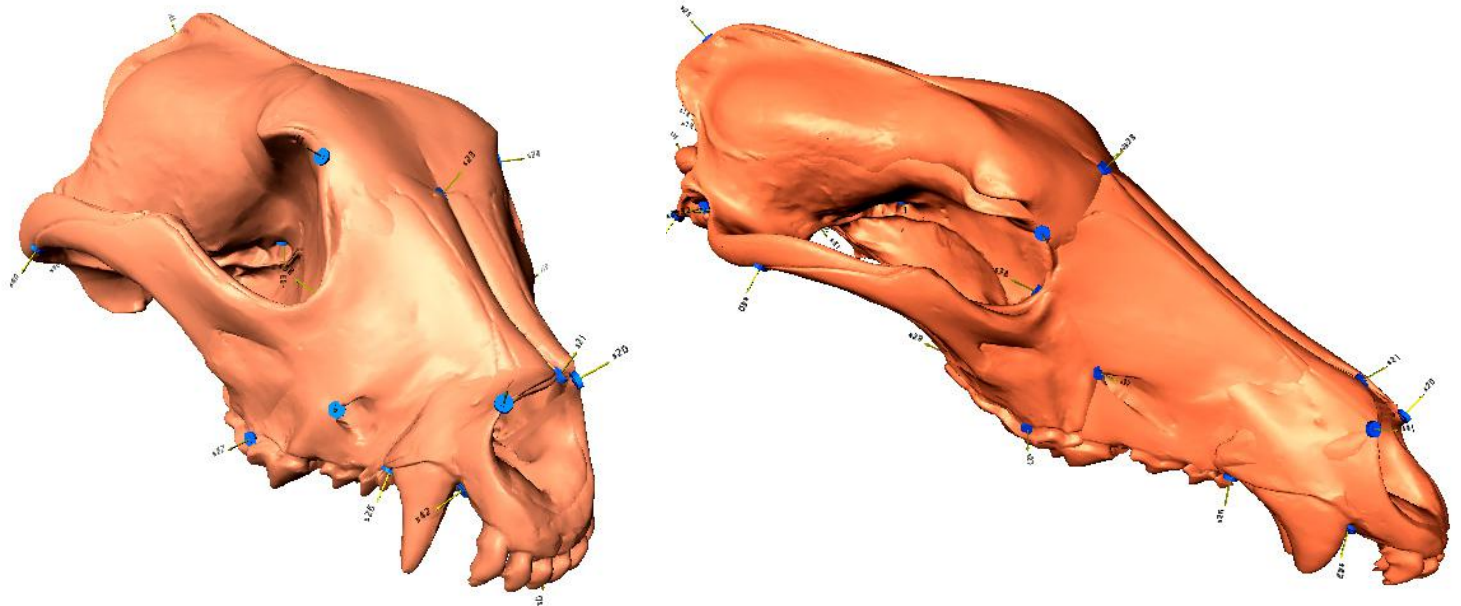


discovering a new rule in the evolution of mammals



UNIVERSITÀ DEGLI STUDI
DI MODENA E REGGIO EMILIA

Andrea Cardini
UNIMORE & UWA



cardini@unimo.it, alcardini@gmail.com

<https://sites.google.com/site/alcardini/home/main>

in mammalian morphological evolution

what's a rule?

trend in change among closely related species

e.g., Bergmann's rule

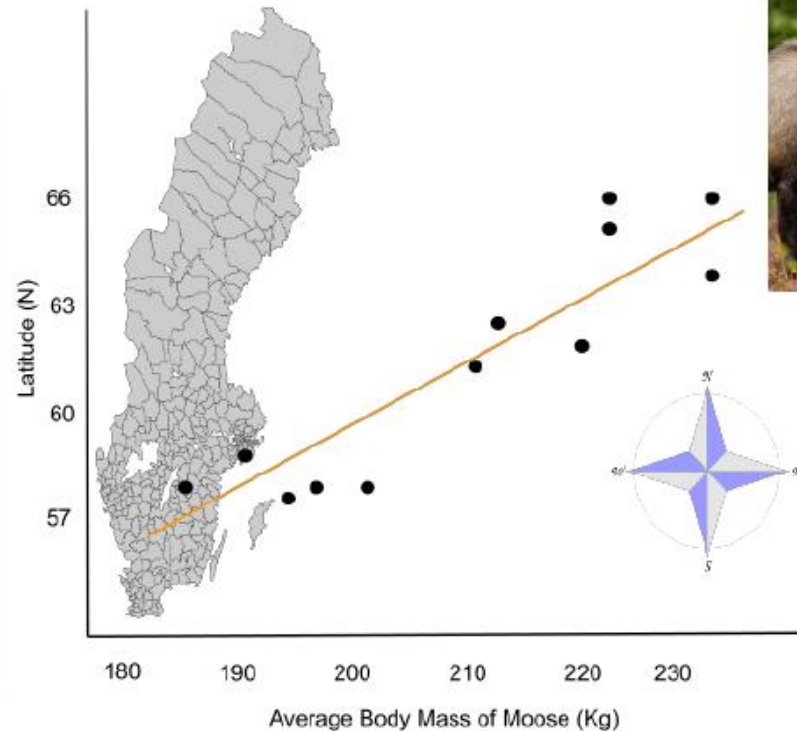


WIKIPEDIA
The Free Encyclopedia

- Main page
- Contents
- Featured content
- Current events
- Random article
- Donate to Wikipedia
- Wikipedia store

Interaction

- Help
- About Wikipedia
- Community portal
- Recent changes
- Contact page



Bergmann's rule is an **ecogeographical rule** that states that within a broadly distributed taxonomic **clade**, populations and species of larger size are found in colder environments, and species of smaller size are found in warmer regions. Although originally formulated in terms of species within a genus, it has often been recast in terms of populations within a species. It is also often cast in terms of latitude.

