

1 **Running head:** CRANIAL MORPHOLOGICAL DISPARITY IN  
2 TENRECS

3 Quantifying cranial morphological  
4 disparity in tenrecs (Afrosoricida,  
5 Tenrecidae) with implications for their  
6 designation as an adaptive radiation

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14 golden moles, adaptive radiation

## 15 Abstract

16 Understanding why some clades are more phenotypically diverse than  
17 others remains a central challenge in evolutionary biology. This issue is  
18 particularly relevant when we consider whether a group represents an  
19 adaptive radiation. However, we must be able to identify exceptionally  
20 diverse clades before we can determine the selective pressures which led  
21 to the evolution of their variety. Tenrecs (Afrosoricida, Tenrecidae) are a  
22 family of small mammals and are often cited as an example of a  
23 phenotypically diverse, adaptively radiated group. However, this  
24 assumption has not been tested. Here we use geometric morphometric  
25 analyses of cranial and mandible shape to test whether tenrecs show  
26 exceptional morphological disparity. We find that tenrecs are no more  
27 morphologically diverse than their sister taxa, the golden moles  
28 (Afrosoricida, Chrysochloridae), casting doubt over whether tenrecs  
29 should be considered to be an exceptionally diverse group.

## 30 Introduction

31 Phenotypically diverse groups have long attracted the attention of  
32 evolutionary biologists, particularly when it comes to the study of  
33 adaptive radiations - 'evolutionary divergence of members of a single  
34 phylogenetic lineage into a variety of different adaptive forms' (Futuyma  
35 1998, cited by Losos, 2010).

36 There are many famous examples of adaptive radiations including  
37 Darwin's finches, Caribbean *Anolis* lizards and cichlid fish (Gavrilets &  
38 Losos, 2009). However, there has been considerable debate about how  
39 adaptive radiations should be defined (Glor, 2010; Losos & Mahler, 2010)  
40 based on the relative importance of speciation rates, species richness and  
41 morphological diversity. One particular issue is whether it is even  
42 meaningful to distinguish a specific group of species as an adaptive  
43 radiation or not based on arbitrary statistical thresholds of variety (Olson  
44 & Arroyo-Santos, 2009).

45 Despite the controversies and disagreements, there does seem to be a  
46 consensus that high morphological diversity is an important criterion for  
47 identifying adaptive radiations (Losos & Mahler, 2010; Olson &  
48 Arroyo-Santos, 2009). One way to test whether a group shows high  
49 morphological diversity is through sister taxa comparisons. For example,  
50 Losos and Miles (2002) used this approach to demonstrate exceptional  
51 diversity in some but not all clades of iguanid lizards. Here we use  
52 sister-taxa comparisons to test whether tenrecs (Afrosoricida, Tenrecidae)  
53 exhibit the high levels of phenotypic diversity that are expected of an  
54 adaptively radiated clade.

55 The tenrec family contains 34 species, 31 of which are endemic to

56 Madagascar (Olson, 2013). Tenrecs are often cited as an example of an  
57 adaptively radiated family which exhibits exceptional morphological  
58 diversity (Soarimalala & Goodman, 2011; Olson & Goodman, 2003). For  
59 example, there are tenrecs which convergently resemble shrews (*Microgale*  
60 tenrecs), moles (*Oryzorictes* tenrecs) and hedgehogs (*Echinops* and *Setifer*  
61 tenrecs) (Eisenberg & Gould, 1969) even though they are not closely  
62 related to these species (Stanhope et al., 1998).

63 However, evidence for claim that tenrecs are exceptionally diverse has  
64 not been tested. Here we present the first quantitative investigation of  
65 morphological diversity in tenrecs, and how this compares to their closest  
66 relatives, the golden moles (Afrosoricida, Chryscholoridae). We apply two  
67 dimensional geometric morphometric techniques (Rohlf & Marcus, 1993;  
68 Adams et al., 2013) to create morphospace plots that depict cranial and  
69 mandible morphological variation in the two Families. We use these  
70 morphospaces to compare the relative morphological disparity (Foote,  
71 1997; Wills et al., 1994; Erwin, 2007) within each Family.

