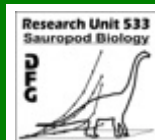




Phylogenetic statistics and biological laws

Marcus Clauss

Clinic for Zoo Animals, Exotic Pets and Wildlife, Vetsuisse Faculty, University of Zurich, Switzerland
Evolutionary Biology, Zurich 2019



University of
Zurich^{UZH}



Clinic
of Zoo Animals, Exotic Pets and Wildlife



Allometry reminder

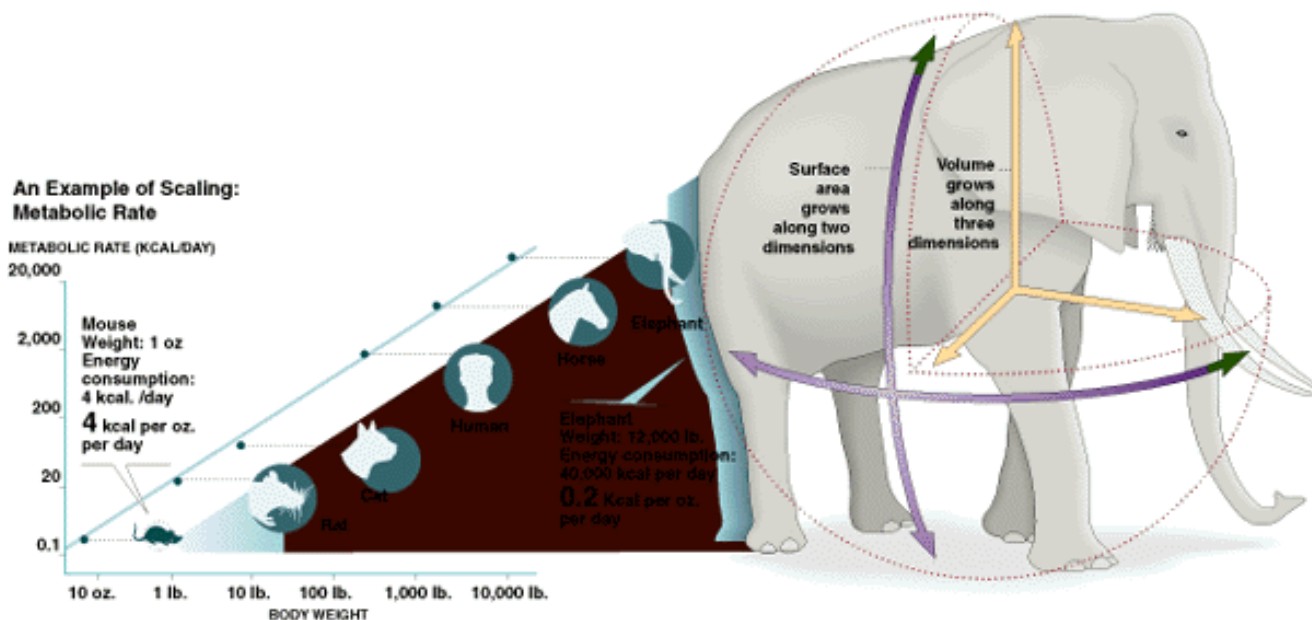


Scaling: fundamental (conceptual) relevance of body mass

Most biologists consider body mass the most important characteristic of an organism. It is also (mostly) easy to measure.

All morphological and physiological traits scale somehow with body mass.

"Scaling is interesting because, aside from natural selection, it is one of the few laws we really have in biology." John Gittleman





Allometries

Morphological, physiological and life history variables scale with body mass.



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Linear scaling: $y = a \text{ BM}^{1.0}$ or $\log y = \log a + 1.0 \text{ BM}$