(general trend; not a strict rule; plenty of exceptions)

'my' rule is about evolutionary allometry

size-related shape changes among species

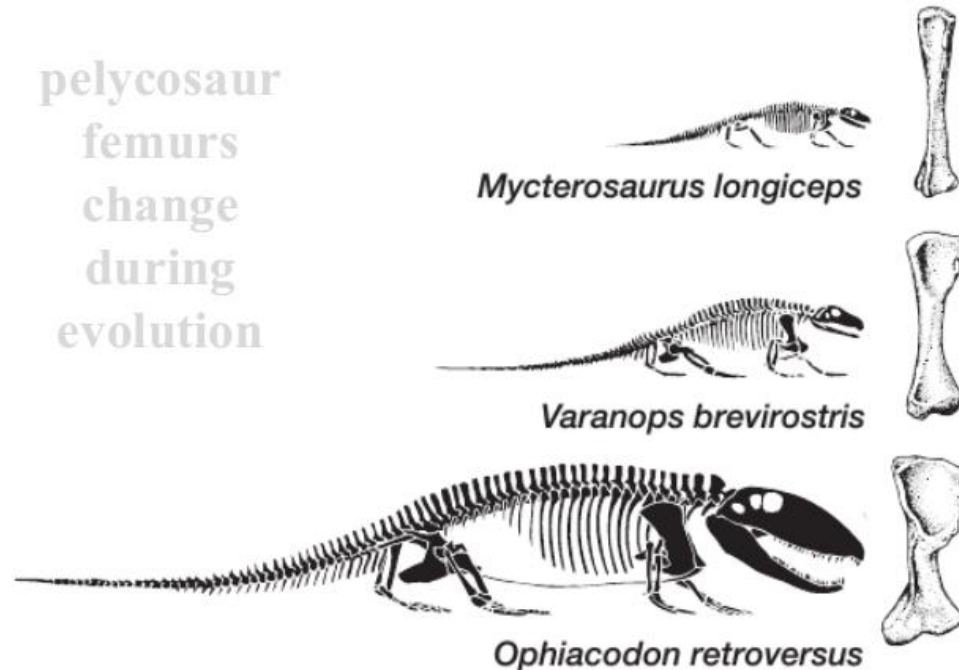
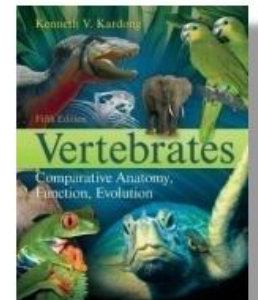
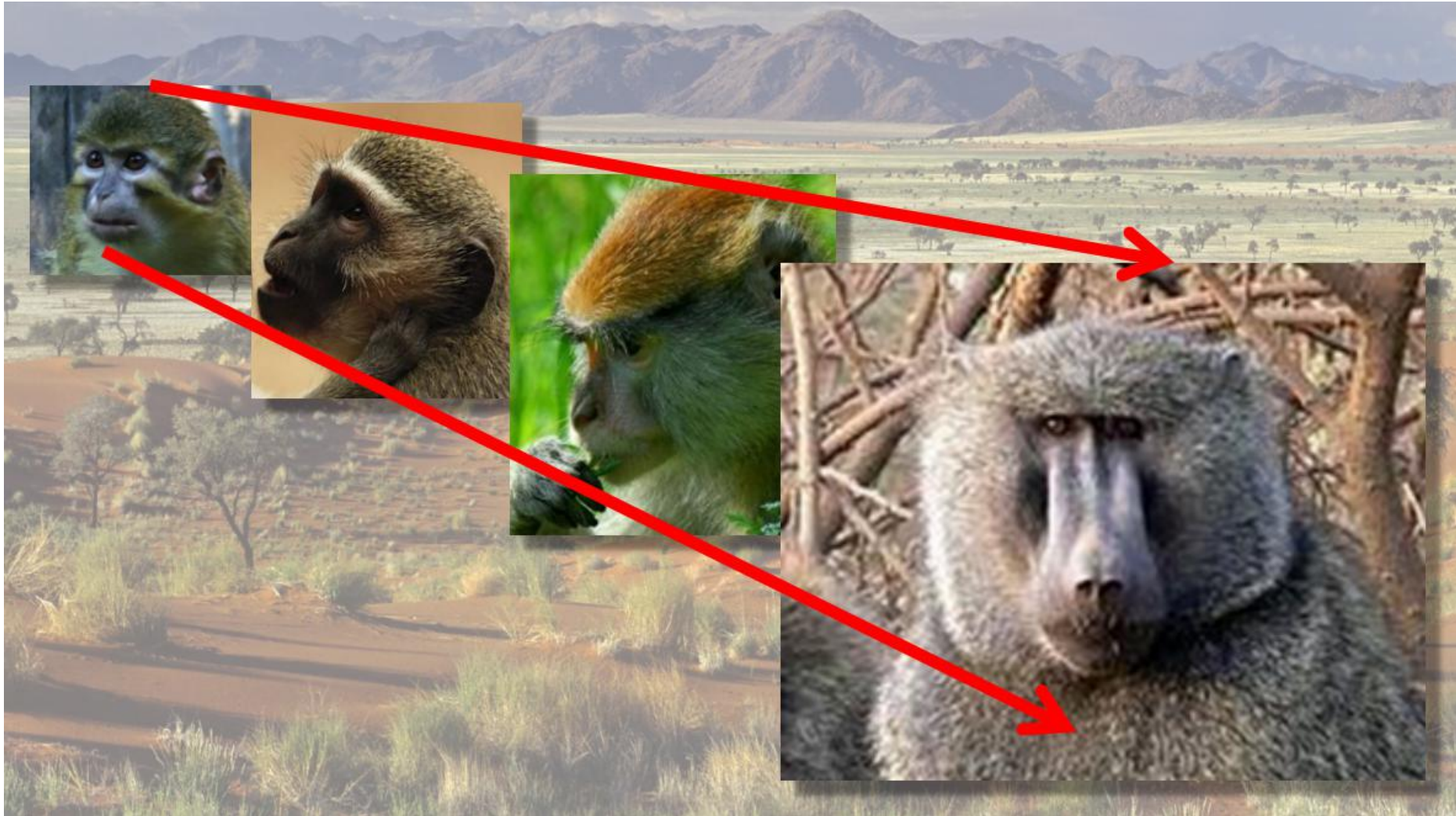