72 Our results show an overall trend for higher morphological diversity  
73 in tenrec crania compared to those of golden moles. However, these  
74 differences are not statistically significant. These findings indicate that,  
75 with regards to cranial shape, tenrecs are no more morphologically  
76 diverse than their closest relatives.

77 In contrast, we found significantly greater morphological disparity in  
78 golden mole mandibles compared to tenrecs. These findings cast doubt  
79 over whether the apparent phenotypic diversity within tenrecs should be  
80 considered to be truly exceptional.

## 81 **Materials and Methods**

### 82 **Morphological data collection**

83 One of us (SF) photographed cranial specimens of tenrecs and golden  
84 moles at the Natural History Museum London (BMNH), the Smithsonian  
85 Institute Natural History Museum (SI), the American Museum of Natural  
86 History (AMNH), Harvard's Museum of Comparative Zoology (MCZ)  
87 and the Field Museum of Natural History, Chicago (FMNH). We  
88 photographed the specimens with a Canon EOS 650D camera fitted with  
89 an EF 100mm f/2.8 Macro USM lens using a standardised procedure to  
90 minimise potential error (see supplementary material for details).

91 We collected pictures of the skulls in dorsal, ventral and lateral views  
92 (right side of the skull) and of the outer (buccal) side of the right  
93 mandibles. A full list of museum accession numbers and details on how  
94 to access the images can be found in the supplementary material.

95 In total we collected pictures from 182 skulls in dorsal view (148  
96 tenrecs and 34 golden moles), 173 skulls in ventral view (141 tenrecs and  
97 32 golden moles), 171 skulls in lateral view (140 tenrecs and 31 golden  
98 moles) and 182 mandibles in lateral view (147 tenrecs and 35 golden  
99 moles), representing 31 species of tenrec (out of the total 34 in the family)  
100 and 12 species of golden moles (out of a total of 21 in the family (Asher  
101 et al., 2010)). We used the taxonomy of Wilson and Reeder (2005)  
102 supplemented with more recent sources (IUCN, 2012; Olson, 2013) to  
103 identify our specimens.

104 We used a combination of both landmarks (type 2 and type 3,  
105 (Zelditch et al., 2012)) and semilandmarks to characterise the shapes of

106 our specimens. Figure 1 shows our landmarks (points) and  
107 semilandmarks (outline curves) for the skulls in dorsal and ventral views  
108 and figure 2 shows the points and curves we used for lateral views of  
109 skulls and mandibles. Corresponding definitions of each of the landmarks  
110 can be found in the supplementary material.

111 We digitised all landmarks and semilandmarks in tpsDIG, version 2.17  
112 (Rohlf, 2013). We re-sampled the outlines to the minimum number of  
113 evenly spaced semilandmark points required to represent each outline  
114 accurately (MacLeod, 2013, details in supplementary material). We used  
115 TPSUtil (Rohlf, 2012) to create sliders files (Zelditch et al., 2012) to define  
116 which points were semilandmarks. We conducted all subsequent analyses  
117 in R version 3.0.2 (R Core Team, 2014) within the geomorph package  
118 (Adams et al., 2013). We used the gpagen function to run a general  
119 Procrustes alignment (Rohlf & Marcus, 1993) of the landmark coordinates  
120 while sliding the semilandmarks by minimising Procrustes distance  
121 (Bookstein, 1997). We used these Procrustes-aligned coordinates of all  
122 species to calculate average shape values for each species ( $n = 43$ ) which  
123 we then used for a principal components (PC) analysis with the  
124 plotTangentSpace function (Adams et al., 2013).

## 125 **Disparity calculations**

126 We calculated morphological disparity separately for golden moles and  
127 tenrecs in each of the morphological datasets. We used the PC axes which  
128 accounted for 95% of the cumulative variation to calculate four disparity  
129 metrics; 1) the sum of the range, 2) the product of the range, 3) the sum of  
130 the variance and 4) the product of the variance of morphospace occupied

131 by each Family (Brusatte et al., 2008; Foth et al., 2012; Ruta et al., 2013).

132 We used two approaches to test whether tenrecs have significantly  
133 different morphologies compared to golden moles. First we compared  
134 morphospace occupation between the two groups with non parametric  
135 MANOVAs (Anderson, 2001) to test whether tenrecs and golden moles  
136 occupy significantly different areas of morphospace (e.g Serb et al., 2011;  
137 Ruta et al., 2013).

