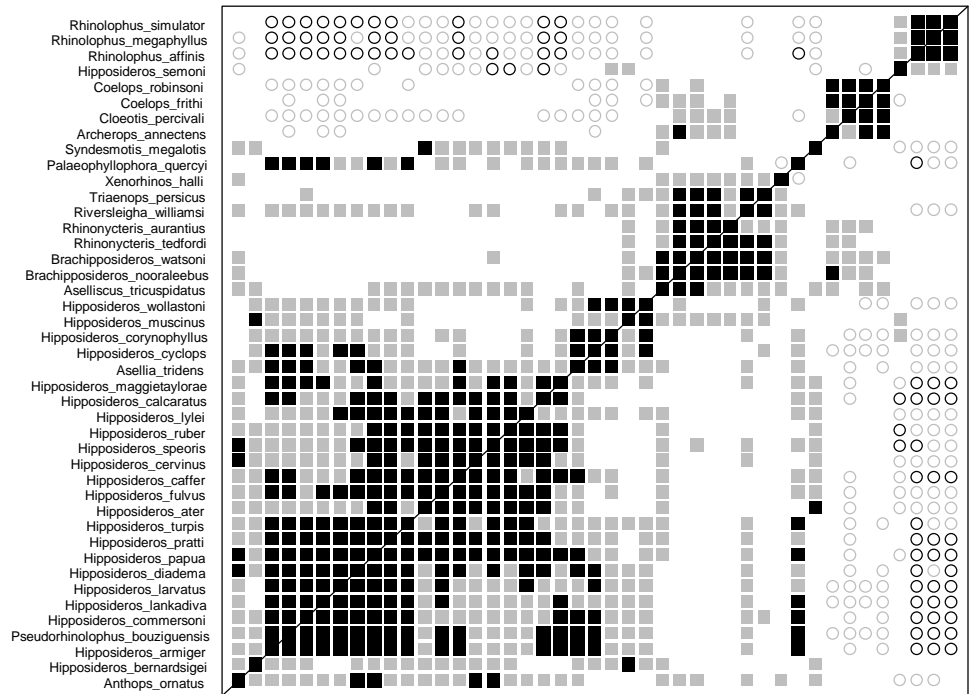
Order Soricomorpha
Family Soricidae

| | |
|----------------------------------|------|
| Published taxa | 15 |
| Published characters | 27 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 15 |
| Characters used for calculations | 21 |
| Median bootstrap value | 79 |
| F ₉₀ | 0.34 |
| Stress of 3D MDS | 0.22 |
| k _{min} | 5 |
| Conclusion | HB? |

Notes: Soricidae is well separated from the outgroup in BDC and MDS results. Soricidae is likely a holobaramin based on these characters.

Hand, S.J. and J.A.W. Kirsch. 2003. *Archerops*, a new annectant hipposiderid genus (Mammalia: Microchiroptera) from the Australian Miocene. *Journal of Paleontology* 77:1139-1151.

Characters: Craniodental and postcranial



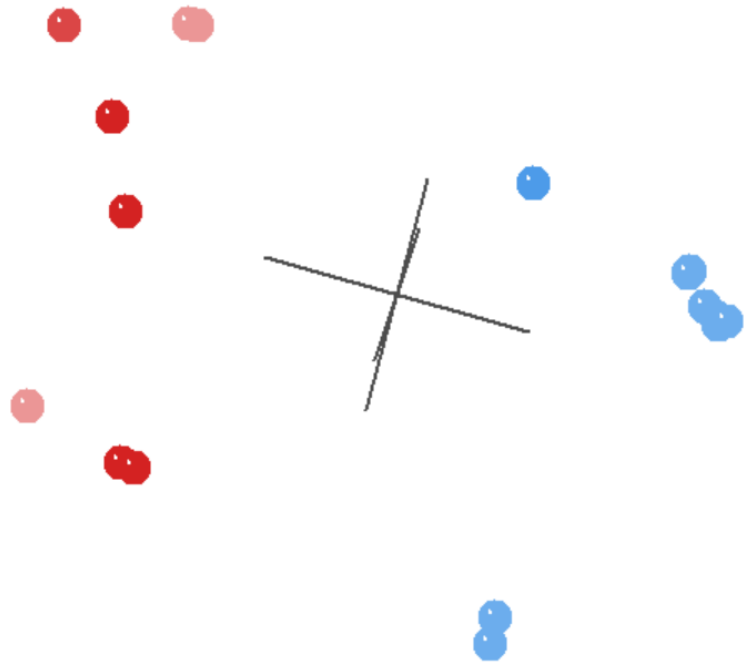
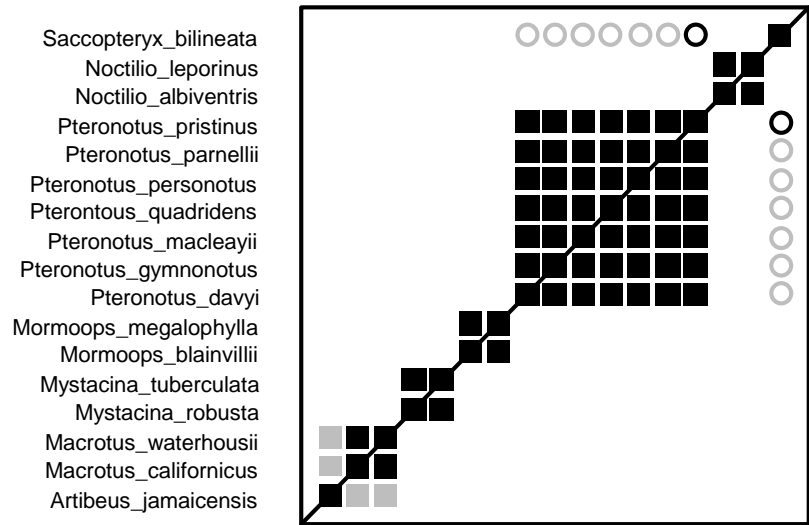
Order Chiroptera
Family Rhinolophidae

| | |
|----------------------------------|------|
| Published taxa | 44 |
| Published characters | 64 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 44 |
| Characters used for calculations | 64 |
| Median bootstrap value | 73 |
| F ₉₀ | 0.2 |
| Stress of 3D MDS | 0.32 |
| k _{min} | 11 |
| Conclusion | HB? |

Notes: BDC and MDS support a separation between Rhinolophidae and outgroup taxa. *Hipposideros semoni* shares positive BDC with all outgroup taxa and two other rhinolophids, but the MDS supports including *H. semoni* in the rhinolophid cluster. Rhinolophidae is likely a holobaramin.

Simmons, N.B. and T.M. Conway. 2001. Phylogenetic relationships of mormoopid bats (Chiroptera: Mormoopidae) based on morphological data. *Bulletin of the American Museum of Natural History* 258:1-97.

Characters: Craniodental, postcranial, soft tissue



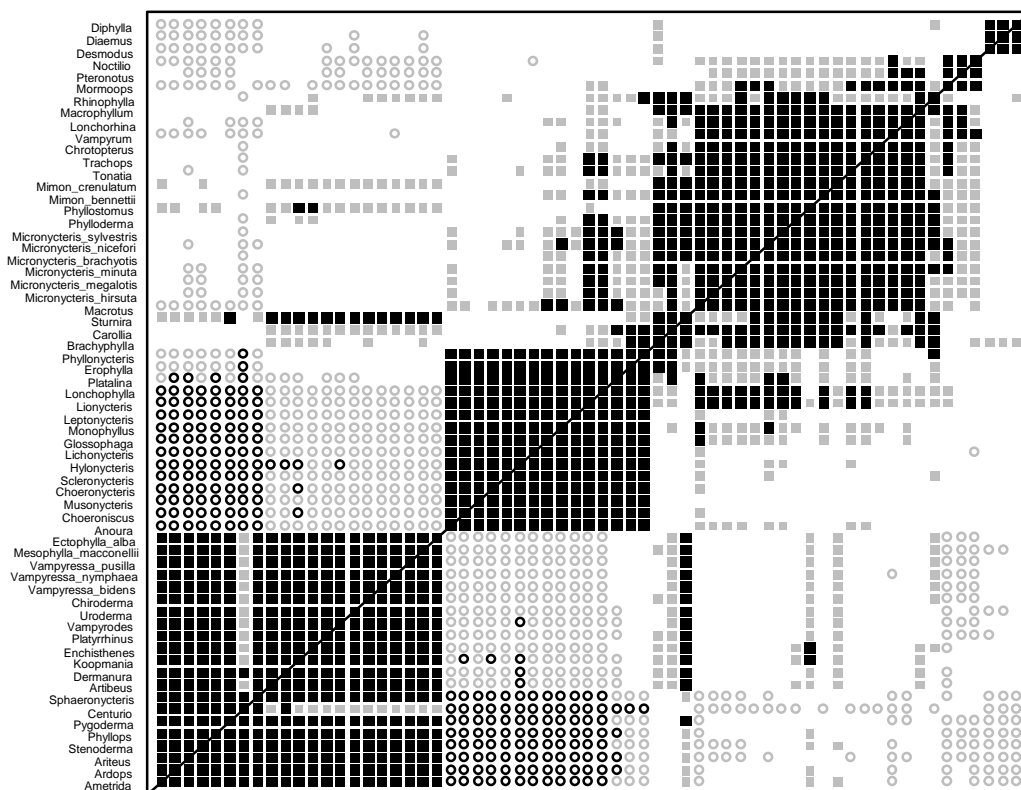
Order Chiroptera
 Family Mormoopidae

| | |
|----------------------------------|------|
| Published taxa | 17 |
| Published characters | 209 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 17 |
| Characters used for calculations | 162 |
| Median bootstrap value | 92 |
| F ₉₀ | 0.54 |
| Stress of 3D MDS | 0.19 |
| k _{min} | 7 |
| Conclusion | MB |

Notes: BDC reveals very little correlation between genera. *Pteronotus* appears to be a well-defined monobaramin in the BDC results. MDS reveals a distinction between ingroup and outgroup taxa, but both are diffuse and poorly clustered. We may provisionally accept *Pteronotus* as a monobaramin. Clear discontinuity is not evident.

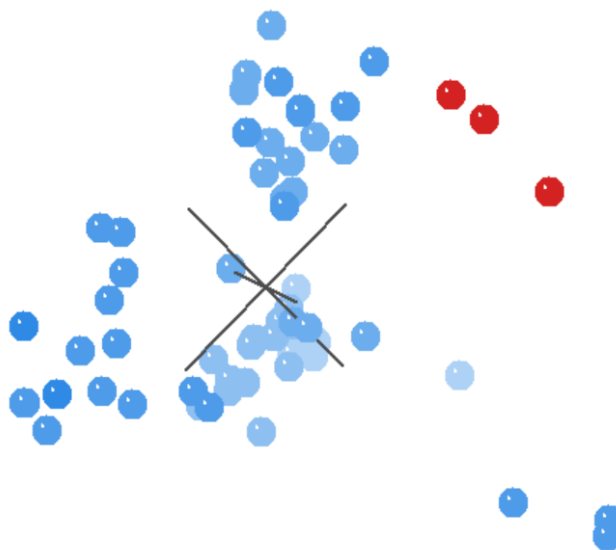
Wetterer, A.L., M.V. Rockman, and N.B. Simmons. 2000. Phylogeny of phyllostomid bats (Mammalia: Chiroptera): data from diverse morphological systems, sex chromosomes, and restriction sites. *Bulletin of the American Museum of Natural History* 248:1-200.

Characters: Craniodental, postcranial, soft issue, karyotype, restriction sites



Order Chiroptera
Family Phyllostomidae

| | |
|----------------------------------|------|
| Published taxa | 63 |
| Published characters | 150 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 63 |
| Characters used for calculations | 98 |
| Median bootstrap value | 77 |
| F ₉₀ | 0.33 |
| Stress of 3D MDS | 0.24 |
| k _{min} | 10 |
| Conclusion | Inc |



Notes: No clear evidence of discontinuity in BDC or MDS results.

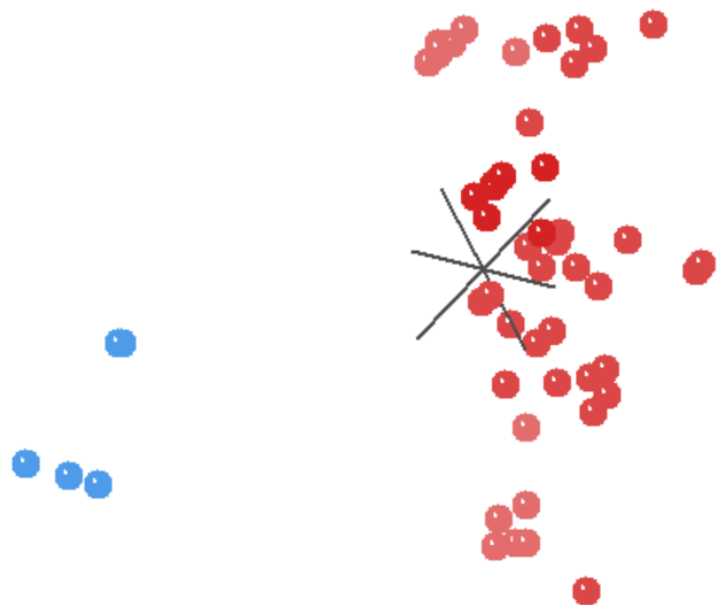
Burger, B.J. 2013. A new species of the archaic primate *Zanycteris* from the late Paleocene of western Colorado and the phylogenetic position of the family Picrodontidae. *PeerJ* 1:e191.

Characters: Dental



Order Primates
Family Picrodontidae

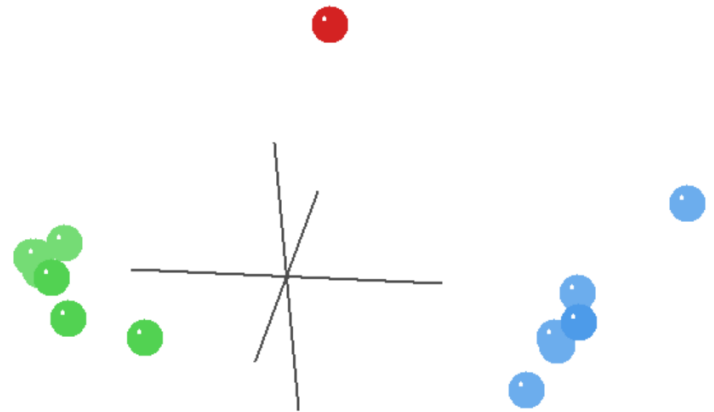
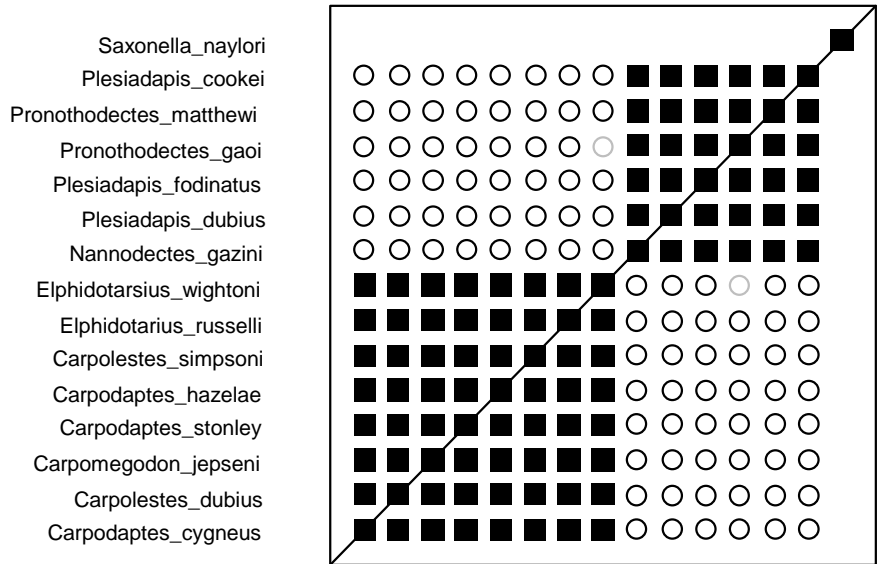
| | |
|----------------------------------|------|
| Published taxa | 58 |
| Published characters | 113 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 58 |
| Characters used for calculations | 98 |
| Median bootstrap value | 79 |
| F ₉₀ | 0.34 |
| Stress of 3D MDS | 0.24 |
| k _{min} | 7 |
| Conclusion | HB |



Notes: The taxa are trimmed to include Picrodontidae, with Plesiadapidae, and Carpolestidae as the outgroup. Picrodontidae is well-separated from the outgroup taxa in both BDC and MDS results. Picrodontidae is probably a holobaramin.

Burger, B.J. 2013. A new species of the archaic primate *Zanycteris* from the late Paleocene of western Colorado and the phylogenetic position of the family Picrodontidae. *PeerJ* 1:e191.

Characters: Dental



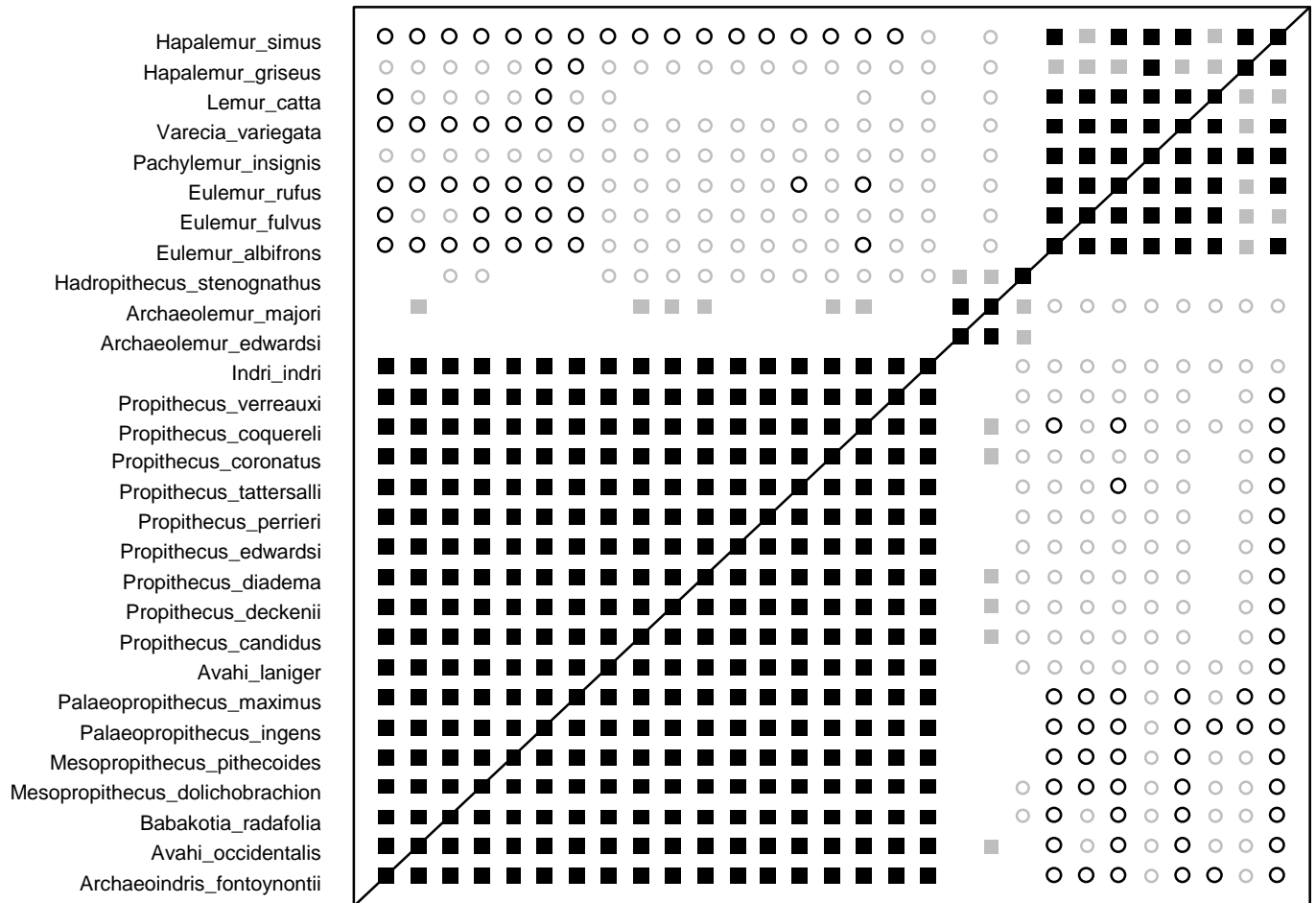
Order Primates
Family Plesiadapidae

| | |
|----------------------------------|------|
| Published taxa | 58 |
| Published characters | 113 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 15 |
| Characters used for calculations | 110 |
| Median bootstrap value | 100 |
| F ₉₀ | 0.86 |
| Stress of 3D MDS | 0.09 |
| k _{min} | 4 |
| Conclusion | HB |

Green: Carpolestidae
Blue: Plesiadapidae
Red: *Saxonella*

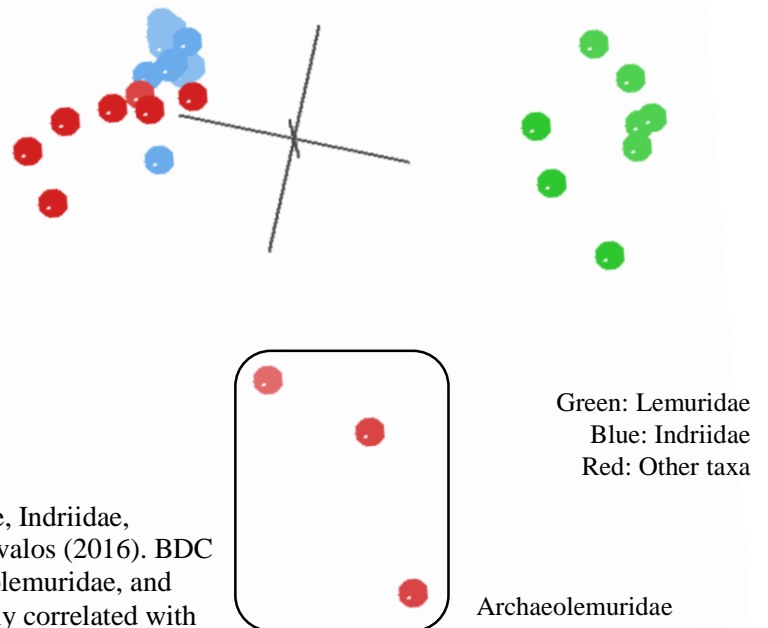
Notes: Taxa from the previous analysis are further subdivided to remove the picrodontids. Plesiadapidae and Carpolestidae are well separated from each other and from the outgroup *Saxonella* in BDC and MDS results. Both are likely holobaramins.