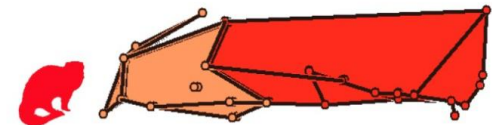
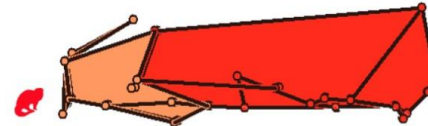
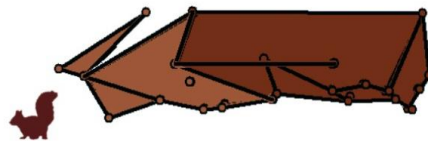
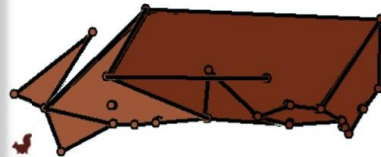
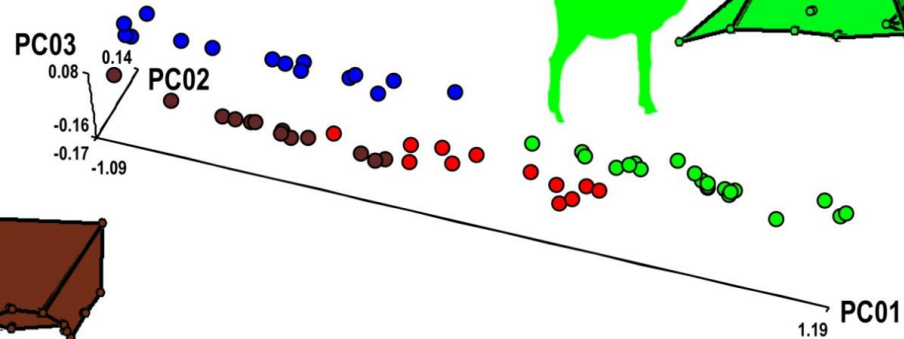
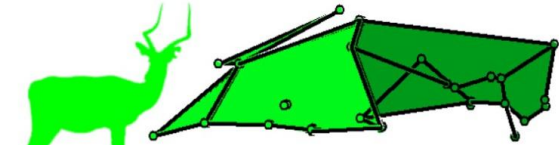
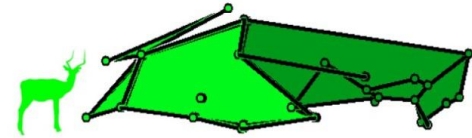
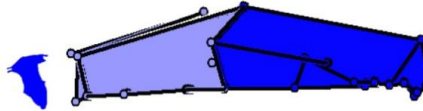
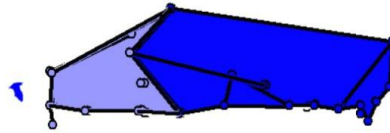
FIGURE 4.8 Body size and limb design in pelycosaurs. Relative sizes of three pelycosaur species are illustrated. The femurs of each, drawn to the same length, are shown to the right of each species. The larger pelycosaur carries a relatively larger mass, and its more robust femur reflects this supportive demand.