138 Secondly, we used pairwise permutation tests to test the null  
139 hypothesis that tenrecs and golden moles have equal disparity. If this  
140 hypothesis was true then the designation of each species as belonging to  
141 either tenrecs or golden moles should be arbitrary. Therefore we  
142 permuted the data by assigning Family identities at random to each  
143 specimen and calculated the differences in disparity for each of the new  
144 Family groupings. We repeated these permutations 1000 times to generate  
145 a null distribution of the expected differences in Family disparity. We  
146 compared our observed (true) measures of the differences in disparity  
147 between tenrecs and golden moles to these permuted distributions to test  
148 whether the families had significantly different levels of disparity.

149 The majority of tenrec species (19 out of 31 in our dataset) are  
150 members of the *Microgale* (shrew-like) Genus which is notable for its  
151 relatively low phenotypic diversity (Soarimalala & Goodman, 2011;  
152 Jenkins, 2003). The strong similarities among these species may mask  
153 signals of higher disparity among other tenrecs. Therefore we repeated  
154 our Family-level comparisons of disparity excluding the *Microgale* species  
155 so that we could compare disparity within the remaining 12 tenrec species  
156 to disparity within the 12 species of golden moles.

## Results

### Morphological disparity in tenrecs and golden moles

Figure 3 depicts the morphospace plots derived from our principal components analyses of average Procrustes-superimposed shape coordinates for each species in our skull and mandible data respectively. We used the principal components axes which accounted for 95% of the cumulative variation (number of axes:  $n = 7$  (dorsal),  $n = 8$  (ventral),  $n = 8$  (lateral) and  $n = 12$  (mandibles)) to calculate the disparity of each Family.

Tenrecs and golden moles clearly have very different cranial and mandible morphologies: in each analysis, the families occupy significantly different areas of morphospace (npMANOVA, table 3). Our comparisons of disparity within each Family yielded different trends for skulls compared to mandibles. In our analyses of the three different views of the skulls, there is an overall trend for tenrecs to have higher disparity than golden moles. However, none of these differences are statistically significant (table 1).

There is a less clear pattern from our analysis of disparity in mandibles. Three of our four metrics indicate that golden moles have significantly higher disparity in the shape of their mandibles than tenrecs (table 1) although one metric (sum of ranges) indicated the opposite result.

The three curves at the back of the mandibles (figure 2) place a particular emphasis on shape variation in the posterior of the bone; the ramus, coronoid, condylar and angular processes. Therefore, higher disparity in golden mole mandibles compared to tenrecs could be driven by greater morphological variation in these structures. To test this idea,



182 we repeated our morphometric analyses of the mandibles with a reduced  
183 dataset of points; just the seven landmark points and one single curve at  
184 the base of the jaw between landmarks 1 and 7 (figure 2). When we  
185 compared disparity with this reduced data set we found that golden  
186 moles no longer had significantly higher disparity than tenrecs (table 1).

## 187 **Morphological disparity in non-*Microgale* tenrecs and** 188 **golden moles**

189 We repeated our disparity comparisons with a subset of the tenrec  
190 specimens to remove the large and phenotypically similar *Microgale* tenrec  
191 Genus. In this case we found that tenrecs have significantly higher  
192 disparity than golden moles when the skulls are analysed in lateral view  
193 (table 2). However, none of the other comparisons in any of the analyses  
194 were significant.

## 195 **Discussion**

196 Our analyses are the first quantitative investigation of morphological  
197 disparity in tenrecs. Our results suggest that phenotypic variation in  
198 tenrecs is not as exceptional as it first appears.

199 When we compared tenrec cranial morphologies to their closest  
200 relatives we found a trend towards higher disparity in tenrecs compared  
201 to golden moles but none of these differences were significant. Even when  
202 we removed the phenotypically similar *Microgale* Genus, tenrecs were still  
203 no more diverse than golden moles in most of the analyses (table 2). In  
204 contrast, our mandible analyses showed that golden moles have more

205 disparate mandibles than tenrecs seemingly due to greater diversity  
206 within their posterior-mandible shapes.