Characters: Craniodental and postcranial



Order Primates
Family Lemuridae

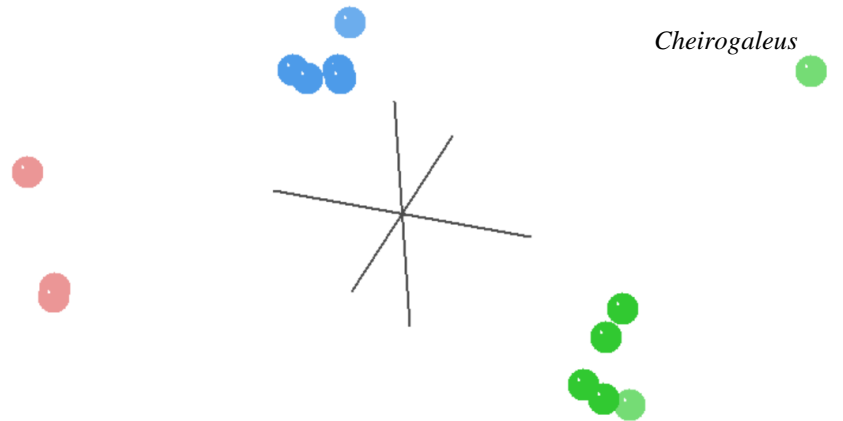
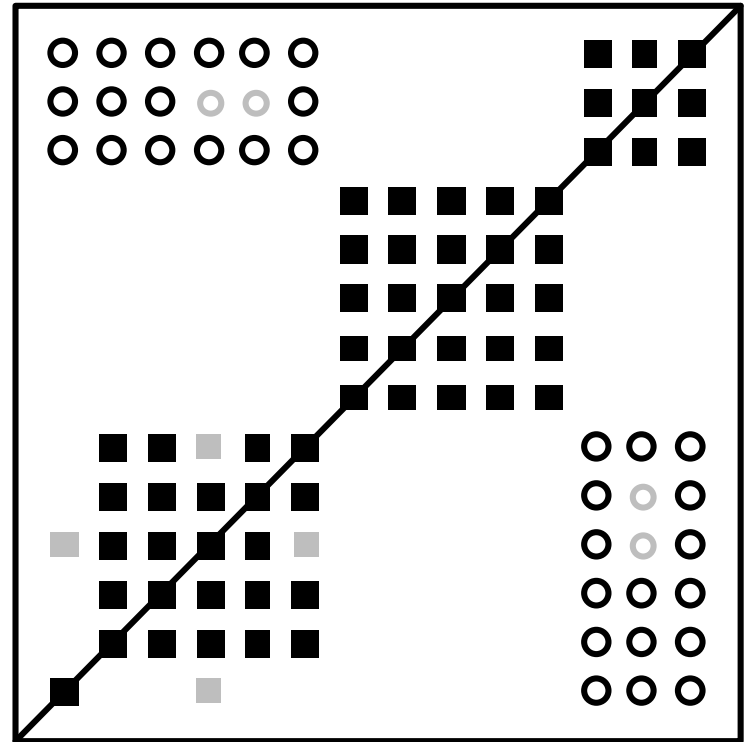
| | |
|----------------------------------|------|
| Published taxa | 29 |
| Published characters | 421 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 29 |
| Characters used for calculations | 157 |
| Median bootstrap value | 98 |
| F ₉₀ | 0.67 |
| Stress of 3D MDS | 0.23 |
| k _{min} | 11 |
| Conclusion | HB |



Notes: Taxon sample has been trimmed here to include Lemuridae, Indriidae, Palaeopropithecidae, and Archaeolemuridae from Herrera and Dávalos (2016). BDC reveals three groups of taxa corresponding to Lemuridae, Archaeolemuridae, and Indriidae + Palaeopropithecidae. *Archaeolemur majori* is positively correlated with six members of the Indriidae + Palaeopropithecidae group, but all but one BDC with *Archaeolemur majori* have poor bootstrap values. All comparisons between the groups Lemuridae and Indriidae + Palaeopropithecidae share significant, negative BDC. The MDS results support the three groups observed in the BDC, but *Archaeolemur majori* is clearly separated from the Indriidae + Palaeopropithecidae. Hence we may conclude that Lemuridae and Indriidae + Palaeopropithecidae are both likely holobaramins.

Craniodental and postcranial

Megaladapis_madagascariensis
 Megaladapis_grandidieri
 Megaladapis_edwardsi
 Lepilemur_mustelinus
 Lepilemur_edwardsi
 Lepilemur_ruficaudatus
 Lepilemur_leucopus
 Lepilemur_dorsalis
 Phaner_furcifer
 Mirza_coquereli
 Microcebus_murinus
 Microcebus_rufus
 Microcebus_griseorufus
 Cheirogaleus_major



Order Primates
 Family Lepilemuridae

| | |
|----------------------------------|-------|
| Published taxa | 14 |
| Published characters | 421 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 14 |
| Characters used for calculations | 161 |
| Median bootstrap value | 93 |
| F ₉₀ | 0.575 |
| Stress of 3D MDS | 0.16 |
| k _{min} | 7 |
| Conclusion | HB |

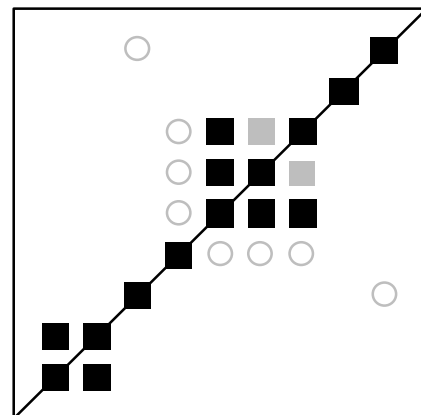
Red: Megaladapidae
 Green: Cheirogaleidae
 Blue: Lepilemuridae

Notes: Taxa have been trimmed to include Lepilemuridae, Cheirogaleidae, and Megaladapidae. BDC reveals three clear groups corresponding to the three families. MDS confirms these three groups. The position of *Cheirogaleus major* is uncertain in both BDC and MDS results. Lepilemuridae and Cheirogaleidae (excluding *Cheirogaleus major*) are likely holobaramins.

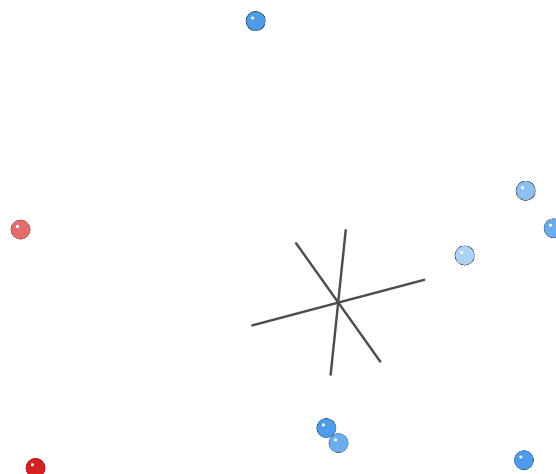
Masters, J.C., N.M. Anthony, M.J. De Wit, and A. Mitchell. 2005. Reconstructing the evolutionary history of the Lorisidae using morphological, molecular, and geological data. *American Journal of Physical Anthropology* 127:465-480.

Characters: Craniodental

Otolemur crassicaudatus
Microcebus murinus
Perodicticus potto
Nycticebus pygmaeus
Nycticebus coucang
Galagoides demidoff
Loris tardigradus
Arctocebus calabarensis
Arctocebus aureus



Microcebus murinus



Order Primates
 Family Loridae

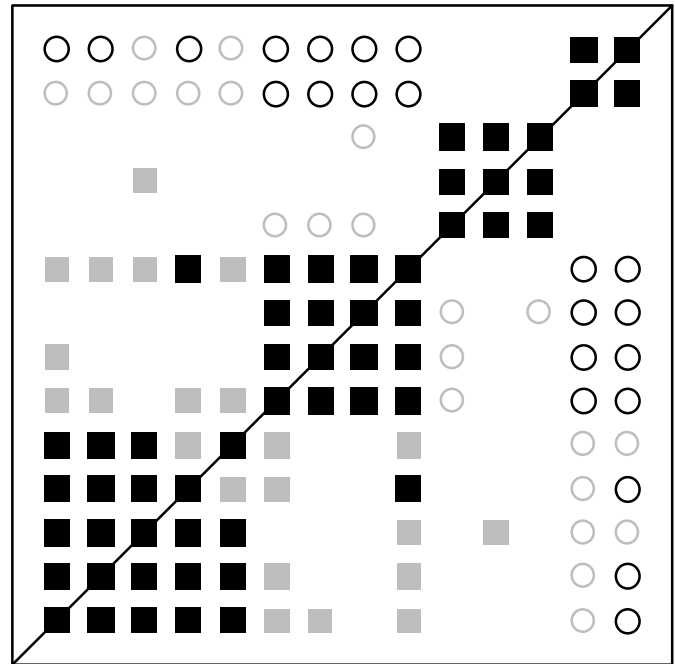
| | |
|----------------------------------|------|
| Published taxa | 9 |
| Published characters | 36 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 9 |
| Characters used for calculations | 36 |
| Median bootstrap value | 95.5 |
| F ₉₀ | 0.64 |
| Stress of 3D MDS | 0.07 |
| k _{min} | 4 |
| Conclusion | Inc |

Notes: BDC has few correlations, and MDS results show a diffuse cluster of taxa. No clear evidence of discontinuity is present.

Bloch, J.I., D.C. Fisher, K.D. Rose, and P.D. Gingerich. 2001. Stratocladistic analysis of Paleocene Carpolestidae (Mammalia, Plesiadapiformes) with description of a new late Tiffanian genus. *Journal of Vertebrate Paleontology* 21:119-131.

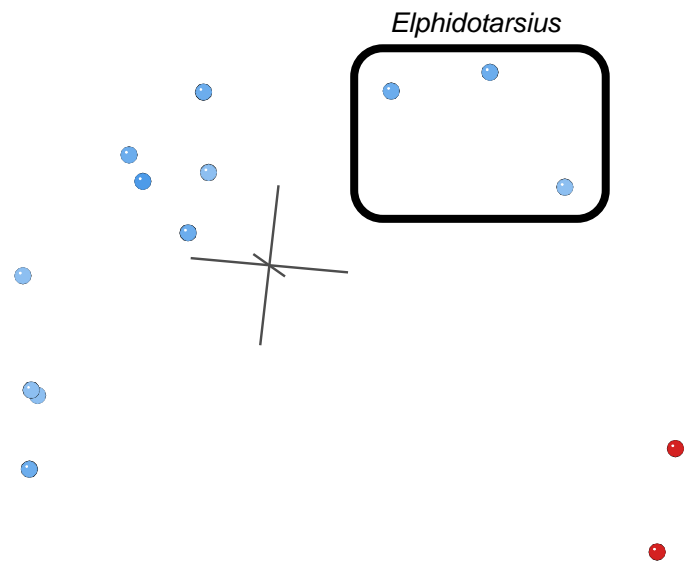
Characters: Dental

Purgatorius
Pronothodectes
Elphidotarsius wightoni
Elphidotarsius shotgunensis
Elphidotarsius florencae
Carpomegodon jepseni
Carpolestes simpsoni
Carpolestes nigridentis
Carpolestes dubius
Carpocristes oriens
Carpodaptes aulacodon
Carpodaptes hazelae
Carpodaptes cygneus
Carpocristes hohackensis

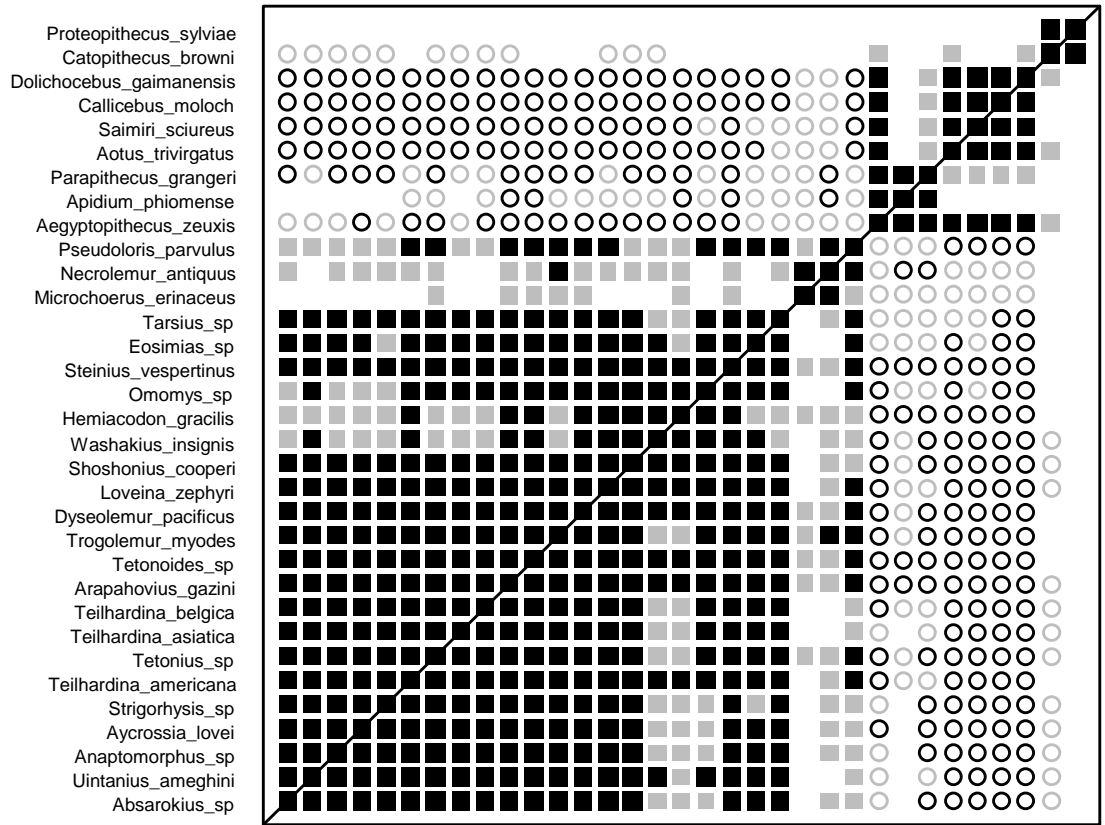


Order Primates
 Family Carpolestidae

| | |
|----------------------------------|------|
| Published taxa | 15 |
| Published characters | 33 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 14 |
| Characters used for calculations | 30 |
| Median bootstrap value | 88 |
| F ₉₀ | 0.46 |
| Stress of 3D MDS | 0.09 |
| k _{min} | 4 |
| Conclusion | HB? |

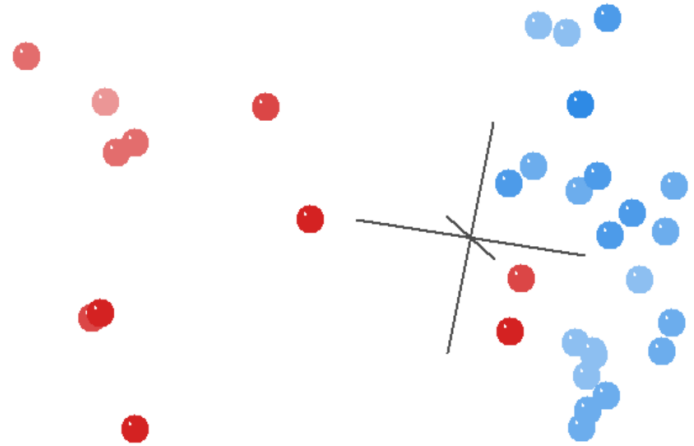


BDC reveals three groups: outgroup taxa, genus *Elphidotarsius*, and the remaining carpolestids. Significant, negative BDC occurs between the outgroup and the remaining carpolestids but not between the outgroup and *Elphidotarsius*. There is very little significant BDC between *Elphidotarsius* and the remaining carpolestids, and none have bootstrap values >90%. MDS results reveal a good separation between the outgroup taxa and the Carpolestidae, including *Elphidotarsius*, which is part of an arc of carpolestid taxa. Taken together, MDS and BDC support recognizing Carpolestidae as a provisional holobaramin.



Order Primates
 Family Omomyidae

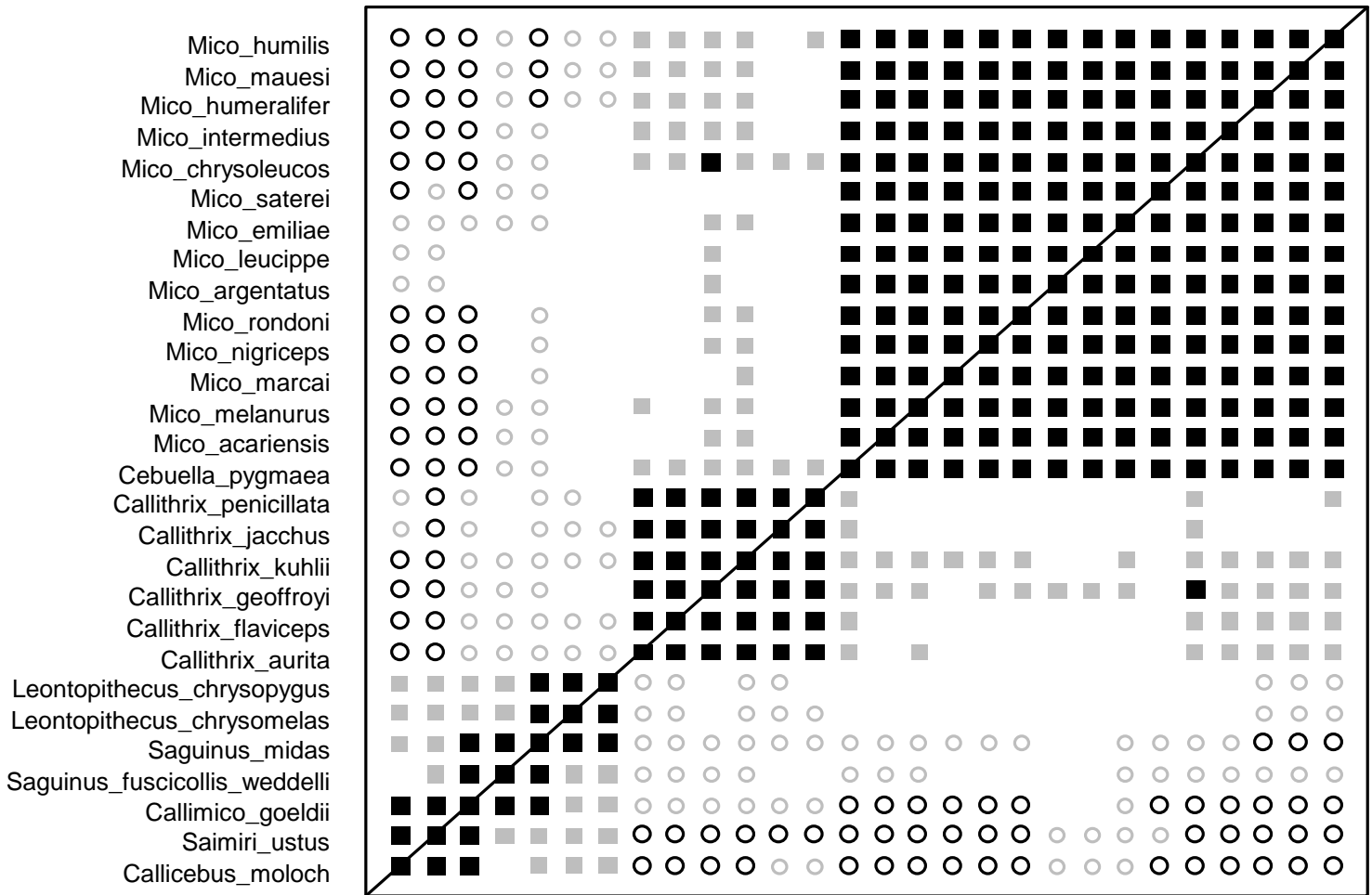
| | |
|----------------------------------|------|
| Published taxa | 52 |
| Published characters | 303 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 33 |
| Characters used for calculations | 152 |
| Median bootstrap value | 97 |
| F ₉₀ | 0.66 |
| Stress of 3D MDS | 0.27 |
| k _{min} | 12 |
| Conclusion | HB |



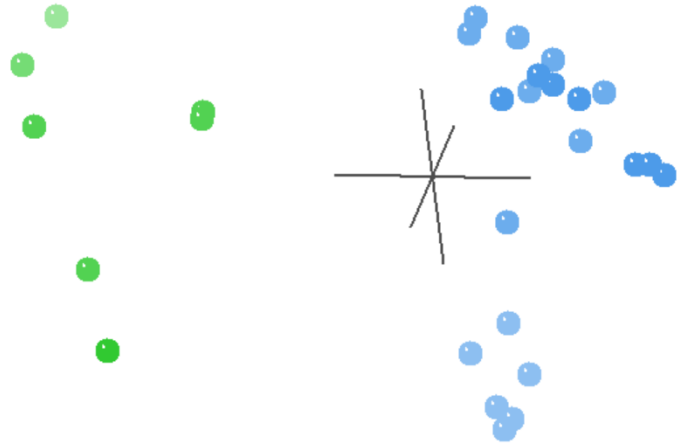
Notes: Taxa have been reduced to the Omomyidae and their sister clade, the anthropoids, from Ni et al.'s (2004) phylogeny. Both BDC and MDS reveal a clear distinction between anthropoids and Omomyidae. *Eosimias* and *Tarsius* cluster with Omomyidae in both BDC and MDS results. Omomyidae + *Eosimias* + *Tarsius* is likely a holobaramin.