'my' rule is about **evolutionary allometry**
of crania (averaged across adults within each species)



1st we showed it in 4 placental orders* (SYNTHESYS 2007)



ARTICLE

Received 1 Feb 2013 | Accepted 19 Aug 2013 | Published 18 Sep 2013

DOI: 10.1038/ncomms3488

Larger mammals have longer faces because of size-related constraints on skull form

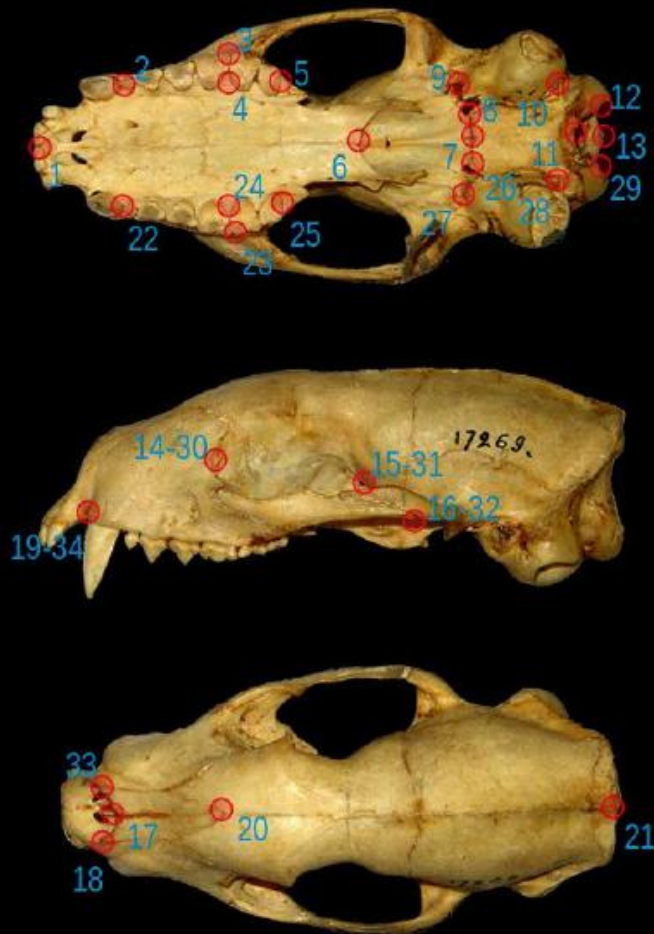
Andrea Cardini^{1,2,3,4} & P. David Polly⁵

Facial length is one of the best known examples of heterochrony. Changes in the timing of facial growth have been invoked as a mechanism for the origin of our short human face from our long-faced extinct relatives. Such heterochronic changes arguably permit great evolutionary flexibility, allowing the mammalian face to be remodelled simply by modifying postnatal growth. Here we present new data that show that this mechanism is significantly constrained by adult size. Small mammals are more brachycephalic (short faced) than large ones, despite the putative independence between adult size and facial length. This pattern holds across four phenotypic lineages: antelopes, fruit bats, tree squirrels and mongooses. Despite the apparent flexibility of facial heterochrony, growth of the face is linked to absolute size and introduces what seems to be a loose but clade-wide mammalian constraint on head shape.

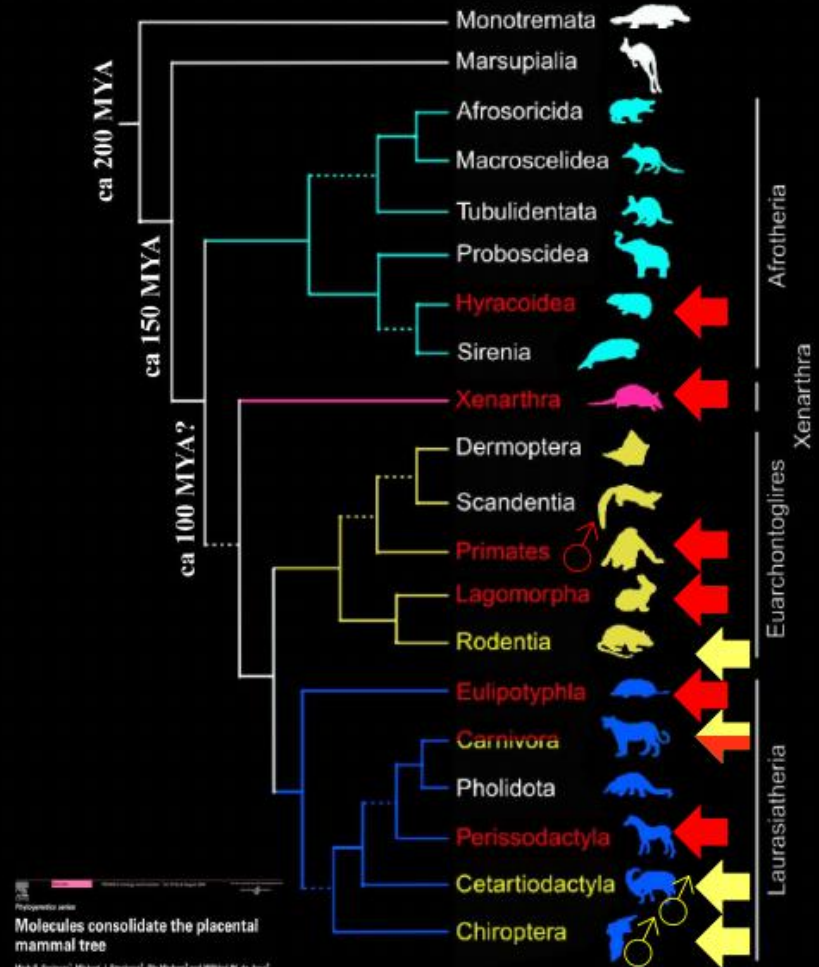
*ca subfam./tribe
representative
of an order