207 It is evident that tenrecs are a diverse group, both phenotypically and  
208 ecologically. Body sizes of extant tenrecs span three orders of magnitude  
209 (2.5 to  $> 2,000\text{g}$ ) which is a greater range than all other Families, and most  
210 Orders, of living mammals (Olson & Goodman, 2003). Within this vast  
211 size range there is striking phenotypic diversity from the spiny *Echinops*,  
212 *Setifer* and *Hemicentetes* to the mole-like *Oryzorictes* and shrew-like  
213 *Microgale*. These diverse forms inhabit a wide variety of ecological niches  
214 and habitats including terrestrial, arboreal, semi-aquatic and  
215 semi-fossorial environments (Soarimalala & Goodman, 2011). In contrast,  
216 although golden moles occupy a wide altitudinal, climatic and  
217 vegetational spectrum of habitats (Bronner, 1995), they are all fossorial  
218 species which, superficially at least, appear to be less phenotypically  
219 diverse than tenrecs.

220 There is a danger when using sister taxa comparisons that a clade's  
221 diversity will be judged to be exceptional just because it is more variable  
222 than an exceptionally non-diverse sister taxon (Losos & Miles, 2002).  
223 However, we compared an apparently phenotypically diverse clade to a  
224 more uniform sister clade yet our results do not indicate that tenrecs are  
225 more morphologically diverse than their closest relatives (table 1). These  
226 unexpected findings highlight the importance of testing our assumptions  
227 about patterns of morphological variety.

228 One apparent anomaly in our results is that we found opposite  
229 patterns for disparity among tenrecs and golden moles in the analyses of  
230 skulls and mandibles. Our landmarks and curves for the mandibles

(figure 2) include aspects of variation in the dentition but they focus particular attention on the ascending ramus (condyloid, condylar and angular processes). Therefore higher disparity in golden moles could reflect greater morphological variability in these posterior mandible structures. To test this idea we deleted the three semi-landmark curves around these structures and repeated our disparity analyses of mandibles using seven landmarks and just one curve at the base of the jaw. In this case we retrieved the opposite pattern: tenrecs had higher morphological disparity than golden moles but not significantly (see supplementary material). Therefore, our results indicate that golden moles have greater morphological variation in the posterior structures of their mandibles compared to tenrecs.

It proved impossible to position reliable landmarks on the corresponding mandibular articulation areas of the skull in lateral view (see supplementary). Therefore we could not test whether higher morphological disparity in the rami were correlated with associated morphological variety in the articulation areas of the skull.

We focused on variation in cranial morphology which is commonly used to study phenotypic variation within species (Blagojević & Milošević-Zlatanović, 2011; Bornholdt et al., 2008), to delineate species boundaries within a clade (e.g. Panchetti et al., 2008) or for cross-taxonomic comparative studies of phenotypic (dis)similarities (e.g. Ruta et al., 2013; Goswami et al., 2011; Wroe & Milne, 2007). However, studies of morphological disparity are inevitably constrained to measure diversity within specific traits rather than overall phenotypes (Roy & Foote, 1997). Disparity calculations based on skull shape can yield similar results compared to analyses of whole-skeleton discrete characters and

limb proportion data sets (Foth et al., 2012). However, we would need to extend our analyses to other morphological proxies of phenotype to test whether the cranial morphological disparity patterns presented here are indicative of overall differences in phenotypic diversity in tenrecs and golden moles

Evidence of exceptional morphological diversity is one criterion for designating a clade as an adaptive radiation (Losos & Mahler, 2010) and our analyses are the first measures of morphological diversity within tenrecs, a group which is commonly cited as an example of an adaptive radiation (Olson, 2013). However, describing phenotypic divergence as the product of an adaptive radiation requires exceptional morphological diversity in traits which have specific and proven adaptive significance (Losos & Mahler, 2010).

The evolution of cranial shape (both upper skull and mandible), particularly dental morphology, has obvious correlations with dietary specialisations and occupation of specific ecological niches (e.g. Wroe & Milne, 2007). Considering the wide ecological diversity of our study species; semi-fossorial, arboreal, terrestrial and semi-aquatic (Soarimalala & Goodman, 2011), we think that it is reasonable to expect that this variety should be reflected in skull morphology.