Garbino, G.S. 2015. How many marmoset (Primates: Cebidae: Callitrichinae) genera are there? A phylogenetic analysis based on multiple morphological systems. *Cladistics* 31:652-678.

Characters: Craniodental, postcranial, soft tissue, vocal



Blue: Callitrichinae
Green: other Cebidae

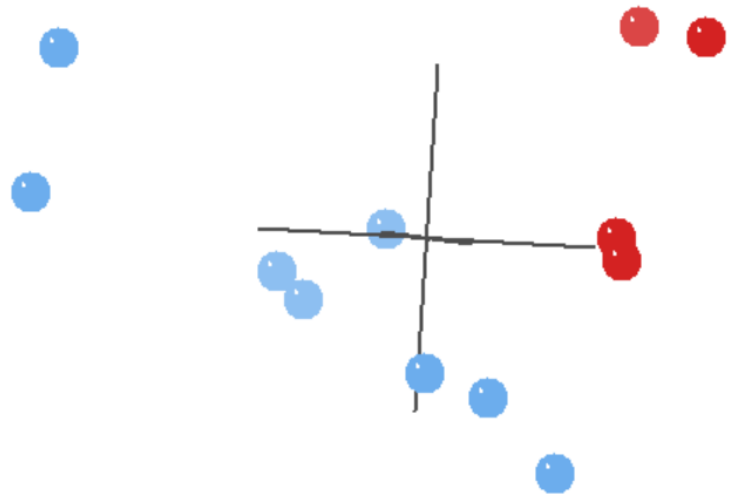
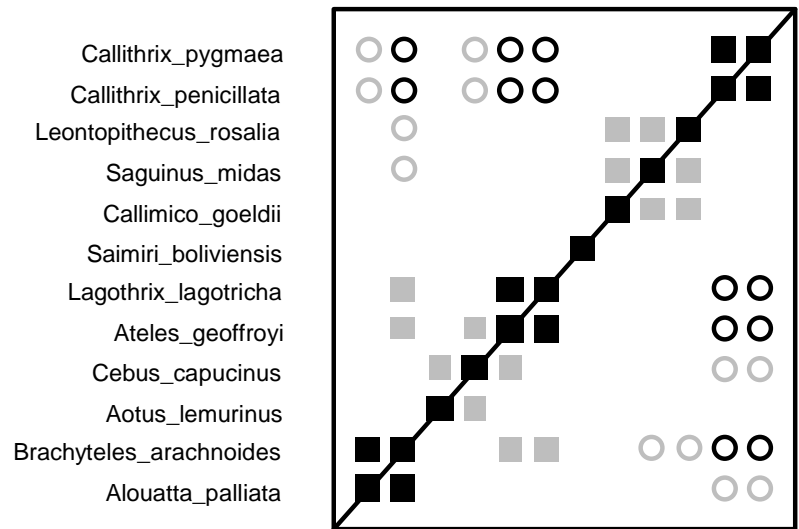


Order Primates
Family Cebidae

| | |
|----------------------------------|------|
| Published taxa | 28 |
| Published characters | 83 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 28 |
| Characters used for calculations | 76 |
| Median bootstrap value | 83.5 |
| F ₉₀ | 0.46 |
| Stress of 3D MDS | 0.14 |
| k _{min} | 7 |
| Conclusion | HB? |

Notes: BDC and MDS reveal two clear groups: Callitrichinae and the rest of the cebids. This suggests that Callitrichinae is a holobaramin.

Characters: Craniodental

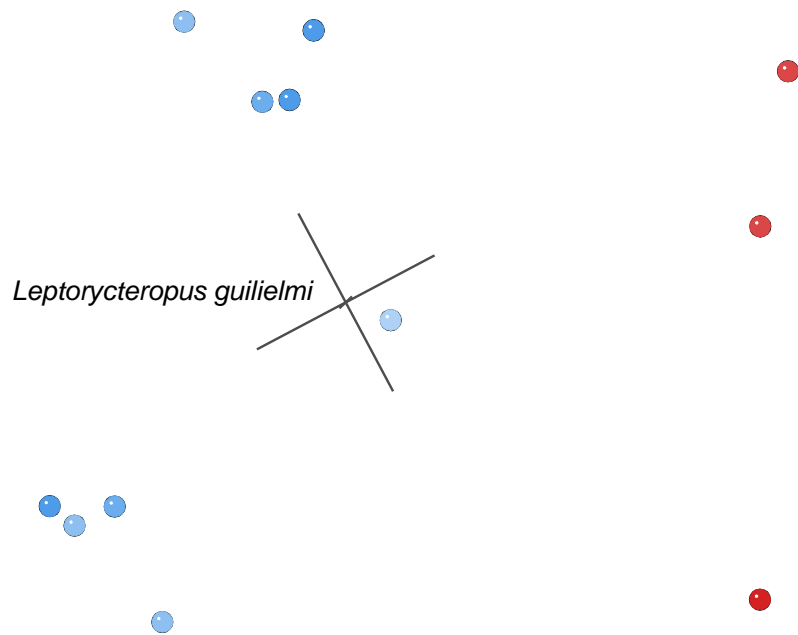
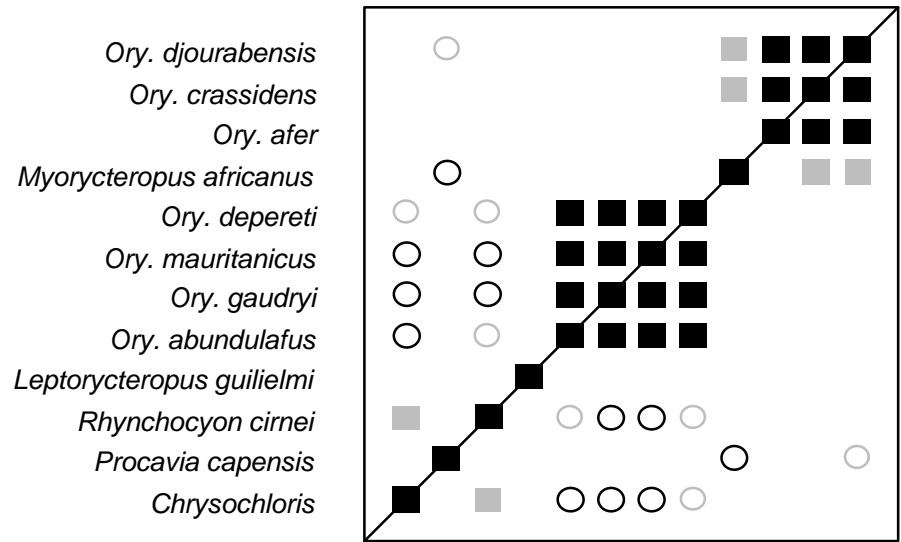


Order Primates
Family Cebidae

| | |
|----------------------------------|------|
| Published taxa | 31 |
| Published characters | 268 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 12 |
| Characters used for calculations | 238 |
| Median bootstrap value | 97 |
| F ₉₀ | 0.61 |
| Stress of 3D MDS | 0.19 |
| k _{min} | 9 |
| Conclusion | Inc |

Notes: BDC results are poor with few significant correlations. MDS reveals a diffuse cluster of taxa. There is no clear evidence of discontinuity.

Characters: Craniodental and postcranial

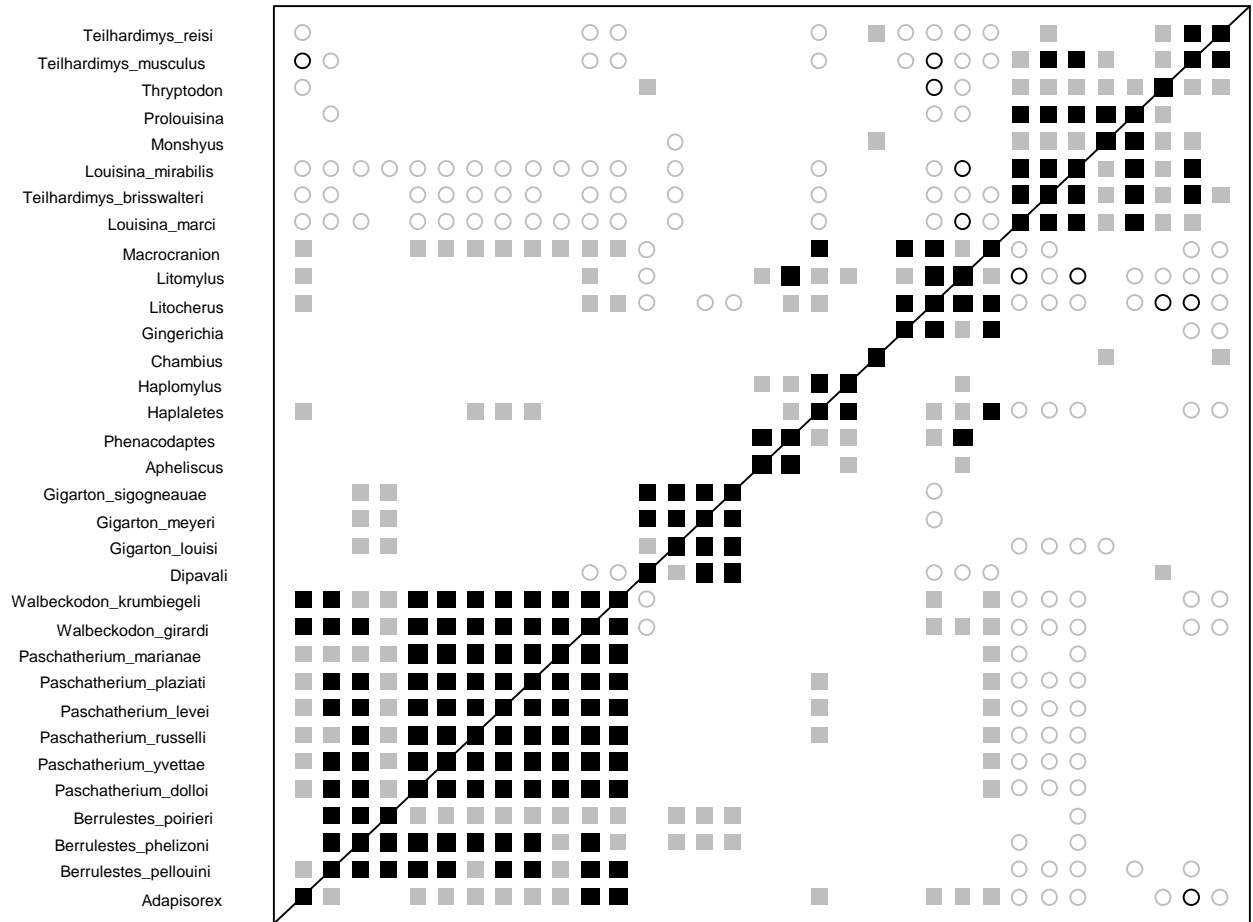


Order Tubulidentata
Family Orycteropodidae

| | |
|----------------------------------|------|
| Published taxa | 12 |
| Published characters | 39 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 12 |
| Characters used for calculations | 32 |
| Median bootstrap value | 89.5 |
| F ₉₀ | 0.47 |
| Stress of 3D MDS | 0.13 |
| k _{min} | 3 |
| Conclusion | Inc |

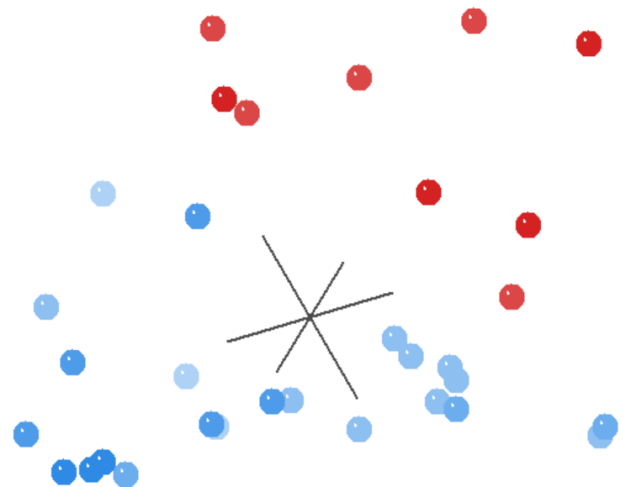
Notes: BDC results reveal two groups of orycteropids with sporadic negative BDC with outgroup taxa. MDS reveals a highly diffuse group of orycteropids that are separated from the outgroup. There is little conclusive evidence of discontinuity or continuity among the orycteropids.

Characters: Craniodental and postcranial



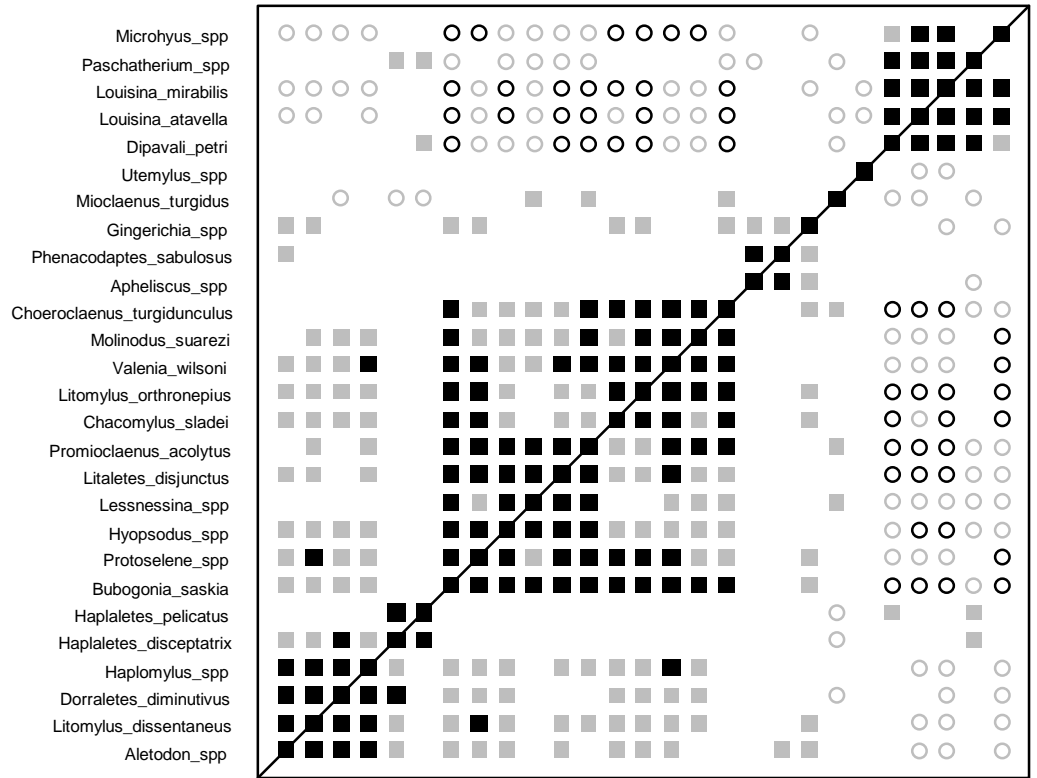
Order Condylarthra
Family Louisinidae

| | |
|----------------------------------|------|
| Published taxa | 37 |
| Published characters | 89 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 37 |
| Characters used for calculations | 67 |
| Median bootstrap value | 79 |
| F_{90} | 0.19 |
| Stress of 3D MDS | 0.14 |
| k_{min} | 5 |
| Conclusion | Inc |

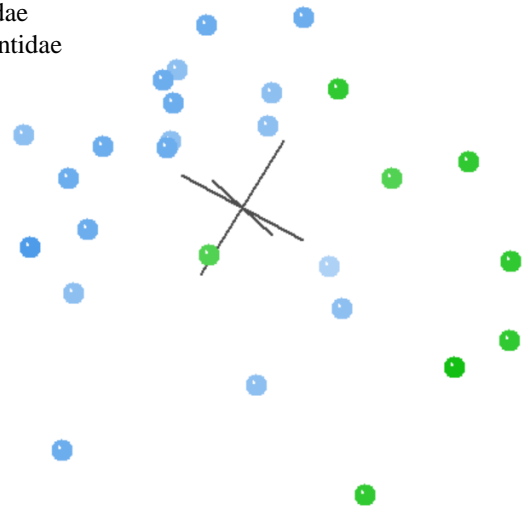


Notes: BDC implies discontinuity within Louisinidae, but no clustering evident in MDS

Characters: Dental and postcranial



Green: Louisinidae
Blue: Hyopsodontidae



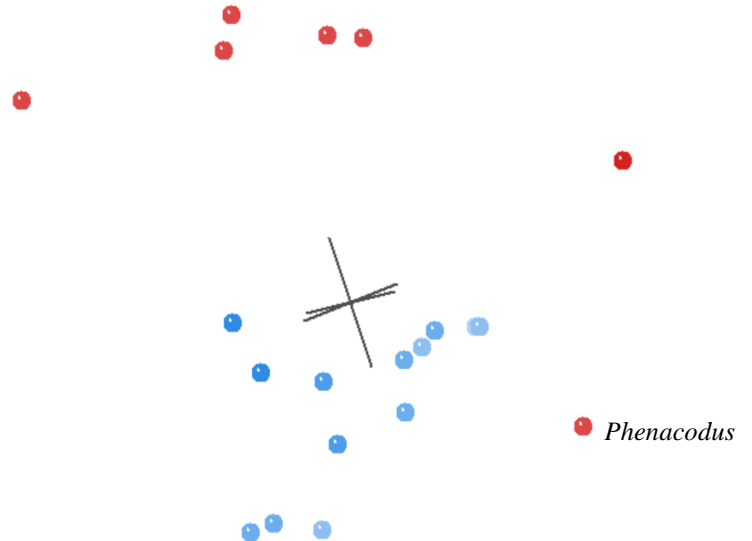
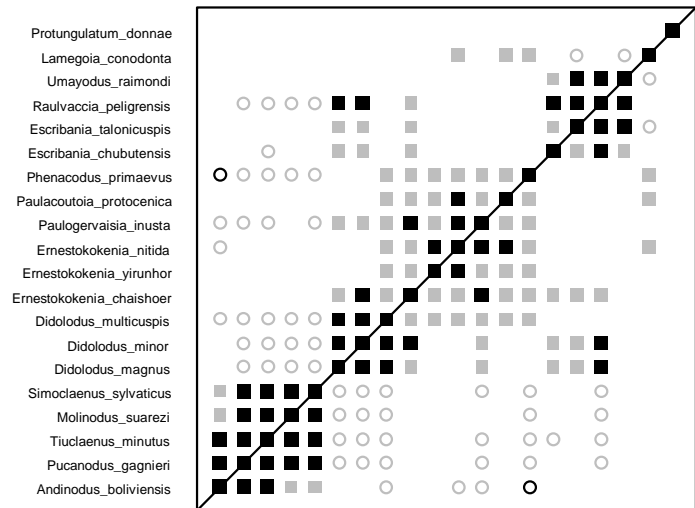
Order Condylarthra
Family Hyopsodontidae

| | |
|----------------------------------|------|
| Published taxa | 30 |
| Published characters | 59 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 27 |
| Characters used for calculations | 44 |
| Median bootstrap value | 78 |
| F ₉₀ | 0.22 |
| Stress of 3D MDS | 0.22 |
| k _{min} | 8 |
| Conclusion | Inc |

Notes: BDC reveals two groups of taxa, but MDS shows a diffuse cluster. Clear discontinuity is not evident.

Gelfo, J.N. and B. Sigé. 2011. A new didolodontid mammal from the Late Paleocene-Earliest Eocene of Laguna Umayo, Peru. *Acta Palaeontologica Polonica* 56:665-678.