now hoping to show it in several more (SYNTHESESYS 2015 + 2007 + Leverhulme 2004) samples

3D configuration



new & old – $N_{\text{total}} = 6483$



and it looks promising!

allometric analyses
RESULTS?

e.g.,

$N_{\text{sp.}} = 13$

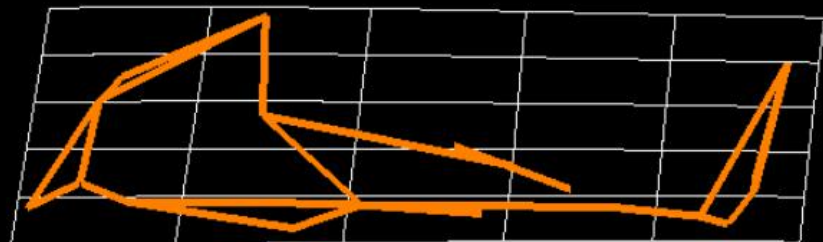
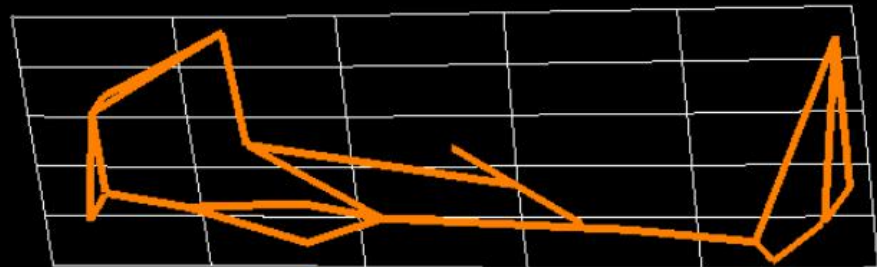
size max/min = 2.3

R^2 (allom. var.) = 16%




visualization ca. x4

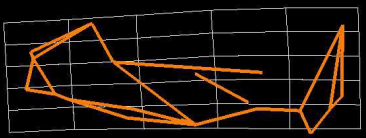
(allometric predictions min-max
in side view – wireframe/TPS)





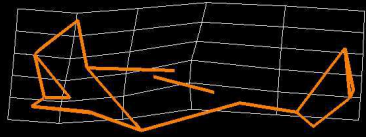
hyraxes

  
 N_{sp} max/min R2
 4 1.3 36%



sloths? (NB above family level!)

  
 N_{sp} max/min R2
 5 1.5 87%



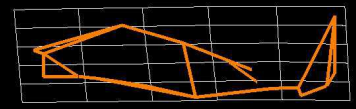
armadillos

  
 N_{sp} max/min R2
 16 5.3 24%



Dasypus (subfam. with min. allom.)