However, we have not included any measures of the ‘adaptiveness’ of cranial shape in our analyses and therefore our analyses should not be considered to be an explicit test of whether or not tenrecs are an adaptive radiation (Losos & Mahler, 2010). Instead we have made the first step towards understanding the apparent phenotypic diversity within tenrecs within a quantitative framework. Future work should focus on explicit

284 measures of the ‘adaptiveness’ and functional importance of tenrec cranial  
285 and post-cranial morphologies to understand the significance of  
286 morphological diversity within the Family (e.g. Mahler et al., 2010).

287 We have presented the first quantitative study which tests the common  
288 claim that tenrecs are an exceptionally diverse group (Olson, 2013;  
289 Soarimalala & Goodman, 2011; Eisenberg & Gould, 1969). Focusing on  
290 cranial diversity is only one aspect of morphological variation and further  
291 analyses are required to test whether other morphological traits yield  
292 similar patterns. However, our current results provide a clear indication  
293 that phenotypic variety within tenrecs is perhaps not as exceptional as it  
294 first seems.

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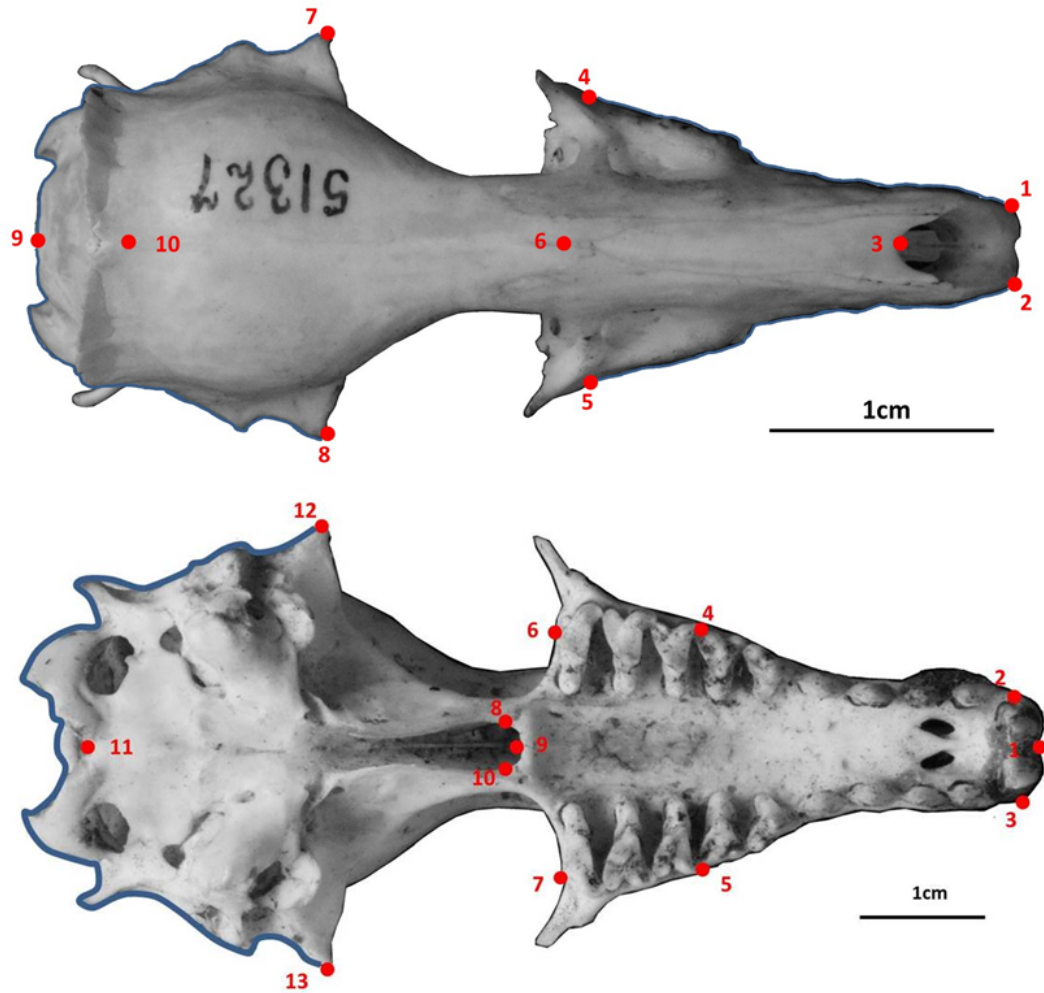