Allometric scaling: $y = a \text{ BM}^b$ or $\log y = \log a + b \text{ BM}$



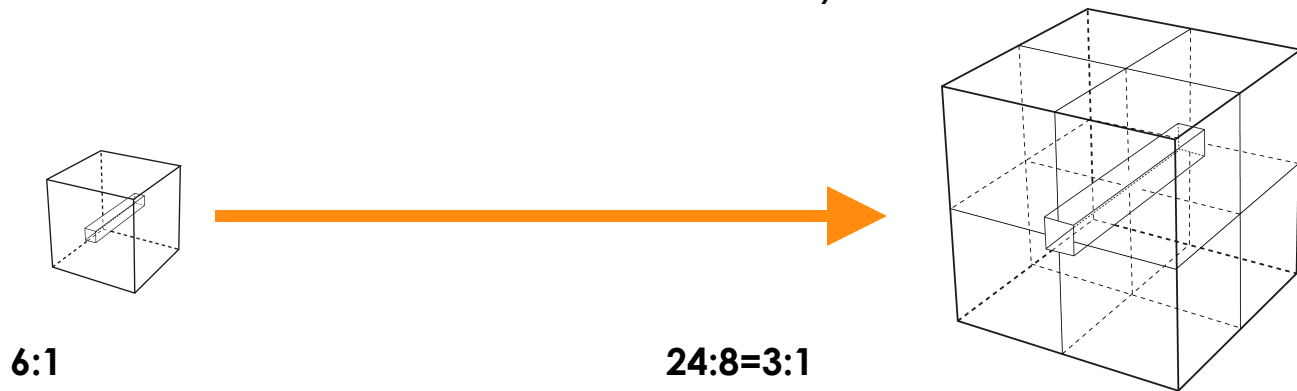
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(allometric scaling mostly explained by geometry – e.g. surface-volume shifts, distribution networks etc.)