  
 N_{sp} max/min R2
 5 1.8 17%

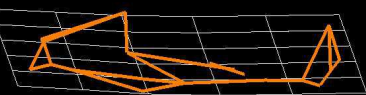


x8




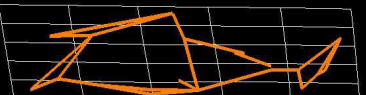
canids

  
 N_{sp} max/min R2
 31 2.7 22%





equids

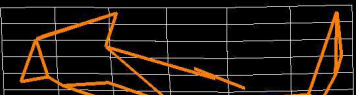
  
 N_{sp} max/min R2
 7 1.3 25%



x8

Erinaceinae

  
 N_{sp} max/min R2
 11 1.4 23%



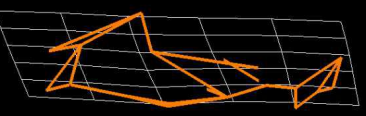
herpestids

  
 N_{sp} max/min R2
 13 2.3 16%



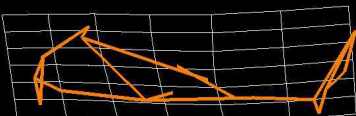
Antilopinae-Cephalophinae + melopes

  
 N_{sp} max/min R2
 22 2.6 17%





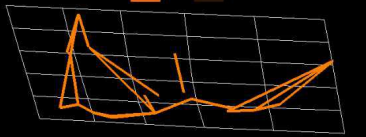
African Pteropodidae

  
 N_{sp} max/min R2
 15 2.9 32%



cercopithecini

  
 N_{sp} max/min R2
 21 1.9 64%



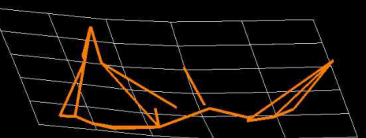
leporids

  
 N_{sp} max/min R2
 32 1.5 24%



papionini

  
 N_{sp} max/min R2
 33 2.0 60%



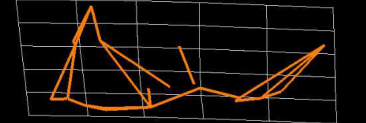
Protoxerini squirrels

  
 N_{sp} max/min R2
 12 2.3 39%





African colobines

  
 N_{sp} max/min R2
 19 1.4 30%





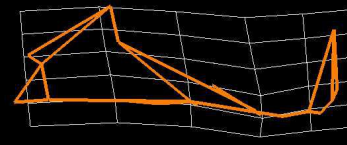
hyraxes

  
 N_{sp} max/min R2
 4 1.3 36%



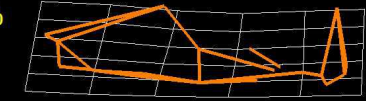
sloths? (NB above family level!)

  
 N_{sp} max/min R2
 5 1.5 87%



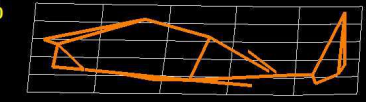
armadillos

  
 N_{sp} max/min R2
 16 5.3 24%






Dasyopus (subfam. with min. allom.)