Figure 1: Landmarks (red points) and curves (blue lines) used to capture the morphological shape of skulls in dorsal and ventral views respectively. Curves were re-sampled to the same number of evenly-spaced points. See Supplementary Material for descriptions of the curves and landmarks. The specimens belong to two different *Potamogale velox* (Tenrecidae) skulls: accession number AMNH 51327 (dorsal) and NHML 1934.6.16.2 (ventral)

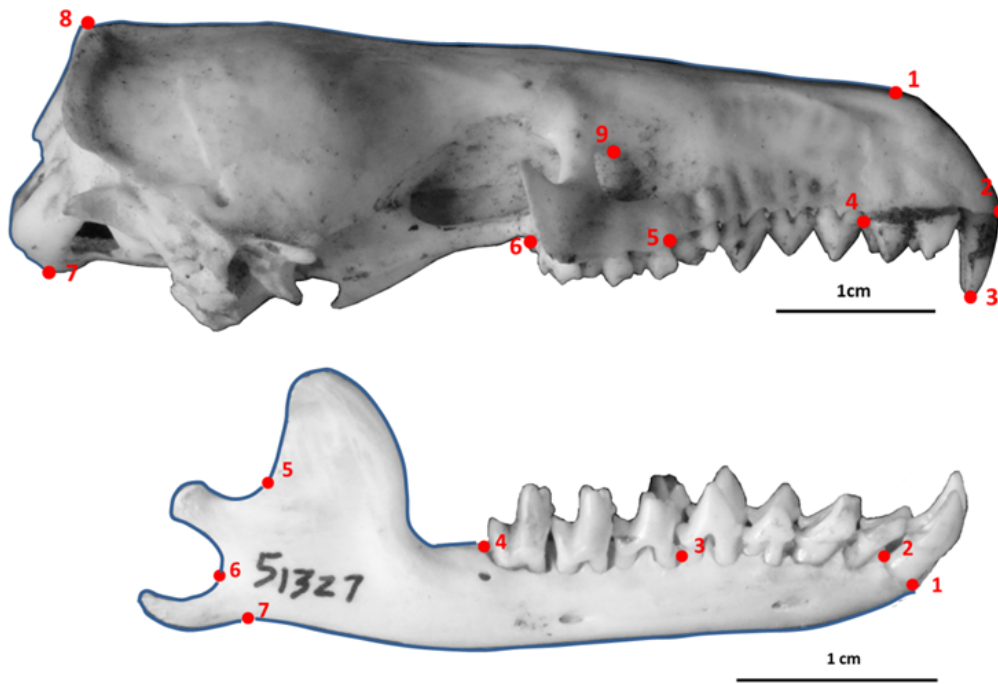


Figure 2: Landmarks (red points) and curves (blue lines) used to capture the morphological shape of lateral views of skulls and mandibles respectively. Curves were re-sampled to the same number of evenly-spaced points. See Supplementary Material for descriptions of the curves and landmarks. The specimens belong to two different *Potamogale velox* (Tenrecidae) skulls: accession number AMNH 51327 (dorsal) and NHML 1934.6.16.2 (ventral)