Characters: Dental

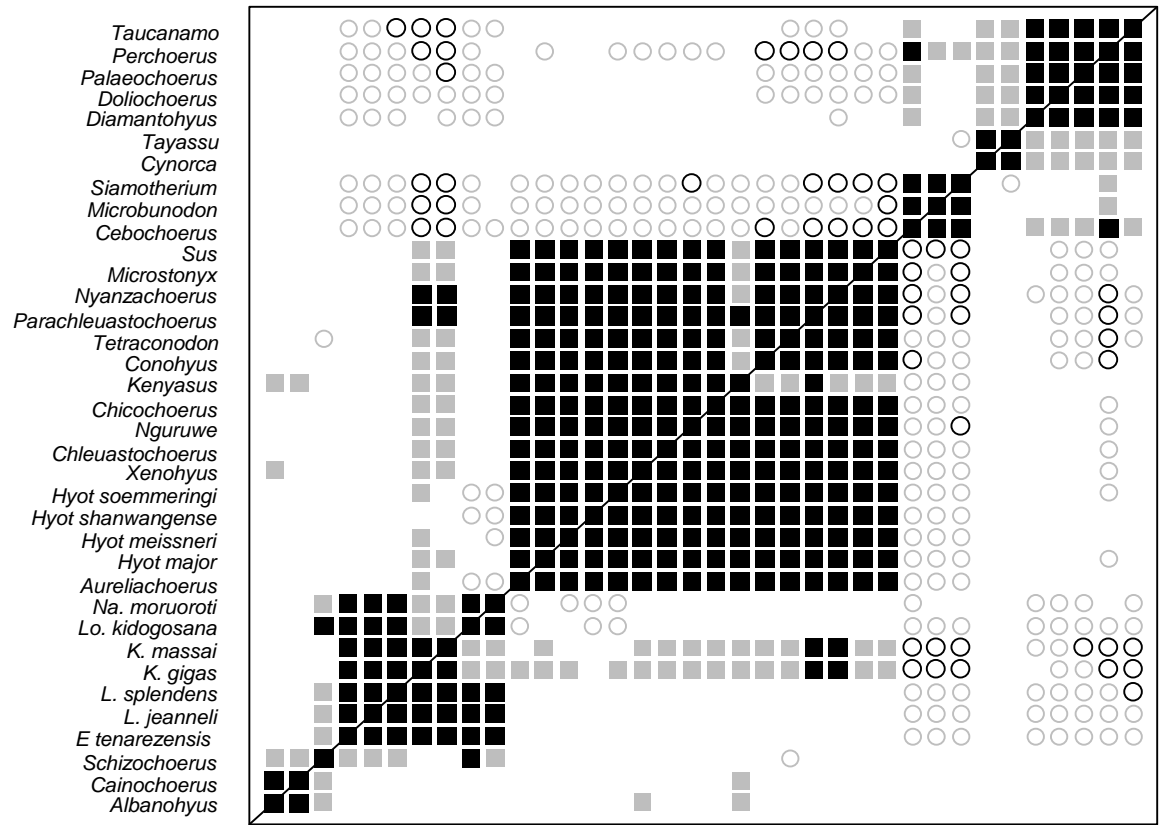


Order Condylarthra
Family Didolodontidae

| | |
|----------------------------------|------|
| Published taxa | 20 |
| Published characters | 41 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 20 |
| Characters used for calculations | 25 |
| Median bootstrap value | 74 |
| F ₉₀ | 0.13 |
| Stress of 3D MDS | 0.2 |
| k _{min} | 5 |
| Conclusion | HB? |

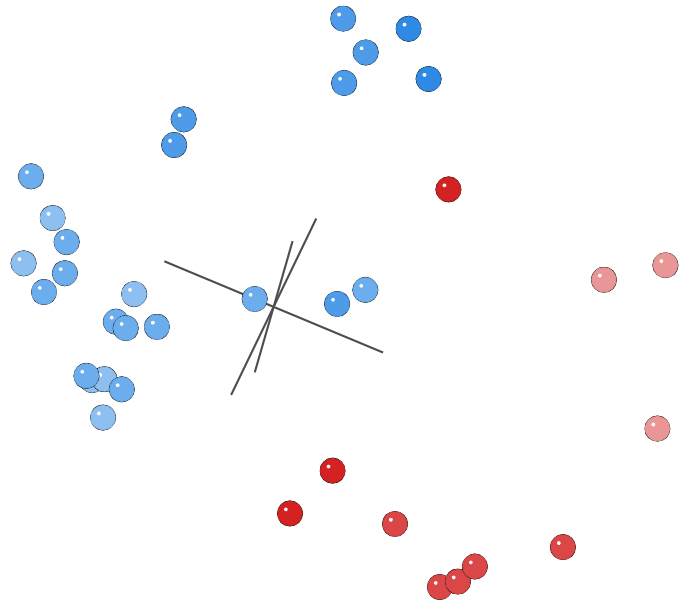
Notes: Didolodontidae is well separated from the outgroup in both BDC and MDS results. *Phenacodus* is also part of the Dipolodontidae in both analyses. Didolodontidae + *Phenacodus* is likely a holobaramin.

Characters: Dental



Order Artiodactyla
Family Suidae

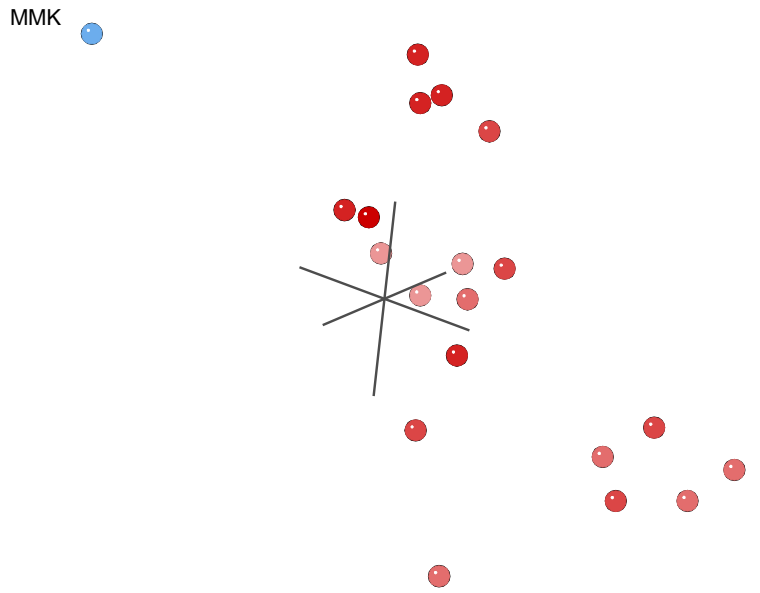
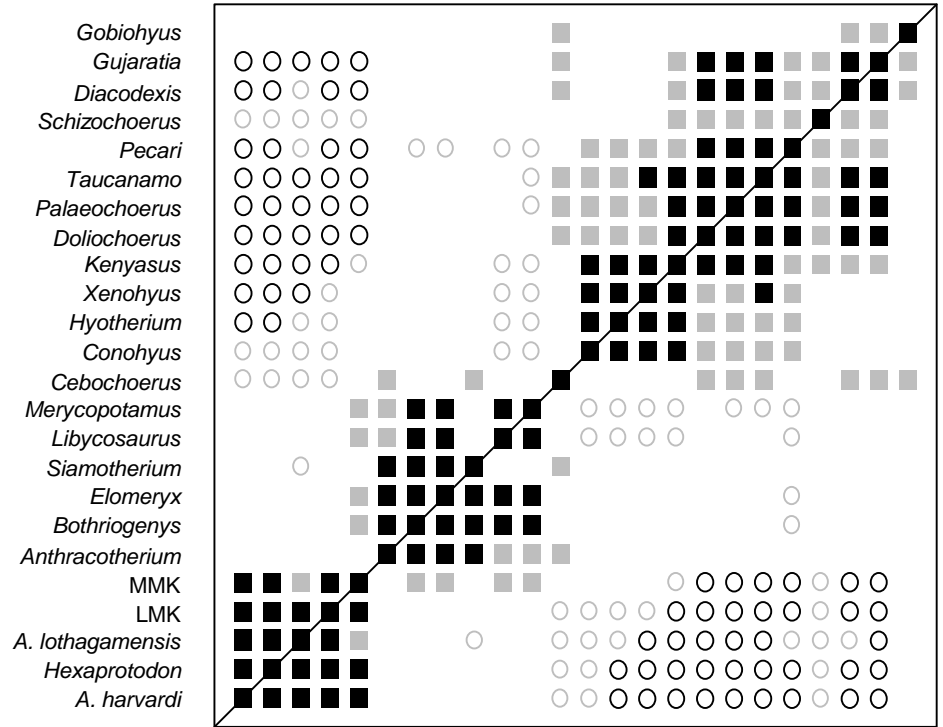
| | |
|----------------------------------|------|
| Published taxa | 41 |
| Published characters | 125 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 36 |
| Characters used for calculations | 87 |
| Median bootstrap value | 79 |
| F ₉₀ | 0.28 |
| Stress of 3D MDS | 0.2 |
| k _{min} | 8 |
| Conclusion | HB |



Notes: BDC supports recognizing Suidae as a holobaramin. MDS is less clear, but the suids are definitely separated from the outgroup taxa.

Boisserie, J.R., F. Lihoreau, M. Orliac, R.E. Fisher, E.M. Weston, and S. Ducrocq. 2010. Morphology and phylogenetic relationships of the earliest known hippopotamids (Cetartiodactyla, Hippopotamidae, Kenyapotaminae). *Zoological Journal of the Linnean Society* 158:325-366.

Characters: Craniodental and postcranial



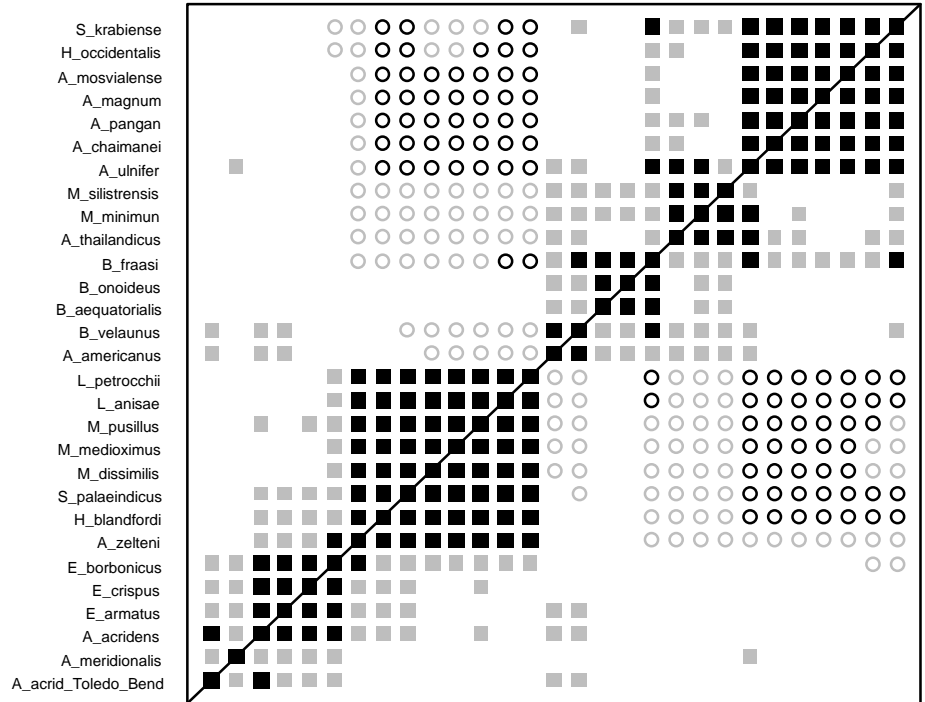
Order Artiodactyla
Family Hippopotamidae

| | |
|----------------------------------|------|
| Published taxa | 24 |
| Published characters | 87 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 24 |
| Characters used for calculations | 81 |
| Median bootstrap value | 84 |
| F ₉₀ | 0.38 |
| Stress of 3D MDS | 0.16 |
| k _{min} | 7 |
| Conclusion | HB? |

Notes: Both MDS and BDC reveal a distinct cluster of hippopotamids separate from other taxa. In the BDC results, the composite taxon “Middle Miocene Kenyapotamines” (MMK) shares significant, positive BDC with four outgroup taxa, but the correlations all have low bootstrap values. The MDS results reveal MMK well separated from the outgroup taxa. Thus, Hippopotamidae is probably a holobaramin.

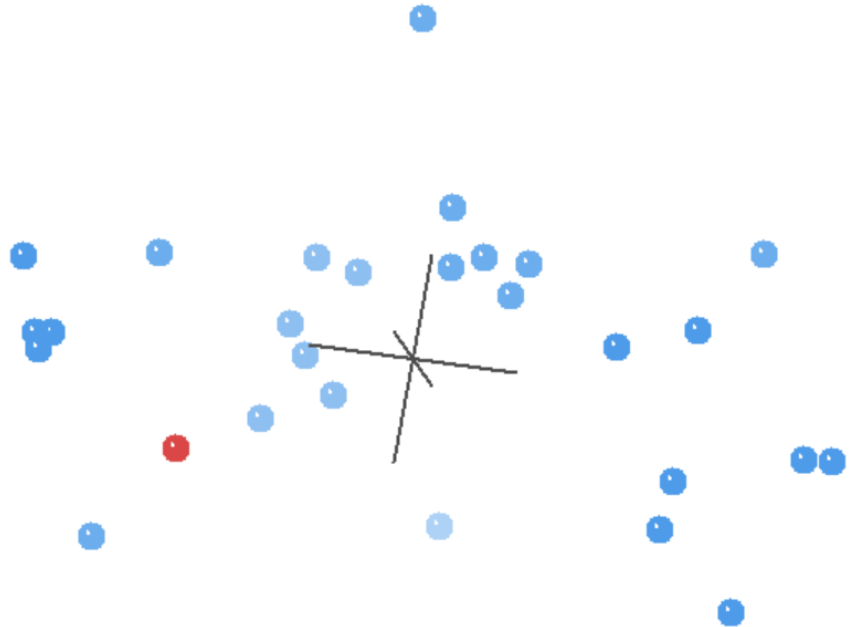
Rincon, A.F., J.I. Bloch, B.J. MacFadden, and C.A. Jaramillo. 2013. First central American record of Anthracotheriidae (Mammalia, Bothriodontinae) from the early Miocene of Panama. *Journal of Vertebrate Paleontology* 33:421-433.

Characters: Craniodental



Order Artiodactyla
Family Anthracotheriidae

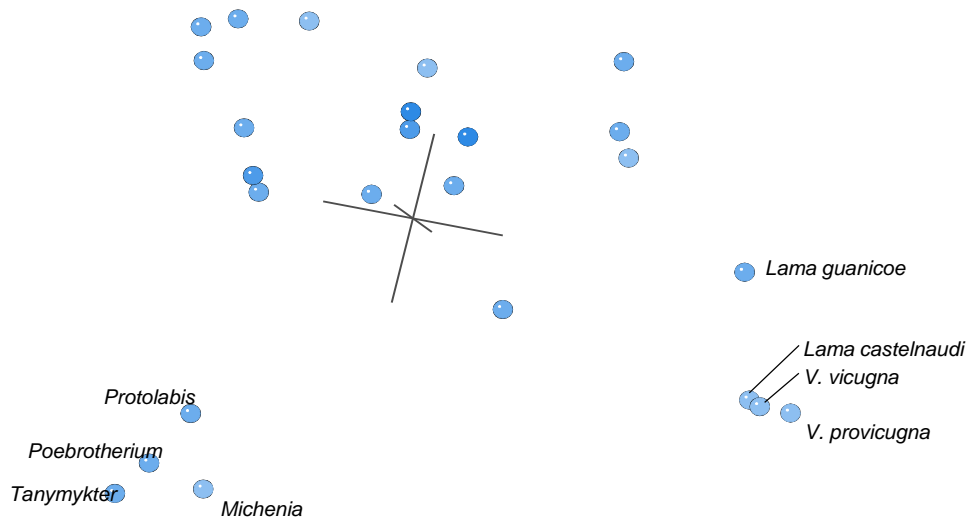
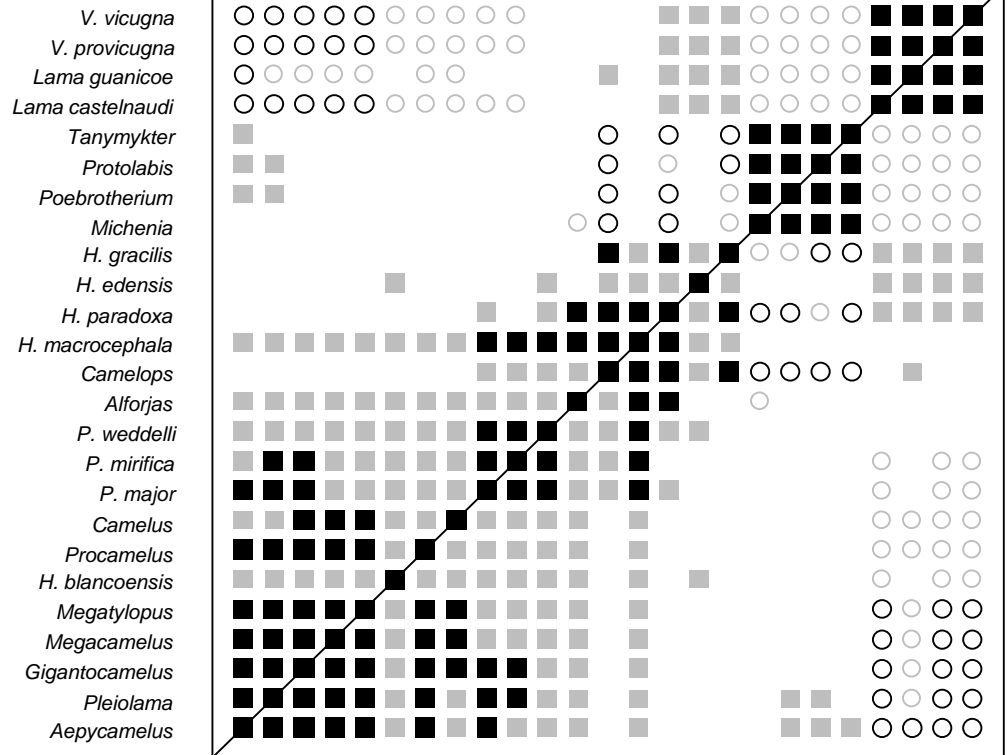
| | |
|----------------------------------|------|
| Published taxa | 29 |
| Published characters | 51 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 29 |
| Characters used for calculations | 41 |
| Median bootstrap value | 73 |
| F ₉₀ | 0.28 |
| Stress of 3D MDS | 0.15 |
| k _{min} | 4 |
| Conclusion | Inc |



Notes: No evidence of discontinuity.

Scherer, C.S. 2013. The Camelidae (Mammalia, Artiodactyla) from the Quaternary of South America: cladistic and biogeographic hypotheses. *Journal of Mammalian Evolution* 20:45-56.

Characters: Craniodental and postcranial



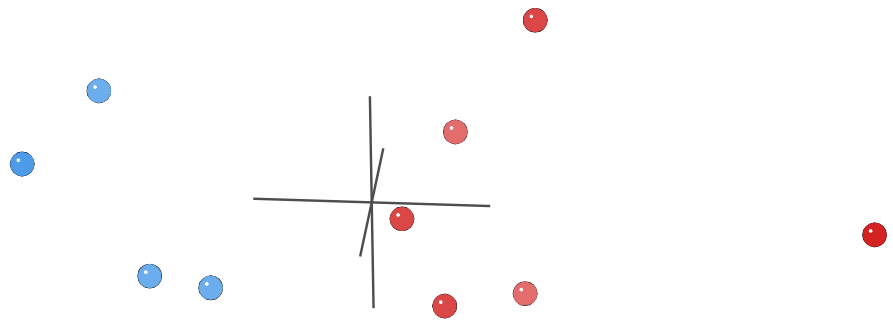
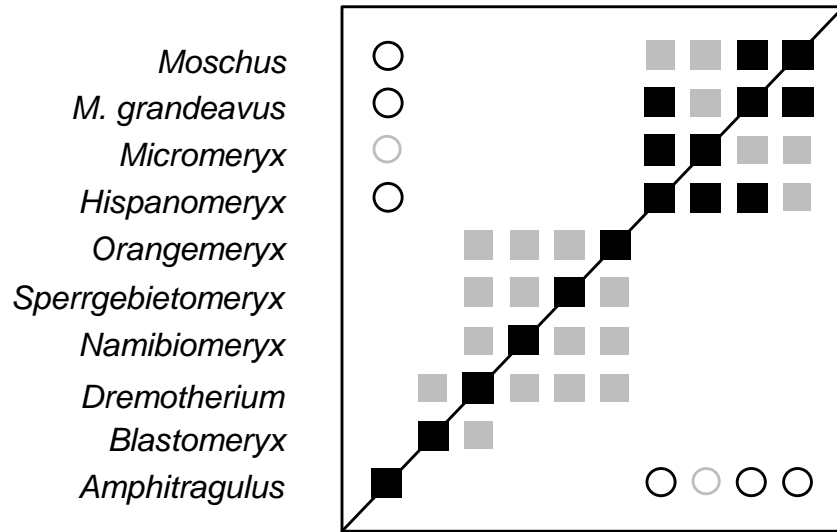
Order Artiodactyla
Family Camelidae

| | |
|----------------------------------|------|
| Published taxa | 25 |
| Published characters | 35 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 10 |
| Characters used for calculations | 25 |
| Median bootstrap value | 71 |
| F ₉₀ | 0.23 |
| Stress of 3D MDS | 0.14 |
| k _{min} | 3 |
| Conclusion | HB? |

Notes: Somewhat ambiguous BDC, with significant, negative BDC between basal camelids *Protolabis*, *Poebrotherium*, *Tanymykteer*, and *Michenia* and the rest of the camelids. MDS reveals clearer evidence of discontinuity, suggesting provisionally that Camelidae *sensu stricto* (excluding basal camelids) is a holobaramin.