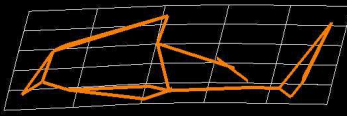
  
 N_{sp} max/min R2
 5 1.8 17%






x8

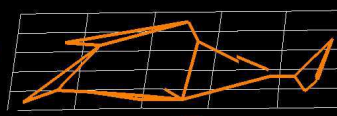
canids

  
 N_{sp} max/min R2
 31 2.7 22%






equids

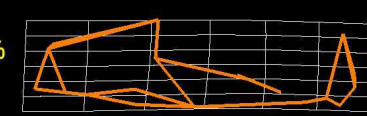
  
 N_{sp} max/min R2
 7 1.3 25%






x8

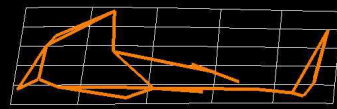
Erinaceinae

  
 N_{sp} max/min R2
 11 1.4 23%



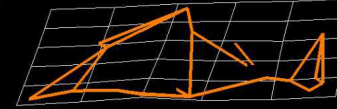
herpestids

  
 N_{sp} max/min R2
 13 2.3 16%






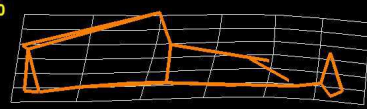
Antilopinae-Cephalophinae: *Neotopos*

  
 N_{sp} max/min R2
 22 2.6 17%



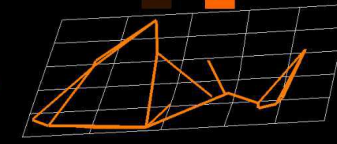
African Pteropodidae

  
 N_{sp} max/min R2
 15 2.9 32%



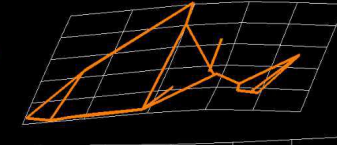
cercopithecini

  
 N_{sp} max/min R2
 21 1.9 64%




papionini

  
 N_{sp} max/min R2
 33 2.0 60%






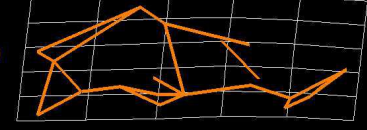
African colobines

  
 N_{sp} max/min R2
 19 1.4 30%



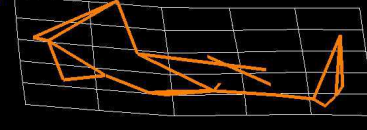
leporids

  
 N_{sp} max/min R2
 32 1.5 24%



Protoxerini squirrels

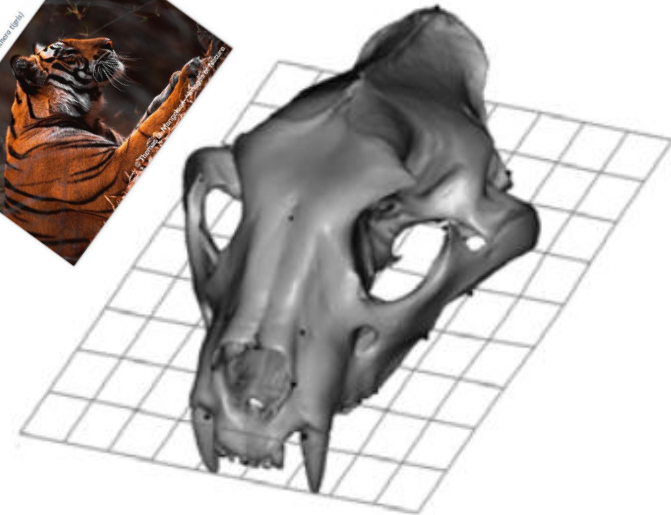
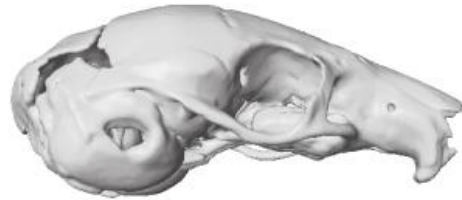
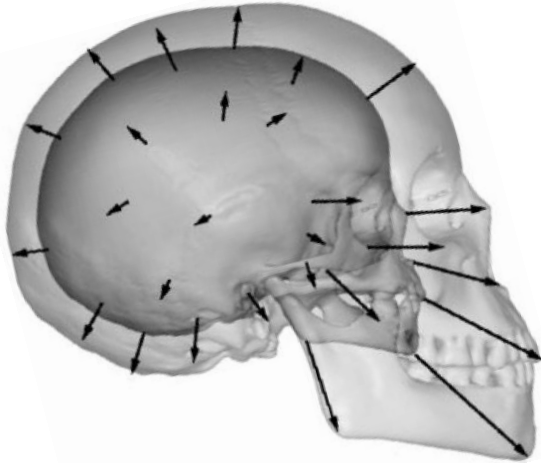
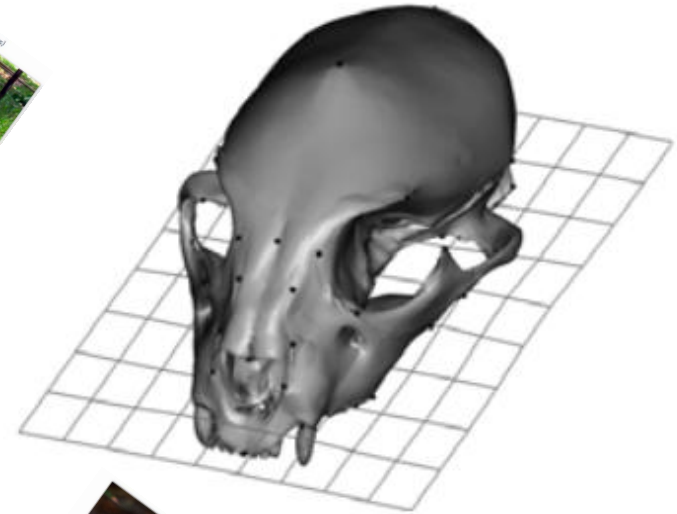
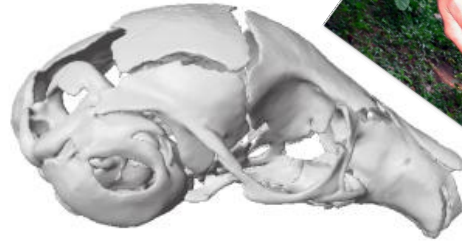
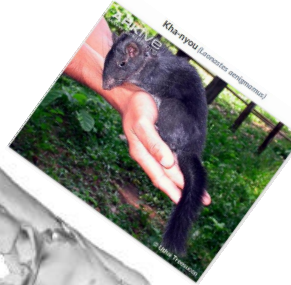
  
 N_{sp} max/min R2
 12 2.3 39%



why??? don't ask me: I am a *patternologist*
'measuring the shape of hills behind the fog'
(modified from Bookstein)



my 'favourite' guess: evodevo-constraints



Ontogeny of the cranial system in *Laonastes aenigmamus*

Anthony Herrel,¹ Anne-Claire Fabre,²⁻⁴ Jean-Pierre Hugot,⁵ Kham Keovitch,⁶ Dominique Adriaens,⁷ Loes Brabant,⁸ Luc Van Hoorebeke⁸ and Raphael Cornette^{5,9}

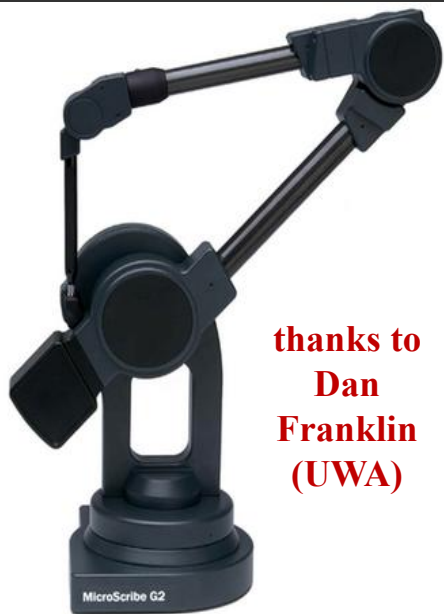
Three-dimensional cranial ontogeny in pantherines (*Panthera leo*, *P. onca*, *P. pardus*, *P. tigris*; Carnivora; Felidae)