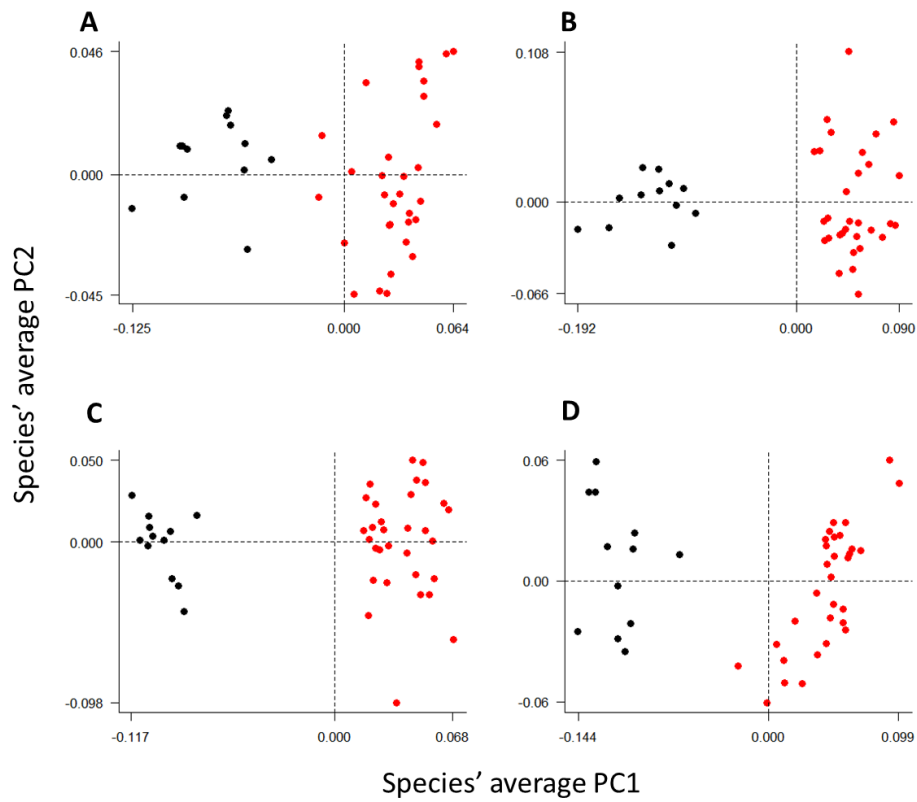


Figure 3: Principal components plots of the morphospaces occupied by tenrecs (red,  $n = 31$  species) and golden moles (black,  $n = 12$ ) for the skulls: dorsal (A), ventral (B), lateral (C) and mandibles (D) analyses. Axes are PC1 and PC2 of the average scores from a PCA analysis of mean Procrustes shape coordinates for each species.

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Table 1: Disparity comparisons between tenrecs (T) and golden moles (G) for each of our data sets(rows) and four disparity metrics (columns). ‘Mandibles:one curve’ refers to our shape analysis of mandibles excluding the three curves around the posterior structures of the jaw (figure 2). Significant differences are highlighted in bold with the corresponding p value in brackets. Disparity metrics are: sum of variance, product of variance, sum of ranges and product of ranges

<b>Disparity metric</b>	<b>SumVar</b>	<b>ProdVar</b>	<b>SumRange</b>	<b>ProdRange</b>
Skulls dorsal	T>G	T>G	T>G	T>G
Skulls lateral	T>G	T>G	T>G	T>G
Skulls ventral	T>G	G>T	T>G	T>G
Mandibles	G>T	<b>G&gt;T* (0.008)</b>	<b>T&gt;G* (0.025)</b>	<b>T&gt;G* (0.009)</b>
Mandibles	G>T	G>T	T>G	T>G



Table 2: Disparity comparisons between non-*Microgale* tenrecs (T) and golden moles (G) for each of our data sets(rows) and four disparity metrics (columns). Significant differences are highlighted in bold with the corresponding p value in brackets. Disparity metrics are; sum of variance, product of variance, sum of ranges and product of ranges.

<b>Disparity metric</b>	<b>SumVar</b>	<b>ProdVar</b>	<b>SumRange</b>	<b>ProdRange</b>
Skulls dorsal	T>G	T>G	T>G	T>G
Skulls lateral	<b>T&gt;G* (0.014)</b>	T>G	<b>T&gt;G* (0.001)</b>	<b>T&gt;G*(0.003)</b>
Skulls ventral	T>G	T>G	T>G	T>G
Mandibles	T>G	G>T	T>G	G>T

Table 3: npMANOVA comparisons of morphospace occupation for tenrecs and golden moles in each of the four analyses (three views of skulls and mandibles). In each case the two families occupy significantly different areas of morphospace.

<b>Analysis</b>	<b>F</b>	<b>R<sup>2</sup></b>	<b>p value</b>
Skulls dorsal	66.02	0.62	0.001
Skulls ventral	100.74	0.71	0.001
Skulls lateral	75.07	0.65	0.001
Mandibles	59.34	0.59	0.001