Sanchez, I.M., M.S. Domingo, and J. Morales. 2010. The genus *Hispanomeryx* (Mammalia, Ruminantia, Moschidae) and its bearing on musk deer phylogeny and systematics. *Palaeontology* 53:1023-1047.

Characters: Craniodental and postcranial



Order Artiodactyla
Family Moschidae

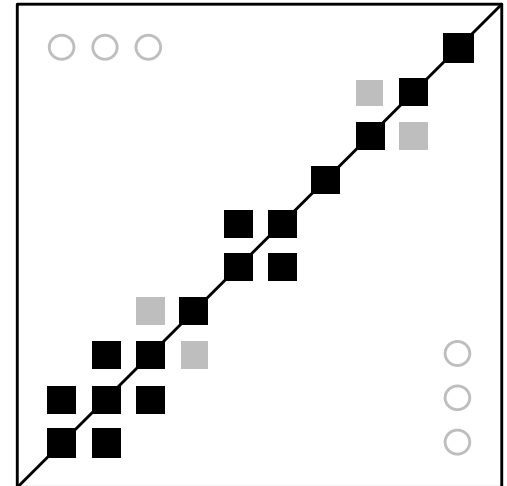
| | |
|----------------------------------|------|
| Published taxa | 10 |
| Published characters | 71 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 10 |
| Characters used for calculations | 40 |
| Median bootstrap value | 86 |
| F ₉₀ | 0.38 |
| Stress of 3D MDS | 0.08 |
| k _{min} | 4 |
| Conclusion | MB |

Notes: Moschidae is a definite cluster in the BDC results, but there is little significant, negative BDC. MDS shows Moschidae adjacent to outgroup taxa. Moschidae is a monobaramin.

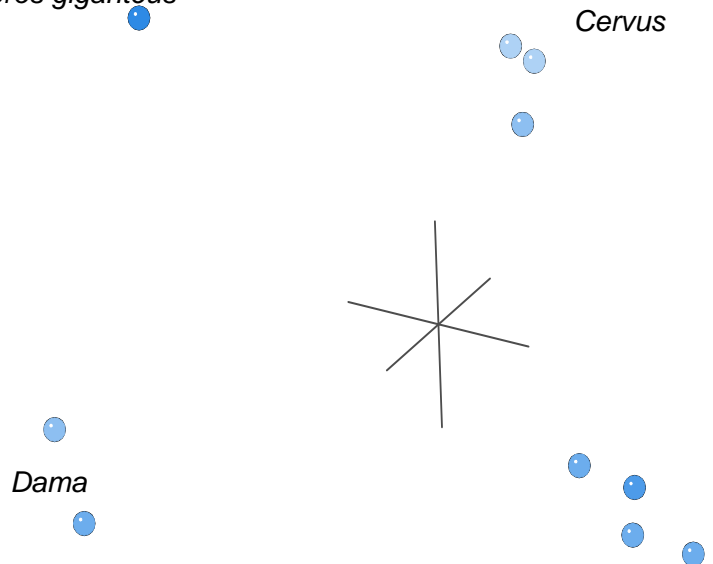
Lister, A.M., C.J. Edwards, D.A.W. Nock, M. Bunce, I.A. Van Pijlen, D.G. Bradley, M.G. Thomas, and I. Barnes. 2005. The phylogenetic position of the ‘giant deer’ *Megaloceros giganteus*. *Nature* 438:850-853.

Characters: Craniodental and postcranial

Megaloceros giganteus
Dama mesopotamica
Dama dama
Cervus nippon
Cervus elaphus
Cervus canadensis
Muntiacus spp.
Cervus eldi
Axis porcinus
Axis axis



Megaloceros giganteus



Order Artiodactyla
 Family Cervidae

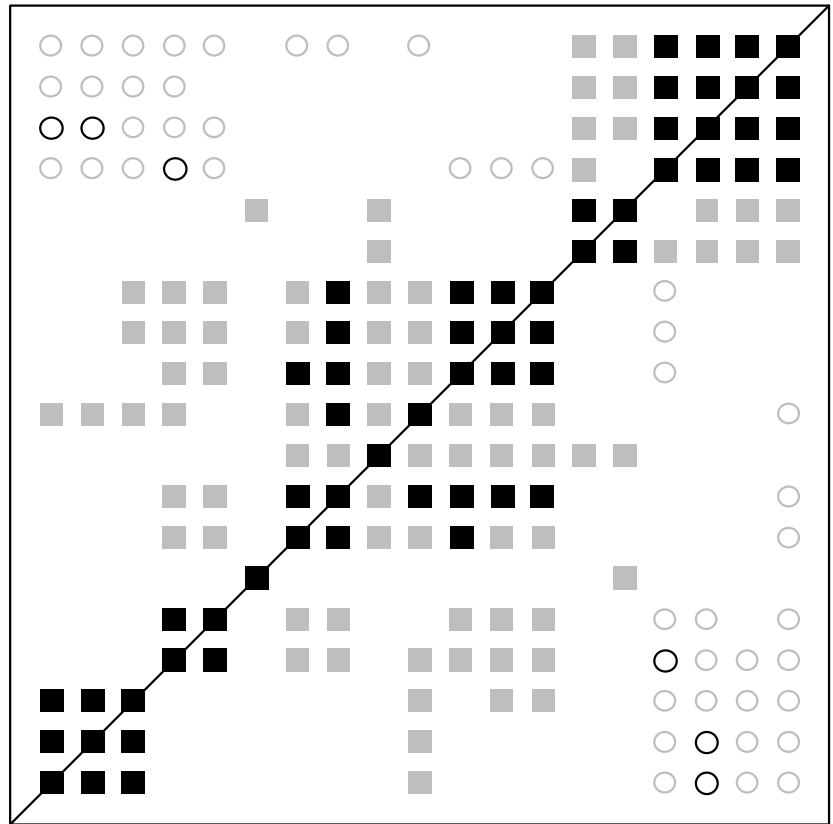
| | |
|----------------------------------|------|
| Published taxa | 10 |
| Published characters | 74 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 10 |
| Characters used for calculations | 74 |
| Median bootstrap value | 100 |
| F ₉₀ | 0.69 |
| Stress of 3D MDS | 0.23 |
| k _{min} | 7 |
| Conclusion | Inc |

Notes: Very few correlations in BDC, and a diffuse cluster of taxa in MDS. No clear discontinuity is evident.

Cerdeño, E. and B. Vera. 2010. *Mendozahippus fierensis*, gen. et sp. nov., new Notohippidae (Notoungulata) from the late Oligocene of Mendoza (Argentina). *Journal of Vertebrate Paleontology* 30:1805-1817.

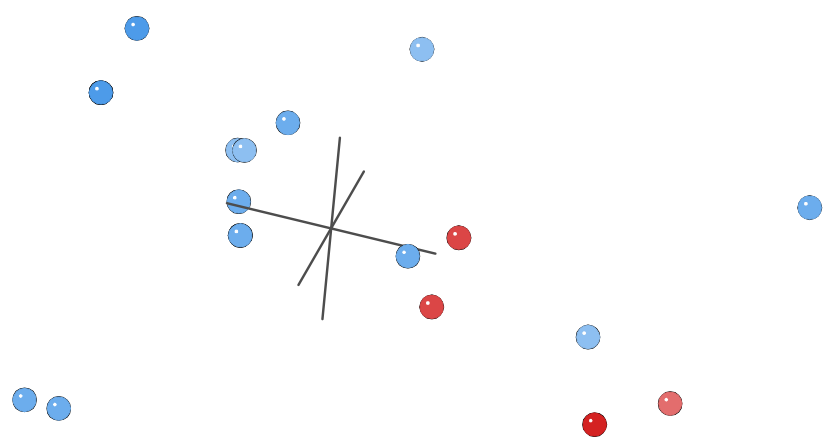
Characters: Craniodental and postcranial

Pleurostylodon modicus
Pampahippus arenalesi
Puelia coarctatus
P. complicatissimus
Scarrittia canquelensis
Leontinia gaudryi
Rhynchippus pumilus
Rhynchippus equinus
R. brasiliensis
P. boliviensis
Mendozahippus fierensis
Eurygenium pacegnum
Eurygenium latirostris
Eomorphippus obscurus
Argyrohippus praecox
Argyrohippus fraterculus
P. leptognathum
Nesodon imbricatus
Adinotherium ovinum



Order Notoungulata
 Family Notohippidae

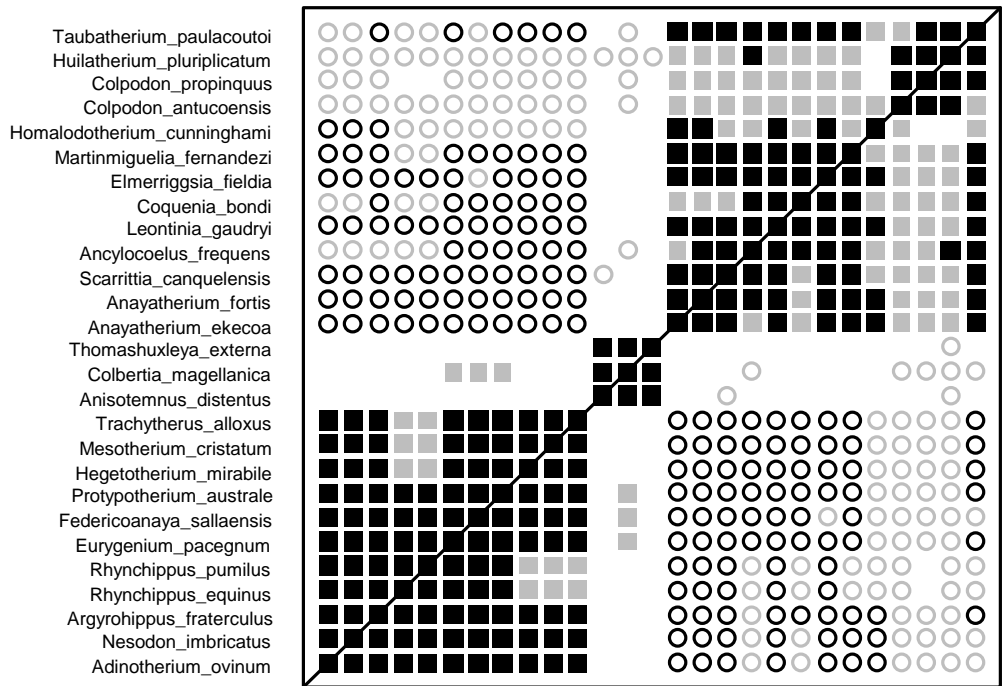
| | |
|----------------------------------|------|
| Published taxa | 19 |
| Published characters | 38 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 19 |
| Characters used for calculations | 29 |
| Median bootstrap value | 73 |
| F ₉₀ | 0.13 |
| Stress of 3D MDS | 0.14 |
| k _{min} | 4 |
| Conclusion | Inc |



Notes: BDC reveals two groups with substantial overlap, and MDS confirms. No clear discontinuity is evident.

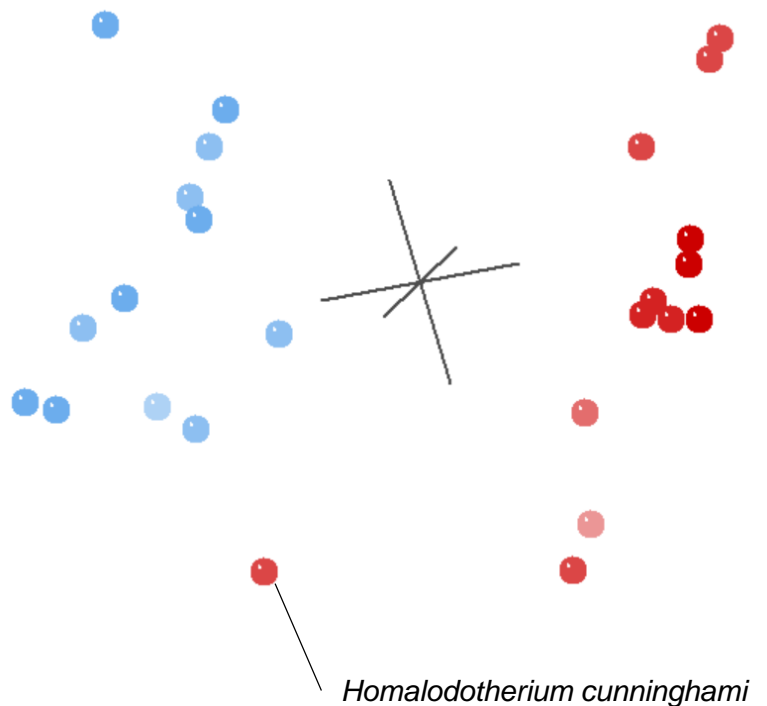
Schockey, B.J., J.J. Flynn, D.A. Croft, P. Gans, and R.A. Wyss. 2012. New leontiniid Notoungulata (Mammalia) from Chile and Argentina: comparative anatomy, character analysis, and phylogenetic hypotheses. *American Museum Novitates* 3737:1-64.

Characters: Craniodental and postcranial



Order Notoungulata
Family Leontiniidae

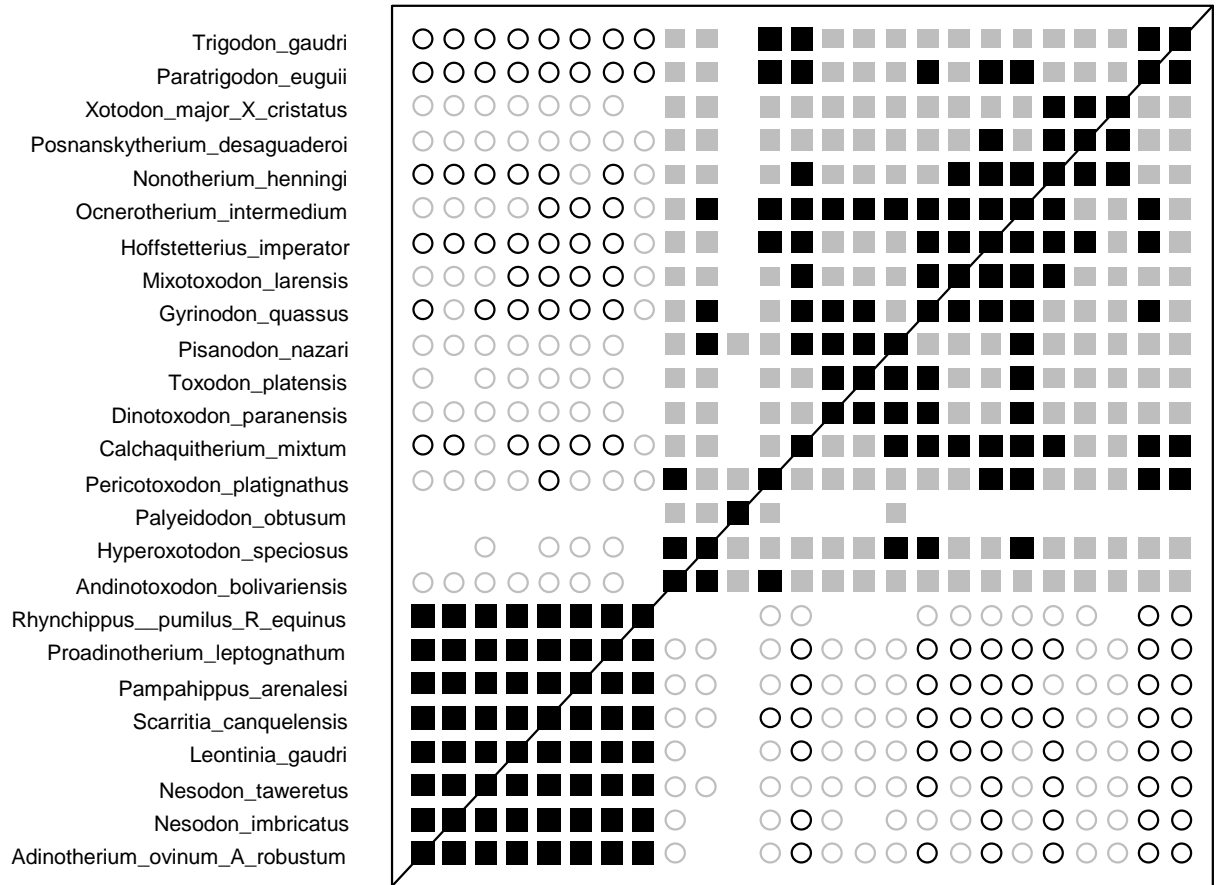
| | |
|----------------------------------|------|
| Published taxa | 27 |
| Published characters | 83 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 27 |
| Characters used for calculations | 36 |
| Median bootstrap value | 90 |
| F ₉₀ | 0.49 |
| Stress of 3D MDS | 0.13 |
| k _{min} | 4 |
| Conclusion | HB |



Notes: BDC and MDS reveal two groups corresponding to Leontiniidae and the outgroup. They are well separated in both analyses. Leontiniidae is likely a holobaramin.

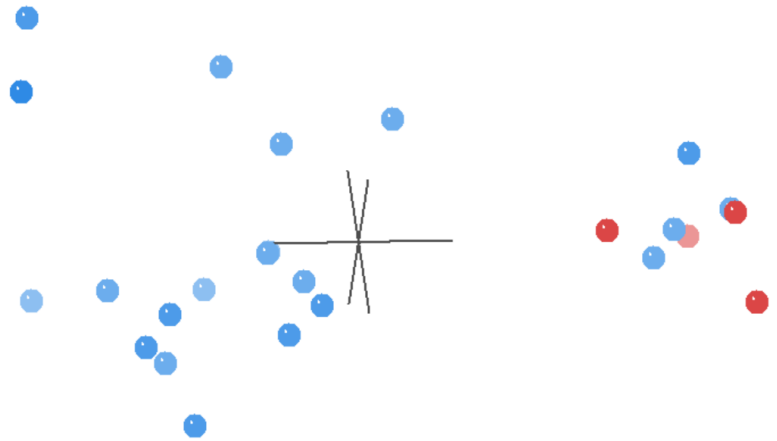
Forasiepi, A.M., E. Cerdeno, M. Bond, G.I. Schmidt, M. Naipauer, F.R. Straehl, A.G. Martinelli, A.C. Garrido, M.D. Schmitz, and J.J. Crowley. 2015. New toxodontid (Notoungulata) from the early Miocene of Mendoza, Argentina. *Paläontologische Zeitschrift* 89:611-634.

Characters: Craniodental



Order Notoungulata
Family Toxodontidae

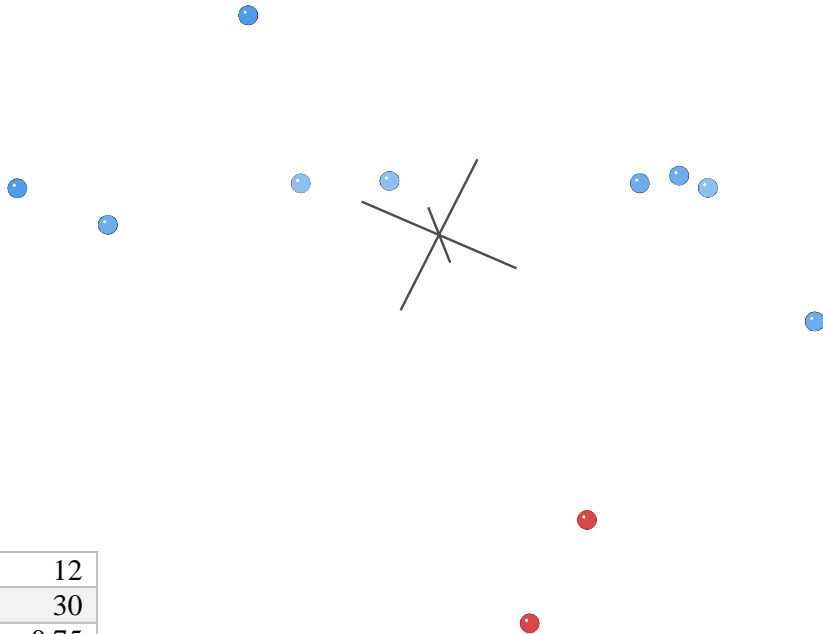
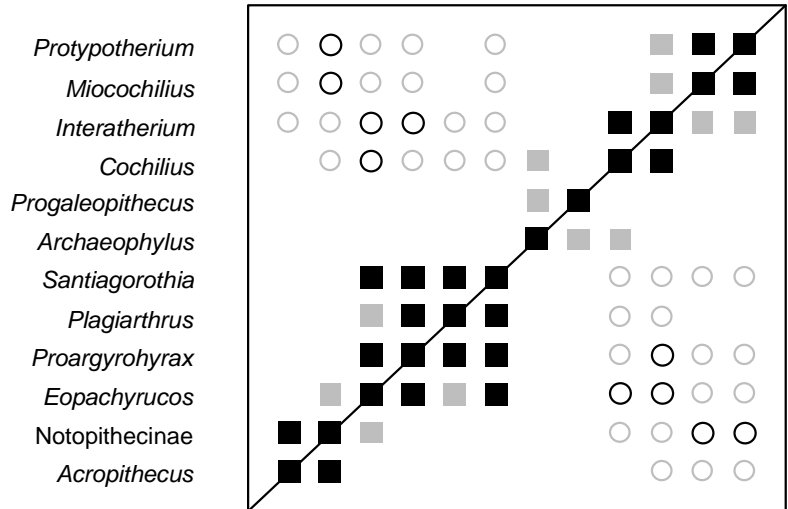
| | |
|----------------------------------|------|
| Published taxa | 25 |
| Published characters | 59 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 25 |
| Characters used for calculations | 33 |
| Median bootstrap value | 86 |
| F ₉₀ | 0.38 |
| Stress of 3D MDS | 0.14 |
| k _{min} | 4 |
| Conclusion | HB |



Notes: Two groups of taxa are distinguishable in the BDC and MDS results. The four outgroup taxa cluster with basal toxodontids. Toxodontidae *sensu stricto* is a holobaramin.