VALENTINA SEGURA^{1,2*}, GUILLERMO H. CASSINI^{1,3,4} and FRANCISCO J. PREVOSTI¹⁻⁵

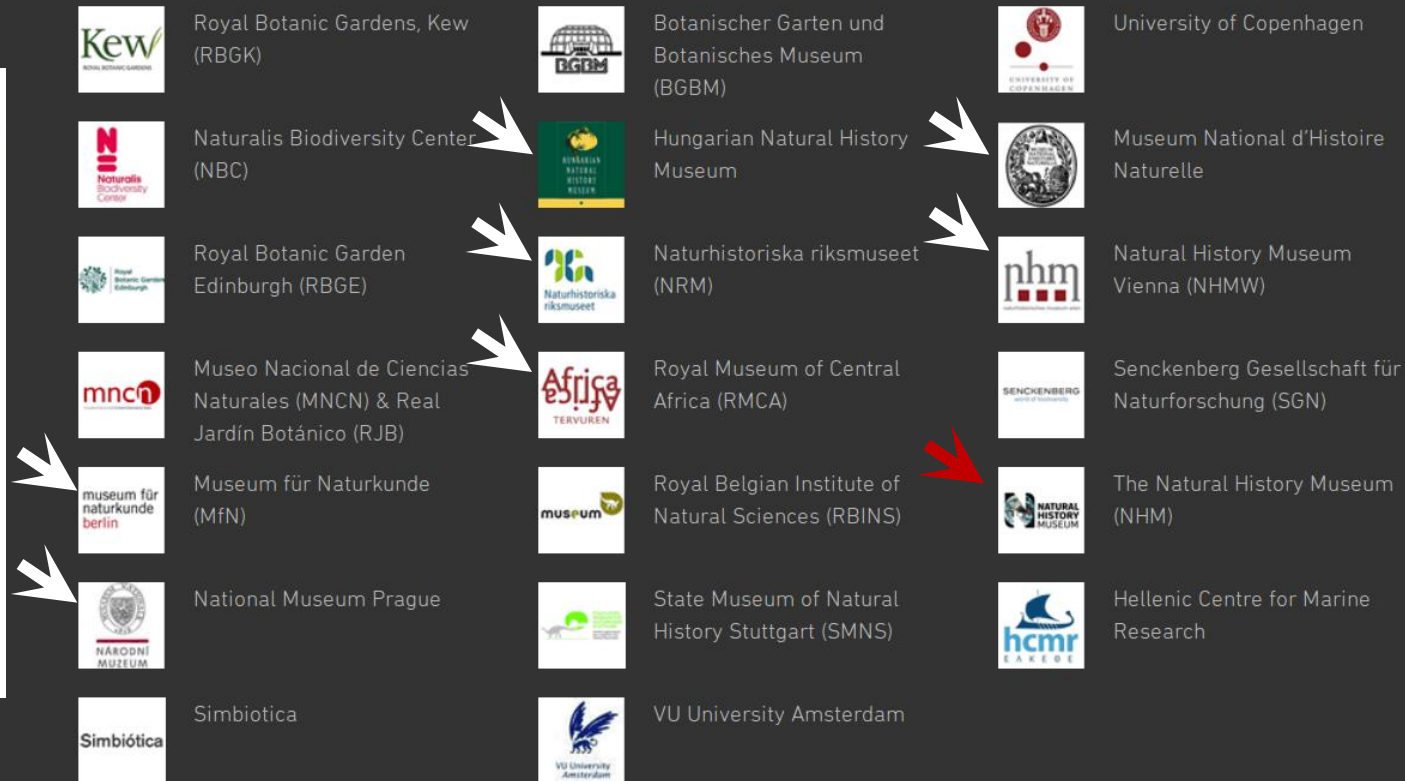


conclusions?

with low cost morphometrics
and great museum collections,
mammals may have another 'evo-rule'



thanks to
Dan
Franklin
(UWA)



**but without support from SYNTHESYS,
this would not have happened**

SYNTHESYS
Synthesis of systematic resources

[Contact us](#) | [FAQs](#) | [Partners](#) | [Previous outputs](#)



[Access](#)

[Joint Research Activities](#)

[Network Activities](#)

SYNTHESYS is an EC-funded project creating an integrated European infrastructure for natural history collections.

[> Find out more](#)



if I may, 2 suggestions for SYNTHEsys 4?

feedback from applicants (even if not funded)

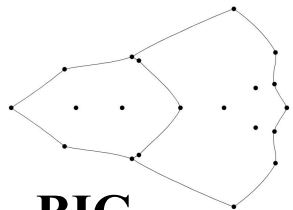
'capitalize' on visits by experts

(maybe in coop. with local universities):

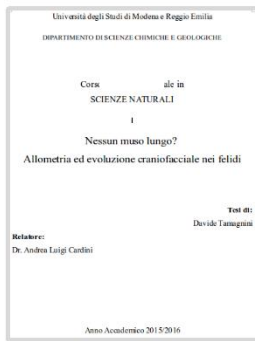
practical demos? lectures? workshops?



many thanks for inviting me! & 1000 thanks to:
SYNTHEsys, all organizers & administrators,
 museum curators and staff, and all collaborators
 including recent ones on big cats and kangaroos



BIG



SMALL