Reguero, M.A., M. Ubilla, and D. Perea. 2003. A new species of *Eopachyrucos* (Mammalia, Notoungulata, Interatheriidae) from the late Oligocene of Uruguay. *Journal of Vertebrate Paleontology* 23:445-457.

Characters: Craniodental and postcranial



Order Notoungulata
Family Interatheriidae

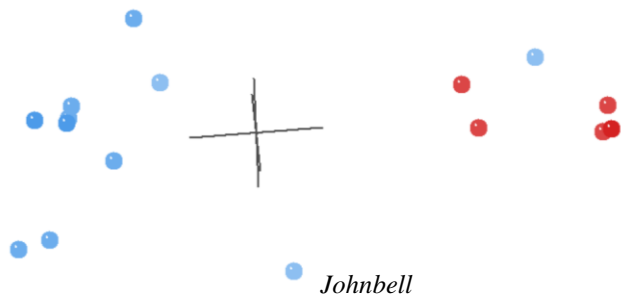
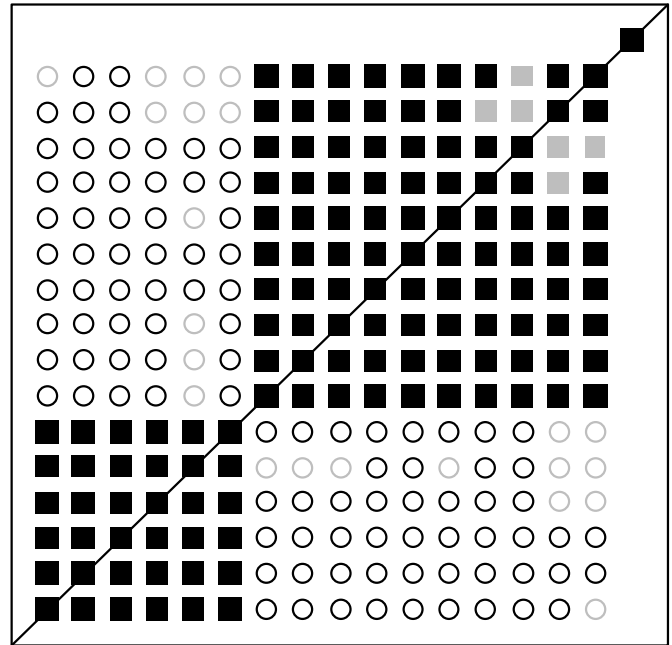
| | |
|----------------------------------|------|
| Published taxa | 12 |
| Published characters | 30 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 12 |
| Characters used for calculations | 26 |
| Median bootstrap value | 78.5 |
| F ₉₀ | 0.23 |
| Stress of 3D MDS | 0.08 |
| k _{min} | 3 |
| Conclusion | Inc |

Notes: BDC reveals two groups with no overlap, neither of which corresponds to a named taxonomic group. The MDS does not confirm these groups but instead shows separation between the ingroup Interatheriidae and two taxa corresponding to the outgroup *Acropithecus* and the composite interatheriid subfamily Notopithecinae. These results are inconclusive.

Hitz, R.B., J.J. Flynn, and A.R. Wyss. 2006. New basal Interatheriidae (Typotheria, Notoungulata, Mammalia) from the Paleogene of Central Chile. *American Museum Novitates* 3520:1-32.

Characters: Craniodental

Johnbell_hatcheri
 Santiagorothia_chiliensis
 Proargyrohyrax
 Protypotherium
 Miocochilius
 Plagiarthrus
 Interatherium
 Cochilius
 Salla_new_taxonB
 Salla_new_taxonA
 Archaeophylus
 Punapithecus
 Transpithecus
 Guiliemoscottia
 Notopithecus
 Colbertia
 Antepithecus



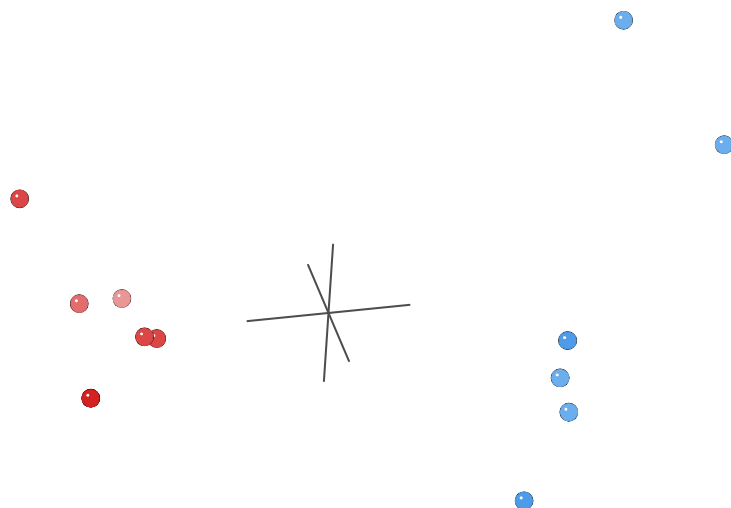
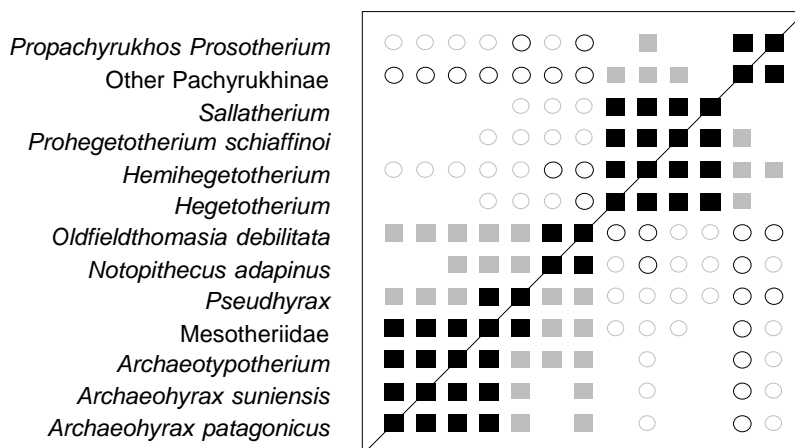
Order Notoungulata
 Family Interatheriidae

| | |
|----------------------------------|------|
| Published taxa | 21 |
| Published characters | 43 |
| Character relevance | 0.75 |
| Taxic relevance | 0.45 |
| Taxa used for calculations | 17 |
| Characters used for calculations | 21 |
| Median bootstrap value | 98 |
| F ₉₀ | 0.76 |
| Stress of 3D MDS | 0.04 |
| k _{min} | 3 |
| Conclusion | HB |

Notes: BDC and MDS support a distinct Interatheriidae cluster. The interatheriid *Punapithecus* clusters with the outgroup, and another, *Johnbell*, shares no BDC with any other taxa. Interatheriidae *sensu stricto* could be a holobaramin.

Billet, G., B. Patterson, and C. De Muizon. 2009. Craniodental anatomy of late Oligocene archaeohyracids (Notoungulata, Mammalia) from Bolivia and Argentina and new phylogenetic hypotheses. *Zoological Journal of the Linnean Society* 155:458-509.

Characters: Craniodental and postcranial

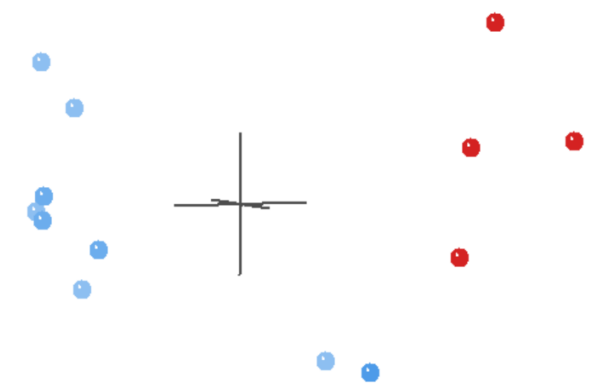
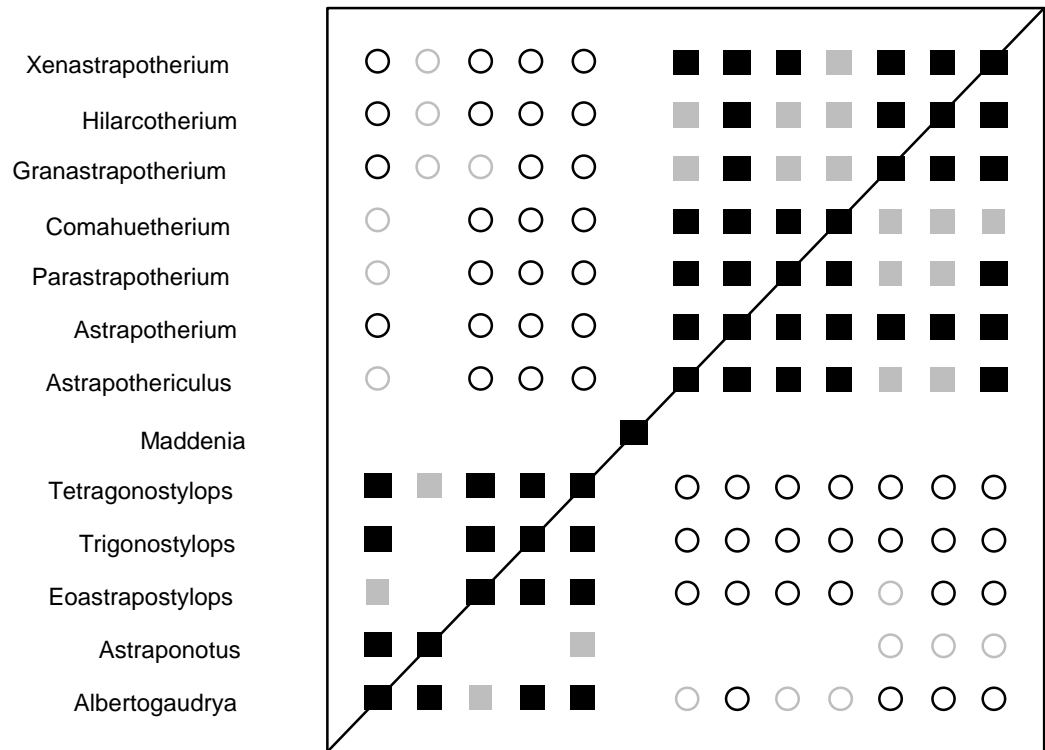


Order Notoungulata
Family Hegetotheriidae

| | |
|----------------------------------|------|
| Published taxa | 15 |
| Published characters | 39 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 13 |
| Characters used for calculations | 32 |
| Median bootstrap value | 82.5 |
| F ₉₀ | 0.29 |
| Stress of 3D MDS | 0.07 |
| k _{min} | 3 |
| Conclusion | HB |

Notes: Hegetotheriidae and outgroup taxa are well-separated in both BDC and MDS results. Hegetotheriidae is likely a holobaramin.

Vallejo-Pareja, M.C., J.D. Carrillo, J.W. Moreno-Bernal, M. Pardo-Jaramillo, D.F. Rodriguez-Gonzalez, and J. Muñoz-Duran. 2015. *Hilarchotherium castanedaii*, gen. et sp. nov., a new Miocene astrapothere (Mammalia, Astrapotheriidae) from the Upper Magdalena Valley, Colombia. *Journal of Vertebrate Paleontology* 35:e903960. Characters: Craniodental



Order Astrapotheria
Family Astrapotheriidae

| | |
|----------------------------------|------|
| Published taxa | 15 |
| Published characters | 64 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 13 |
| Characters used for calculations | 45 |
| Median bootstrap value | 93 |
| F ₉₀ | 0.59 |
| Stress of 3D MDS | 0.09 |
| k _{min} | 5 |
| Conclusion | HB |

Notes: BDC and MDS reveal two well-separated groups. One astrapotheriid *Astraponotus* clusters with the outgroup taxa, and *Maddenia* does not cluster with either larger group. Astrapotheriidae *sensu stricto* (excluding *Astraponotus* and *Maddenia*) is probably a holobaramin.

Antoine, P.O., G. Billet, R. Salas-Gismondi, J.T. Lara, P. Baby, S. Brusset, and N. Espurt. 2015. A new *Carodnia* Simpson, 1935 (Mammalia, Xenungulata) from the early Eocene of Northwestern Peru and a phylogeny of xenungulates at species level. *Journal of Mammalian Evolution* 22:129-140.

Characters: Craniodental

Carodnia inexpectans sp. nov.

Carodnia feruglioi

Carodnia vieirai

Carodnia cf. *feruglioi*

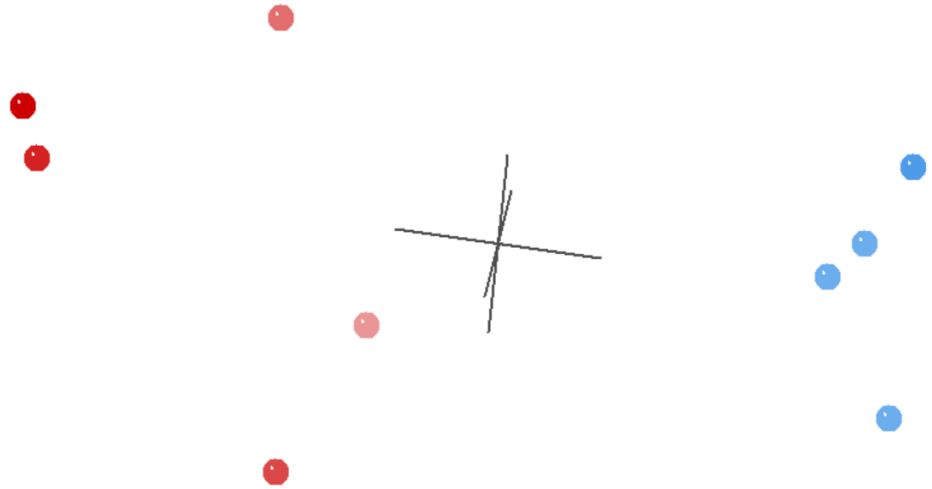
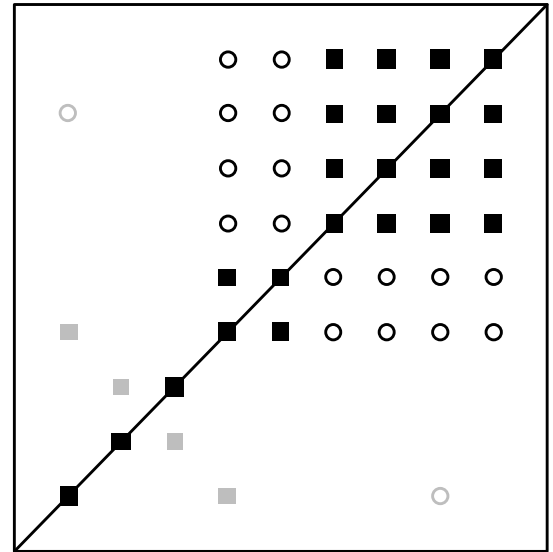
Didolodus multicuspis

Asmithwoodwardia scotti

Notoetayoa gargantuai

Etayoa bacatensis

Alcidedorbignya inopinata



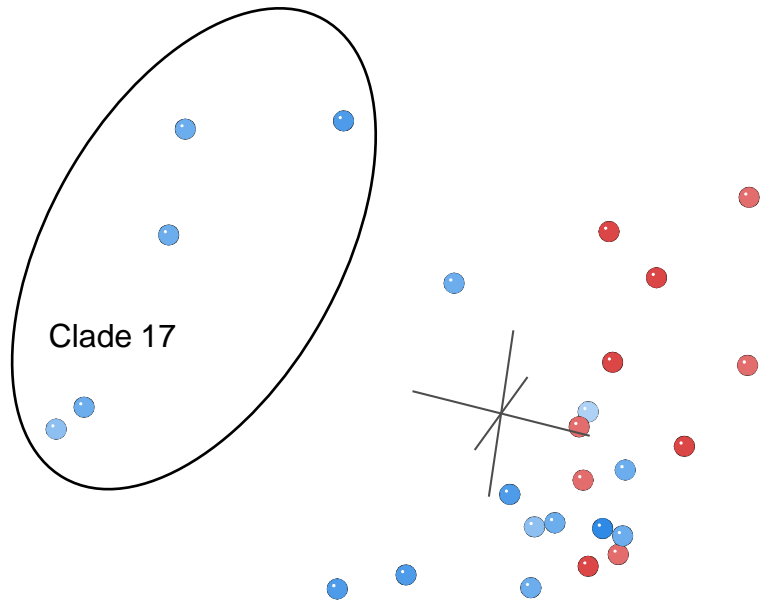
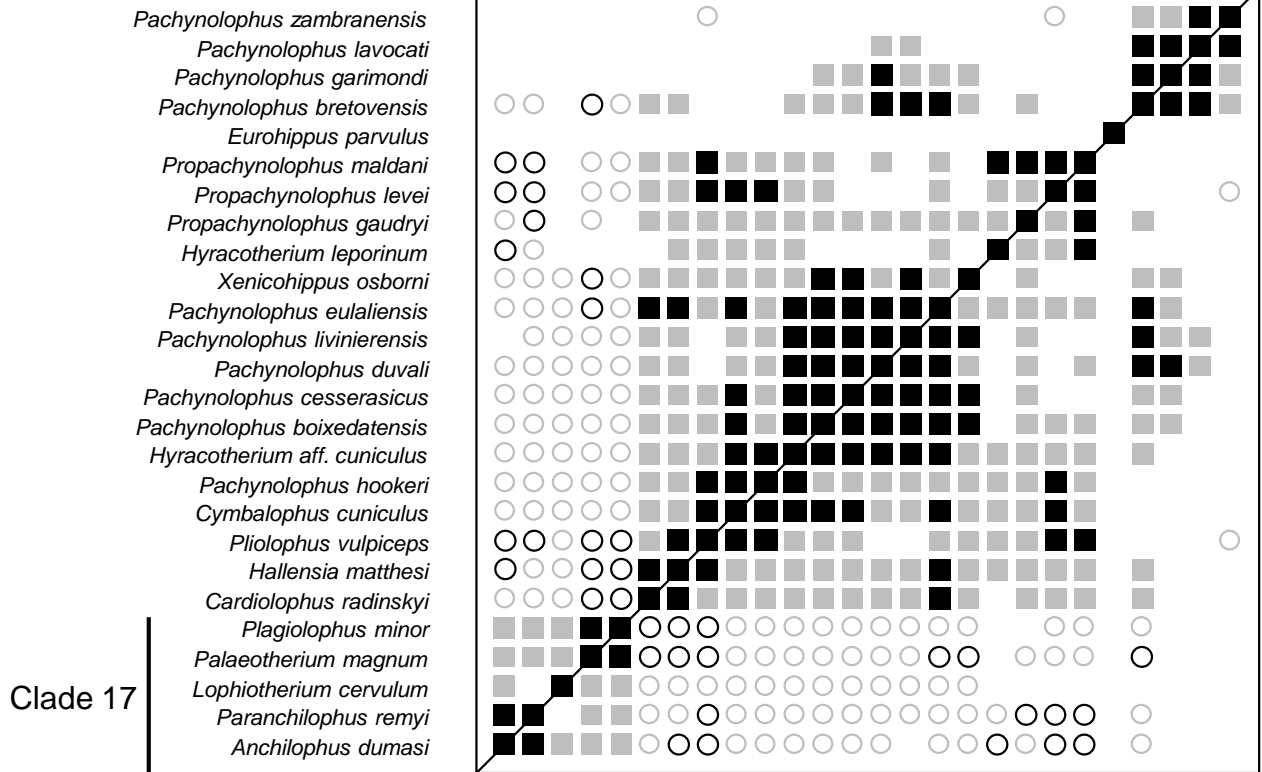
Order Xenungulata
Family Carodniidae

| | |
|----------------------------------|------|
| Published taxa | 9 |
| Published characters | 34 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 9 |
| Characters used for calculations | 28 |
| Median bootstrap value | 92.5 |
| F ₉₀ | 0.53 |
| Stress of 3D MDS | 0.1 |
| k _{min} | 3 |
| Conclusion | HB? |

Notes: BDC and MDS reveal two well-separated groups of Carodniidae and outgroup taxa. Carodniidae is likely a holobaramin.

Danilo, L., J.A. Remy, M. Vianey-Liaud, B. Marandat, J. Sudre, and F. Lihoreau. 2013. A new Eocene locality in southern France sheds light on the basal radiation of Palaeotheriidae (Mammalia, Perissodactyla, Equoidea). *Journal of Vertebrate Paleontology* 33:195-215.

Characters: Craniodental

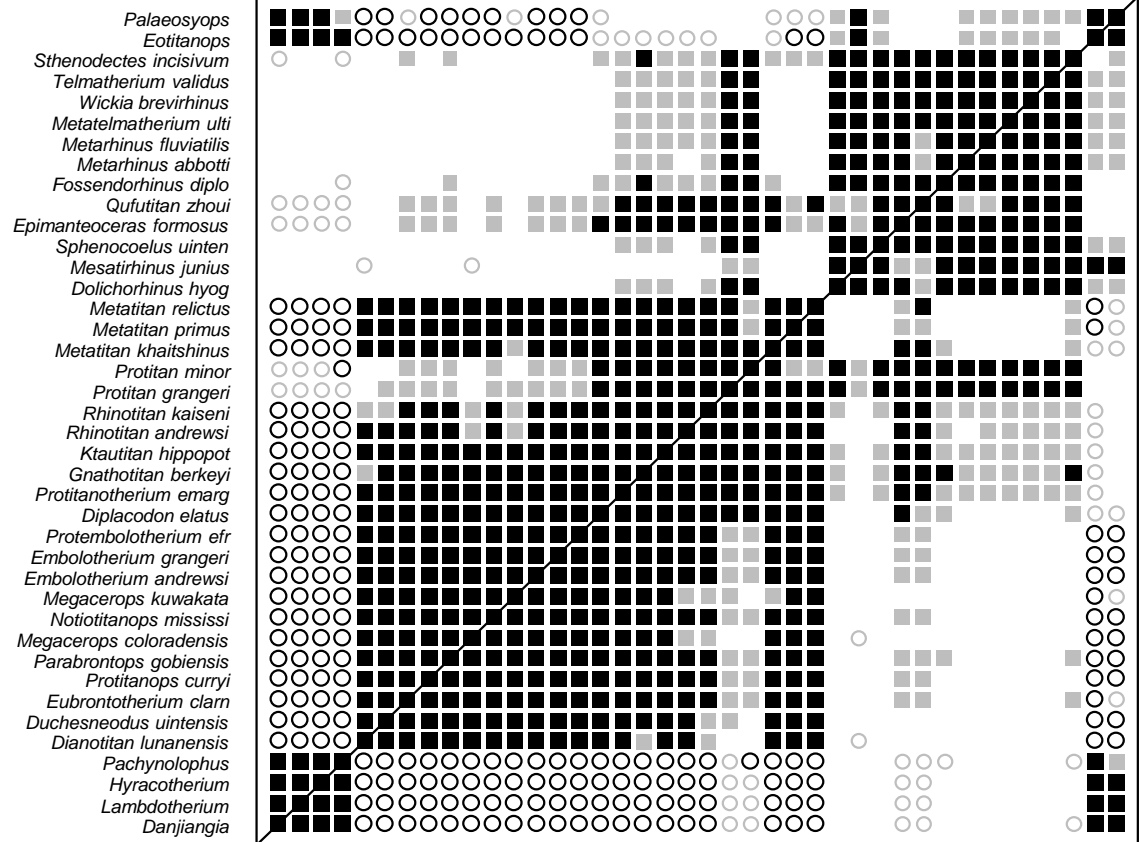


Order Perissodactyla
Family Palaeotheriidae

| | |
|----------------------------------|------|
| Published taxa | 26 |
| Published characters | 72 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 26 |
| Characters used for calculations | 57 |
| Median bootstrap value | 81 |
| F ₉₀ | 0.22 |
| Stress of 3D MDS | 0.25 |
| k _{min} | 8 |
| Conclusion | HB? |

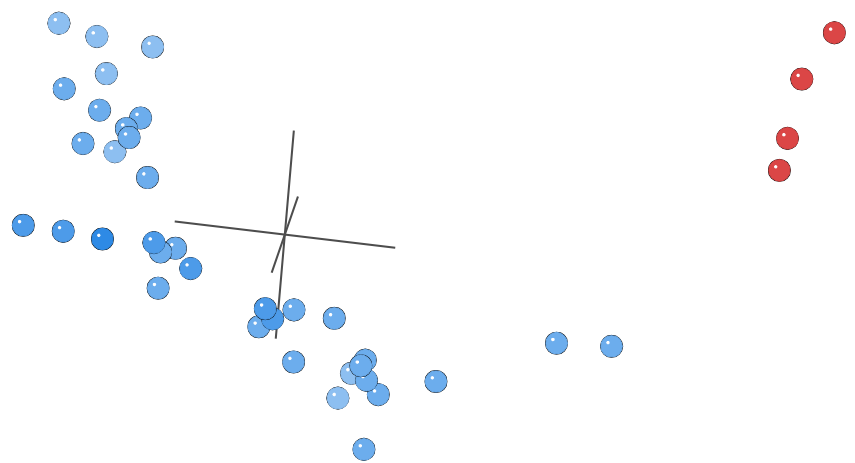
Notes: BDC indicates good evidence for discontinuity around a group of five taxa corresponding to clade 17 in Danilo et al.'s (2013) Figure 9. MDS seems to support the separation of this cluster of taxa. Clade 17 includes Palaeotheriinae (*Plagiolophus minor*, *Palaeotherium magnum*) and three additional taxa included here: *Lophiotherium cervulum*, *Paranchilophus remyi*, and *Anchilophus dumasi*. Clade 17 could be a holobaramin.

Characters: Craniodental



Order Perissodactyla
Family Brontotheriidae

| | |
|----------------------------------|------|
| Published taxa | 51 |
| Published characters | 87 |
| Character relevance | 0.75 |
| Taxic relevance | 0.5 |
| Taxa used for calculations | 40 |
| Characters used for calculations | 81 |
| Median bootstrap value | 94 |
| F ₉₀ | 0.53 |
| Stress of 3D MDS | 0.17 |
| k _{min} | 7 |
| Conclusion | HB? |

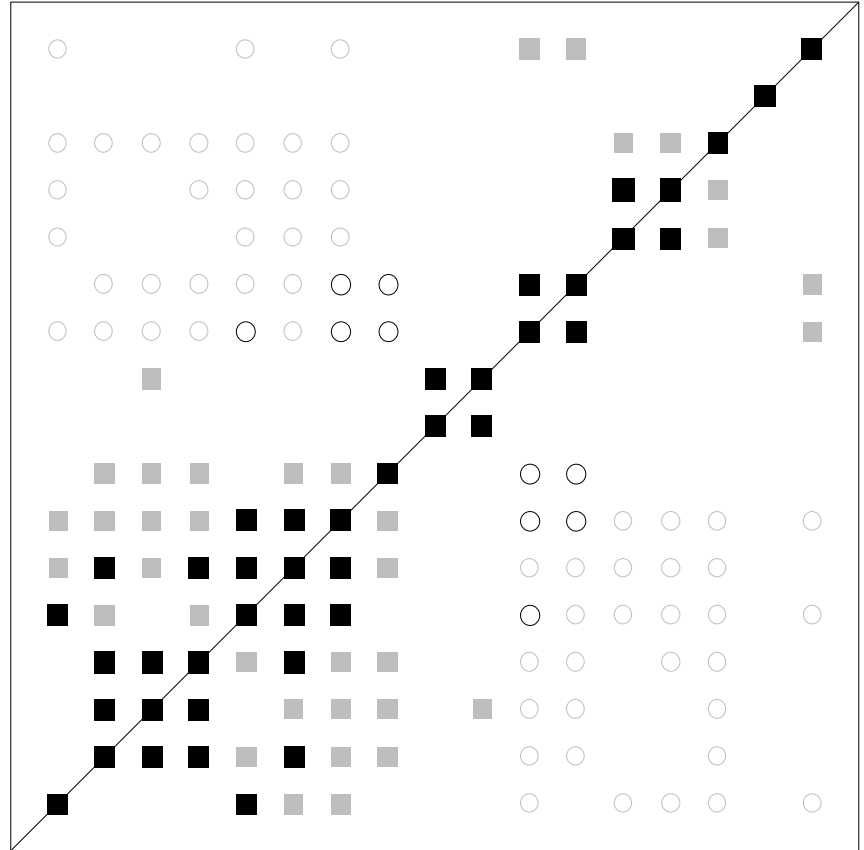


Notes: In BDC, the Brontotheriidae are well separated from the outgroup, except for two brontotheres, *Paleosyops* and *Eotitanops*, which share significant, positive BDC with all outgroup taxa. The MDS shows an arc of taxa but with a separation between the outgroup and the Brontotheriidae. This suggests that the correlations involving the outgroups, *Paleosyops*, and *Eotitanops* are a result of the linear geometry of the brontothere taxa. Taken together, this suggests that Brontotheriidae is likely a holobaramin.

Bai, B., Y. Wang, and J. Meng. 2010. New craniodental materials of *Litolophus gobiensis* (Perissodactyla, “Eomoropidae”) from Inner Mongolia, China, and phylogenetic analyses of Eocene chalicotheres. *American Museum Novitates* 3688:1-27.

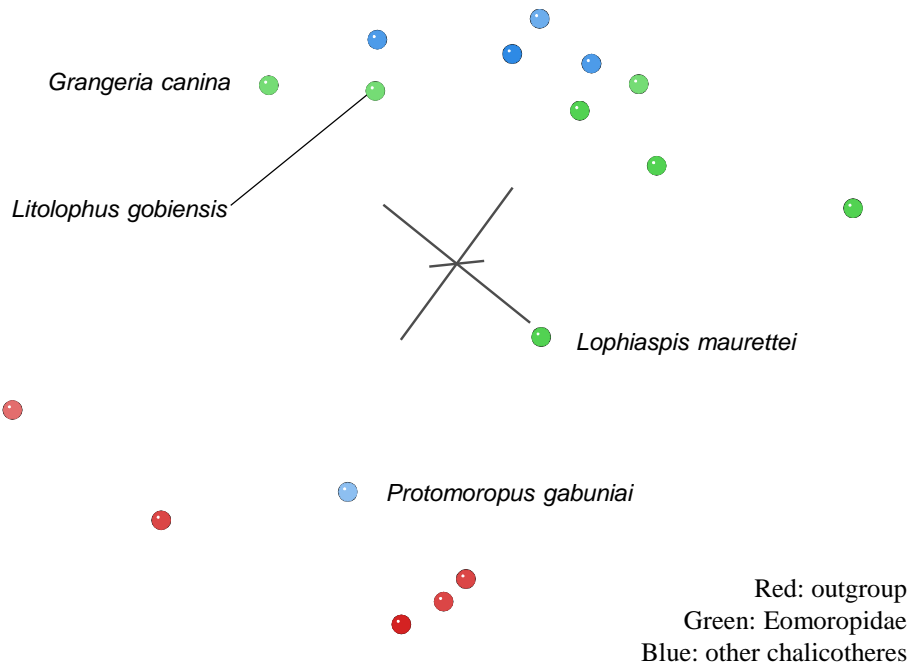
Characters: Craniodental

Protomoropus gabuniai
Lophiaspis maurettei
Lophiodon remensis
Hyrachyus eximius
Heptodon calciculus
Homogalax wutuensis
Cardiolphus radinskyi
Litolophus gobiensis
Grangeria canina
Eomoropus pawnyunti
Schizotherium avitum
SDM 84006
Moropus elatus
Eomoropus quadridentatus
Grangeria anarsius
Eomoropus amarorum
Ansiodon grande



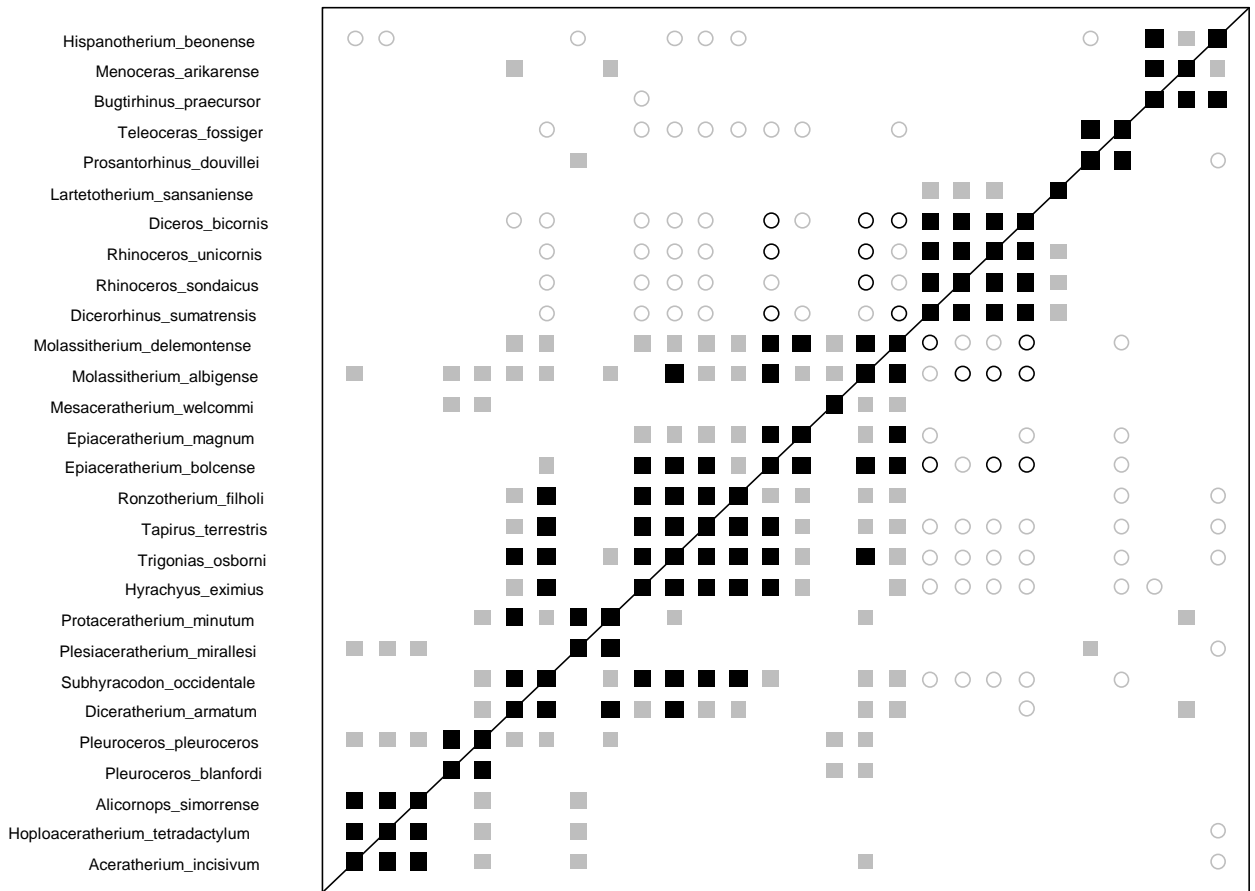
Order Perissodactyla
 Family Chalicotheriidae

| | |
|----------------------------------|------|
| Published taxa | 21 |
| Published characters | 58 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 17 |
| Characters used for calculations | 38 |
| Median bootstrap value | 77 |
| F ₉₀ | 0.16 |
| Stress of 3D MDS | 0.2 |
| k _{min} | 6 |
| Conclusion | HB? |



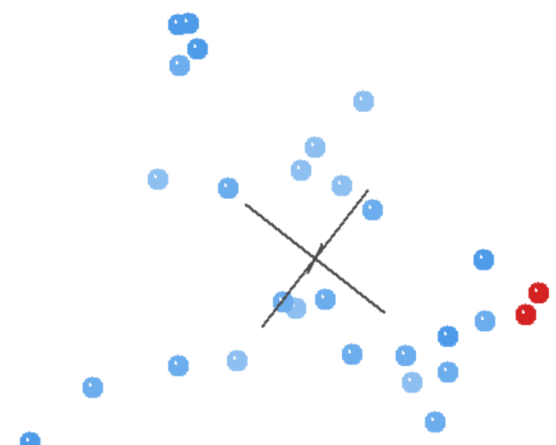
Notes: The BDC results reveal two groups of taxa, corresponding roughly to the chalicotheres and other taxa. The two groups are well-separated, but the chalicothere *Protomoropus* is part of the outgroup. The chalicothere *Lophiaspis* does not share significant BDC with any other taxa. The MDS results reveal the same groups with *Protomoropus* in the outgroup cluster and *Lophiaspis* in between the two clusters. The chalicotheres (excluding *Protomoropus* and *Lophiaspis*) are likely a holobaramin.

Characters: Craniodental and postcranial



Order Perissodactyla
Family Rhinocerotidae

| | |
|----------------------------------|------|
| Published taxa | 30 |
| Published characters | 214 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 28 |
| Characters used for calculations | 161 |
| Median bootstrap value | 89 |
| F ₉₀ | 0.45 |
| Stress of 3D MDS | 0.34 |
| k _{min} | 15 |
| Conclusion | Inc |

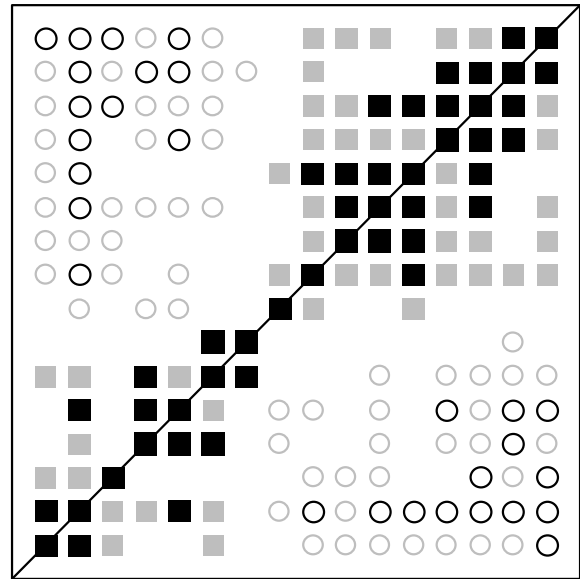


Notes: No evidence of discontinuity.

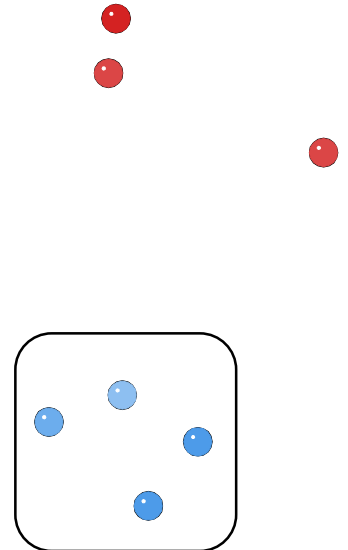
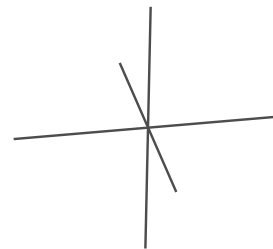
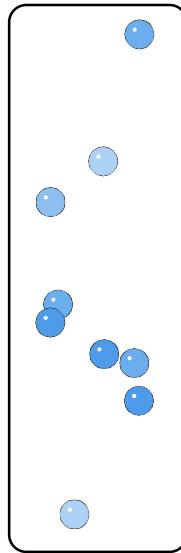
Robinet, C., J.A. Remy, J.A., Y. Laurent, L. Danilo, and F. Lihoreau. 2015. A new genus of Lophiodontidae (Perissodactyla, Mammalia) from the early Eocene of La Borie (Southern France) and the origin of the genus *Lophiodon* Cuvier, 1822. *Geobios* 48:25-38.

Characters: Craniodental

Lophiodon lauricense
Lophiodon rhinocerodes
Lophiodon tapirotherium
Lophiodon remense
Paralophiodon leptorhynchum
Paralophiodon isselense
Paralophiodon buchsowillanum
Lophiodon baroensis
Eolophiodon laboriense
Lophiaspis occitanicus
Lophiaspis maurettei
Litolophus gobiensis
Eomoropus amarorum
Pachynolophus livinierensis
Phenacodus intermedius
Cardiopholophus



Lophiodontinae



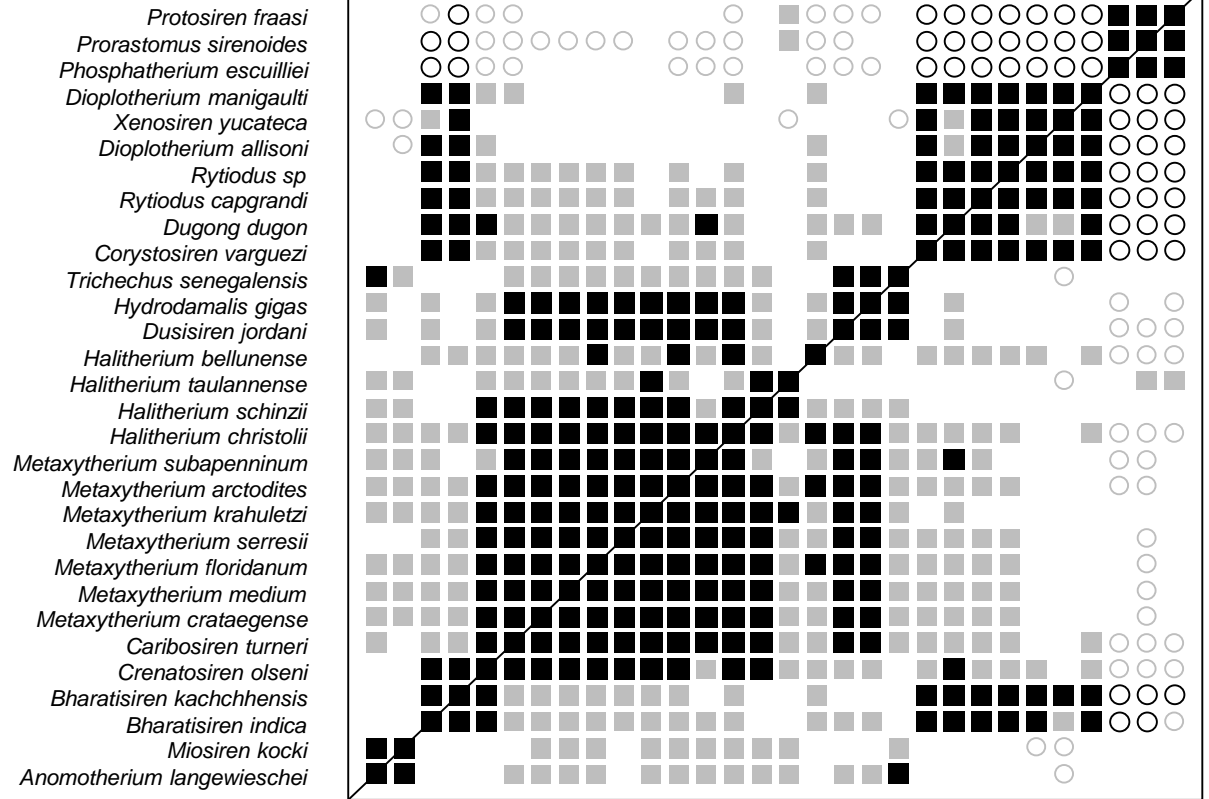
Lophiaspis,
Eomoropus,
Litolophus

Order Perissodactyla
 Family Lophiodontidae

| | |
|----------------------------------|------|
| Published taxa | 18 |
| Published characters | 149 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 16 |
| Characters used for calculations | 118 |
| Median bootstrap value | 77 |
| F ₉₀ | 0.28 |
| Stress of 3D MDS | 0.24 |
| k _{min} | 10 |
| Conclusion | HB |

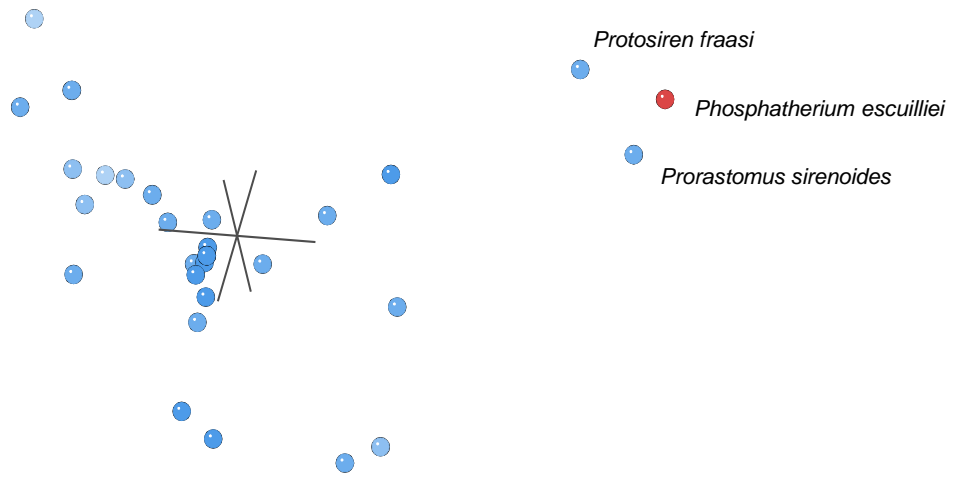
Notes: BDC and MDS both support two distinct groups, one consisting of subfamily Lophiodontinae (*Lophiodon*, *Paralophiodon*, and *Eolophiodon*), and the other containing all other taxa. Lophiodontinae is likely a holobaramin.

Characters: Craniodental



Order Uranotheria
Infraorder Sirenia

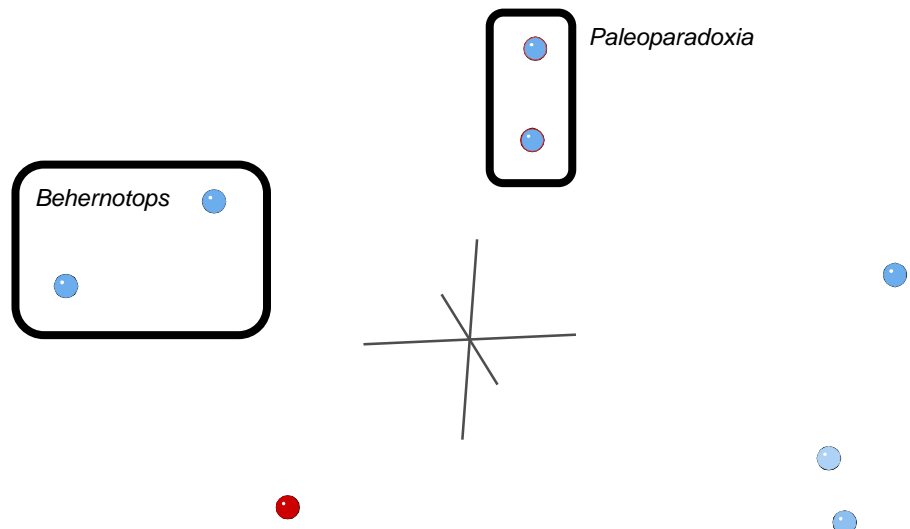
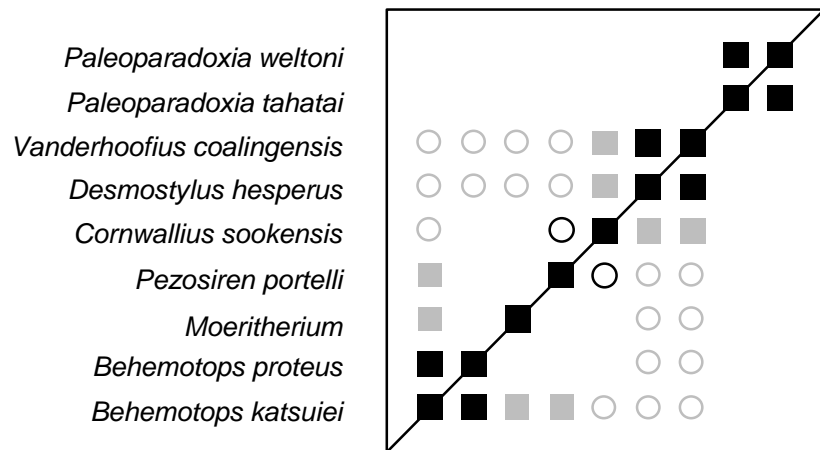
| | |
|----------------------------------|------|
| Published taxa | 32 |
| Published characters | 57 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 7 |
| Characters used for calculations | 50 |
| Median bootstrap value | 78 |
| F ₉₀ | 0.33 |
| Stress of 3D MDS | 0.16 |
| k _{min} | 5 |
| Conclusion | HB |



Notes: BDC and MDS both support two groups: 1. The outgroup *Phosphatherium escuilliei* together with the sirenians *Protosiren fraasi* and *Prorastomus sirenoides*. 2. The rest of the Sirenia. There is significant, positive BDC with low bootstrap values only between *Halitherium taulannense* and the two sirenians that cluster with the outgroup, but in the MDS results, *Halitherium taulannense* is part of the sirenian cluster and separate from the three taxa in the outgroup cluster. We may infer that Sirenia *sensu stricto* (excluding *Protosiren* and *Prorastomus*) is a holobaramin.

Beatty, B.L. 2009. New material of *Cornwallius sookensis* (Mammalia: Desmostylia) from the Yaquina Formation of Oregon. *Journal of Vertebrate Paleontology* 29:894-909.

Characters: Craniodental



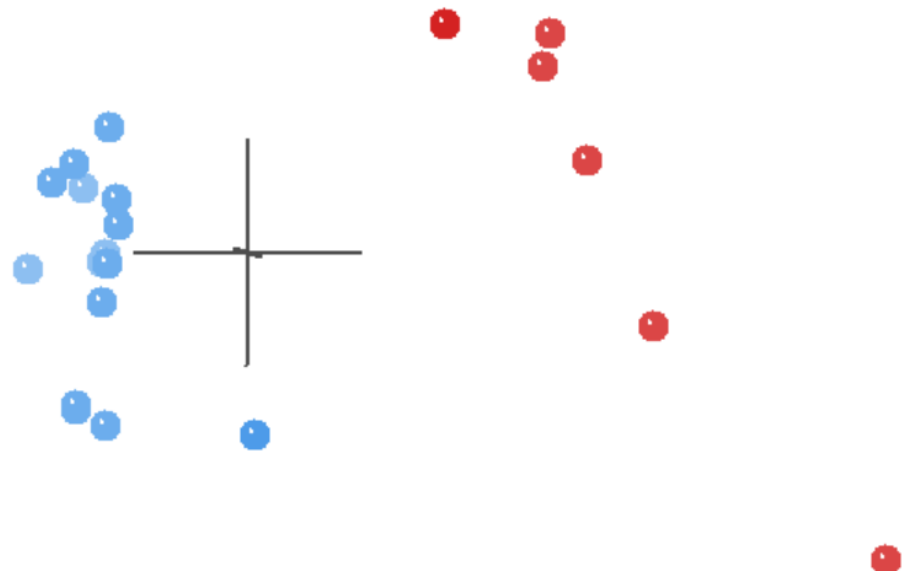
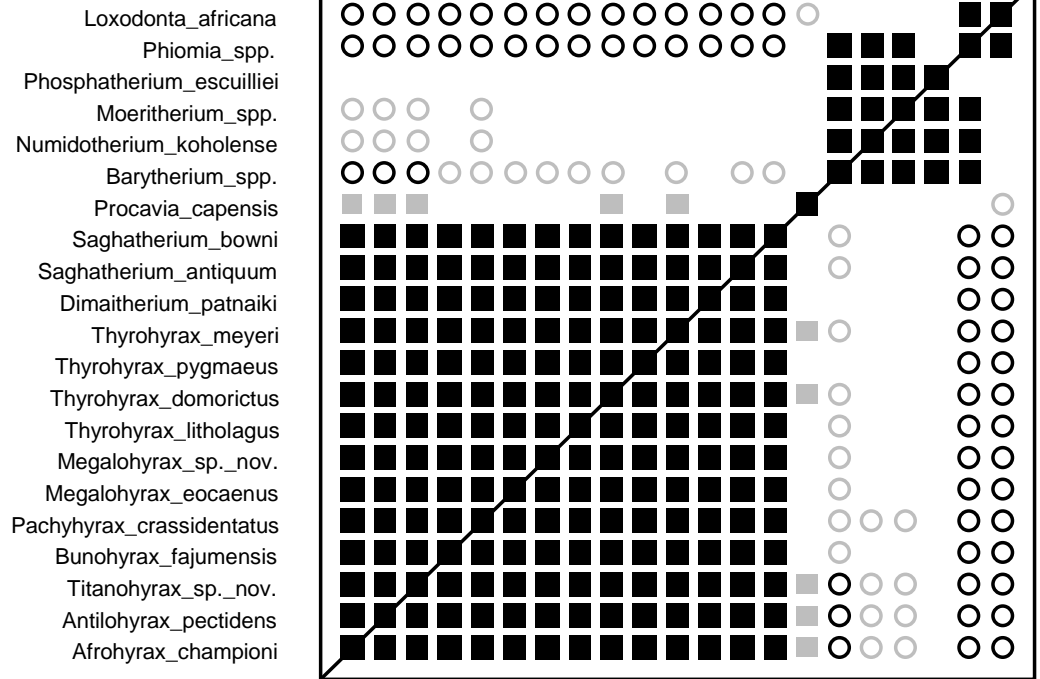
Order Uranotheria
Family Desmostylidae

| | |
|----------------------------------|------|
| Published taxa | 10 |
| Published characters | 37 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 9 |
| Characters used for calculations | 25 |
| Median bootstrap value | 83.5 |
| F ₉₀ | 0.28 |
| Stress of 3D MDS | 0.1 |
| k _{min} | 3 |
| Conclusion | HB? |

Notes: BDC results reveal three clusters: 1. the *Palaeoparadoxia* species, 2. the desmostylids *Vanderhoofius*, *Desmostylus*, and *Cornwallius*, and 3. the remaining taxa. There is significant, negative BDC between the restricted desmostylid group and the outgroup taxa. Bootstrap values for all BDC are poor. The MDS results reveal four clusters, separating the *Behemotops* species from the remaining outgroup taxa. We may provisionally accept the restricted Desmostylidae *sensu stricto* (*Vanderhoofius*, *Desmostylus*, and *Cornwallius*) as a holobaramin.

Seiffert, E.R., S. Nasir, A. Al-Harthy, J.R. Groenke, B.P. Kraatz, N.J. Stevens, and A.R. Al-Sayigh. 2012. Diversity in the later Paleogene proboscidean radiation: a small barytheriid from the Oligocene of Dhofar Governorate, Sultanate of Oman. *Naturwissenschaften* 99:133-141.

Characters: Craniodental, postcranial, soft tissue

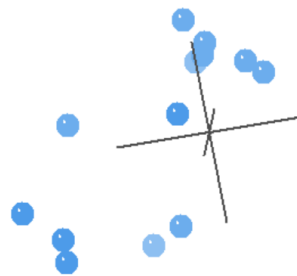
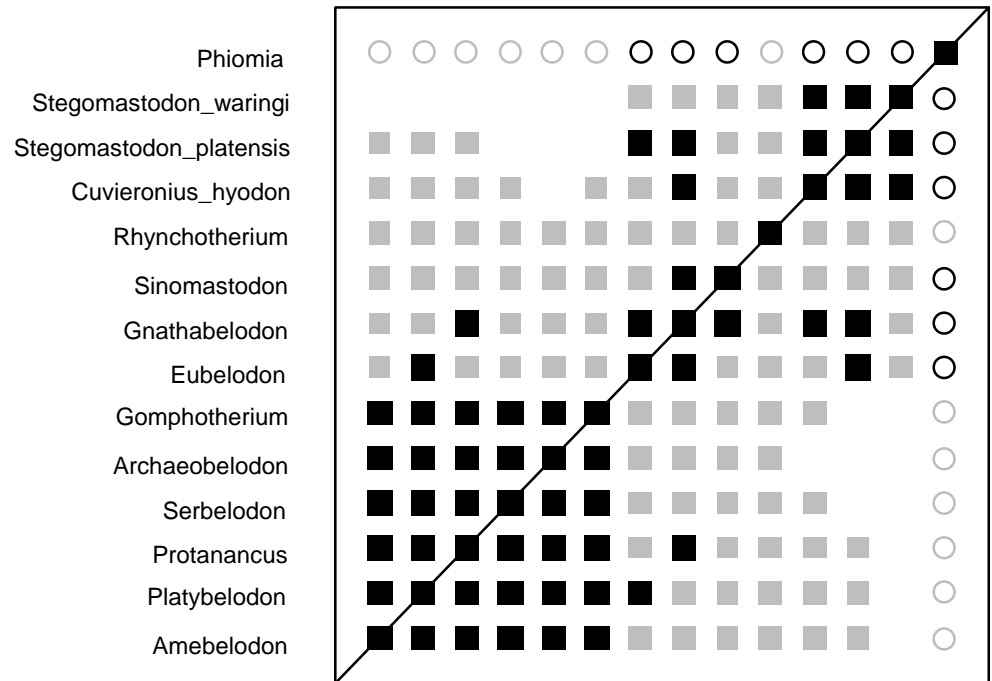


Order Hyracoidea
Family Procaviidae

| | |
|----------------------------------|------|
| Published taxa | 63 |
| Published characters | 422 |
| Character relevance | 0.75 |
| Taxic relevance | 0.4 |
| Taxa used for calculations | 21 |
| Characters used for calculations | 208 |
| Median bootstrap value | 100 |
| F ₉₀ | 0.75 |
| Stress of 3D MDS | 0.25 |
| k _{min} | 15 |
| Conclusion | HB |

Notes: Procaviidae is well separated from the outgroup taxa in BDC and MDS results. Procaviidae is likely a holobaramin.

Characters: Craniodental and postcranial



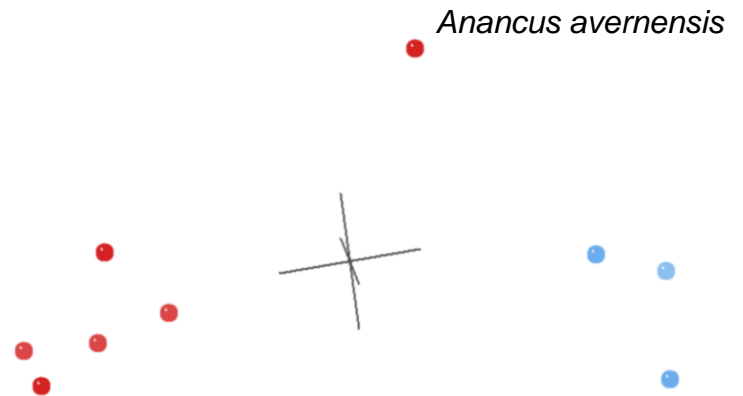
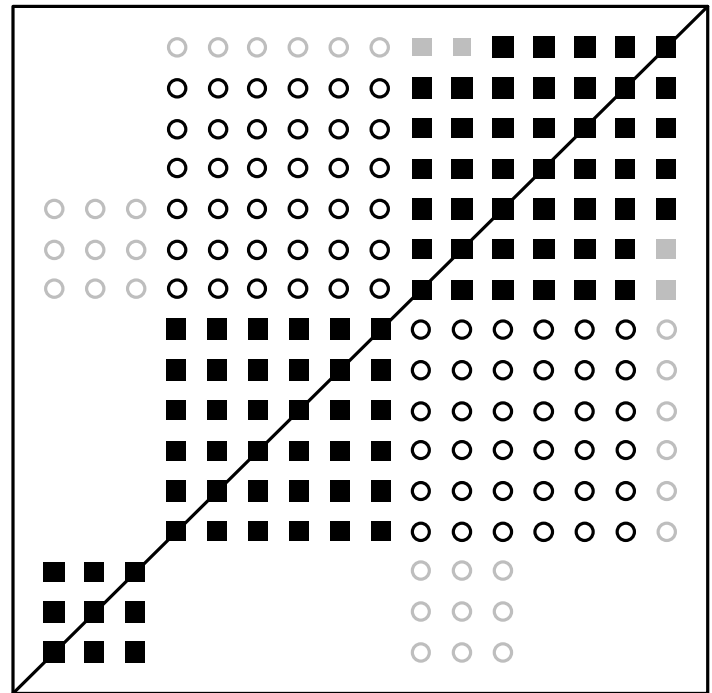
Order Proboscidea
Family Gomphotheriidae

| | |
|----------------------------------|------|
| Published taxa | 14 |
| Published characters | 43 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 14 |
| Characters used for calculations | 34 |
| Median bootstrap value | 85 |
| F ₉₀ | 0.33 |
| Stress of 3D MDS | 0.11 |
| k _{min} | 5 |
| Conclusion | HB? |

Notes: Gomphotheriidae is well separated from the outgroup taxa in BDC and MDS results. Gomphotheriidae is likely a holobaramin; although, the outgroup may not be suitable for these taxa.

Ferretti, M.P. and R. Debruyne. 2010. Anatomy and phylogenetic value of the mandibular and coronoid canals and their associated foramina in proboscideans (Mammalia). *Zoological Journal of the Linnean Society* 161:391-413.
 Characters: Cranial

Prodeinotherium_hobley
 Procavia_capensis
 Phophatherium_escuillei
 Phiomia_serridens
 Moeritherium_trigodon
 Numidotherium_koholense
 Barytherium_grave
 Mammuthus_primigenius
 Mammuthus_meridionalis
 Palaeoloxodon_mnaidriensis
 Loxodonta_africana
 Elephas_planifrons
 Elephas_maximus
 Mammut_americanum
 Gomphotherium_angustidens
 Anancus_avernensis



Order Proboscidea
 Family Elephantidae

| | |
|----------------------------------|------|
| Published taxa | 16 |
| Published characters | 10 |
| Character relevance | 0.75 |
| Taxic relevance | 0 |
| Taxa used for calculations | 16 |
| Characters used for calculations | 10 |
| Median bootstrap value | 97 |
| F ₉₀ | 0.61 |
| Stress of 3D MDS | 0.03 |
| k _{min} | 3 |
| Conclusion | HB? |

Notes: BDC and MDS reveal Elephantidae clearly separated from the outgroup taxa. Elephantidae is likely a holobaramin. The ten characters are not remotely holistic, so the elephantid holobaramin should be considered extremely provisional.