Allometries

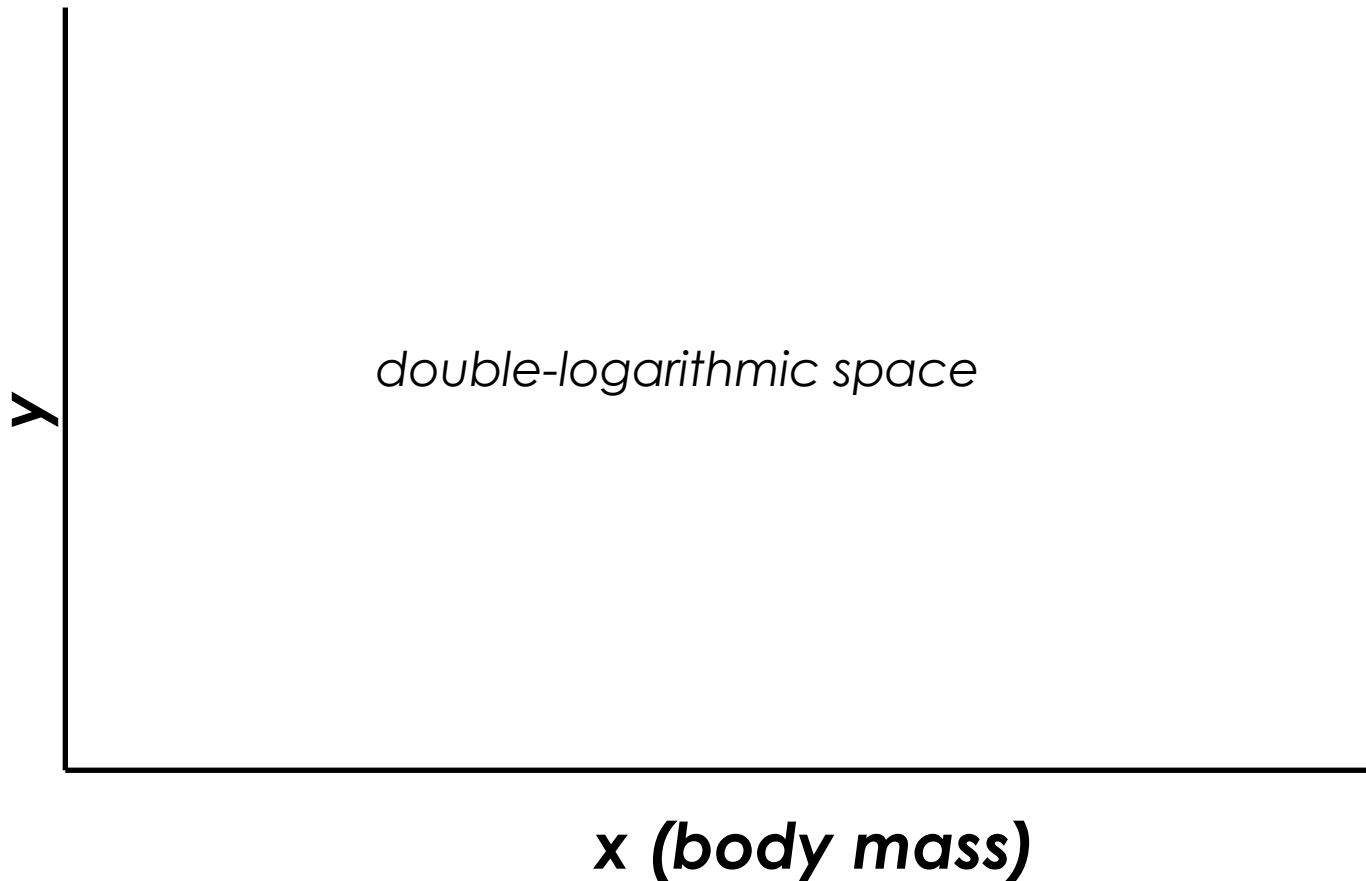
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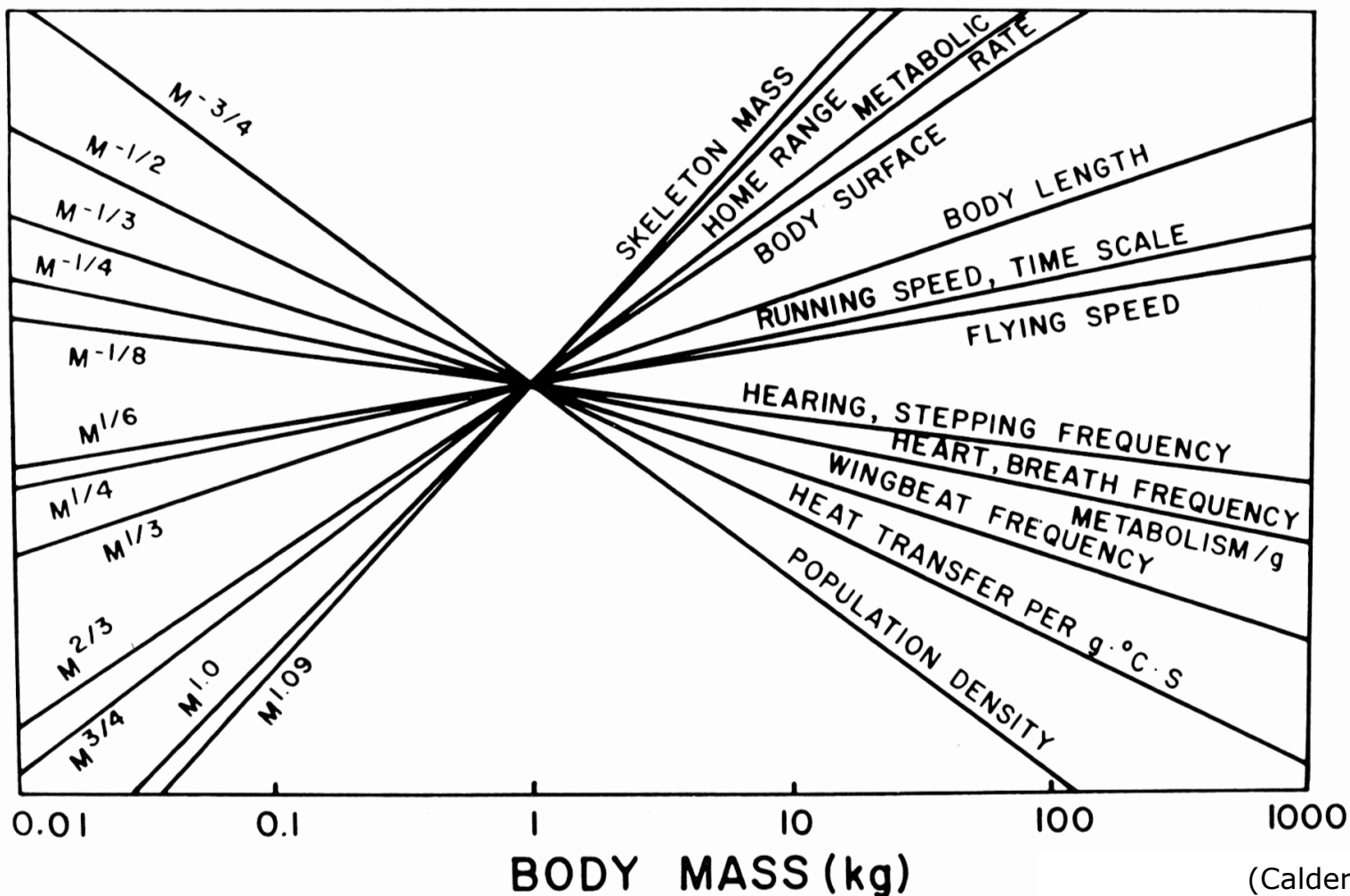
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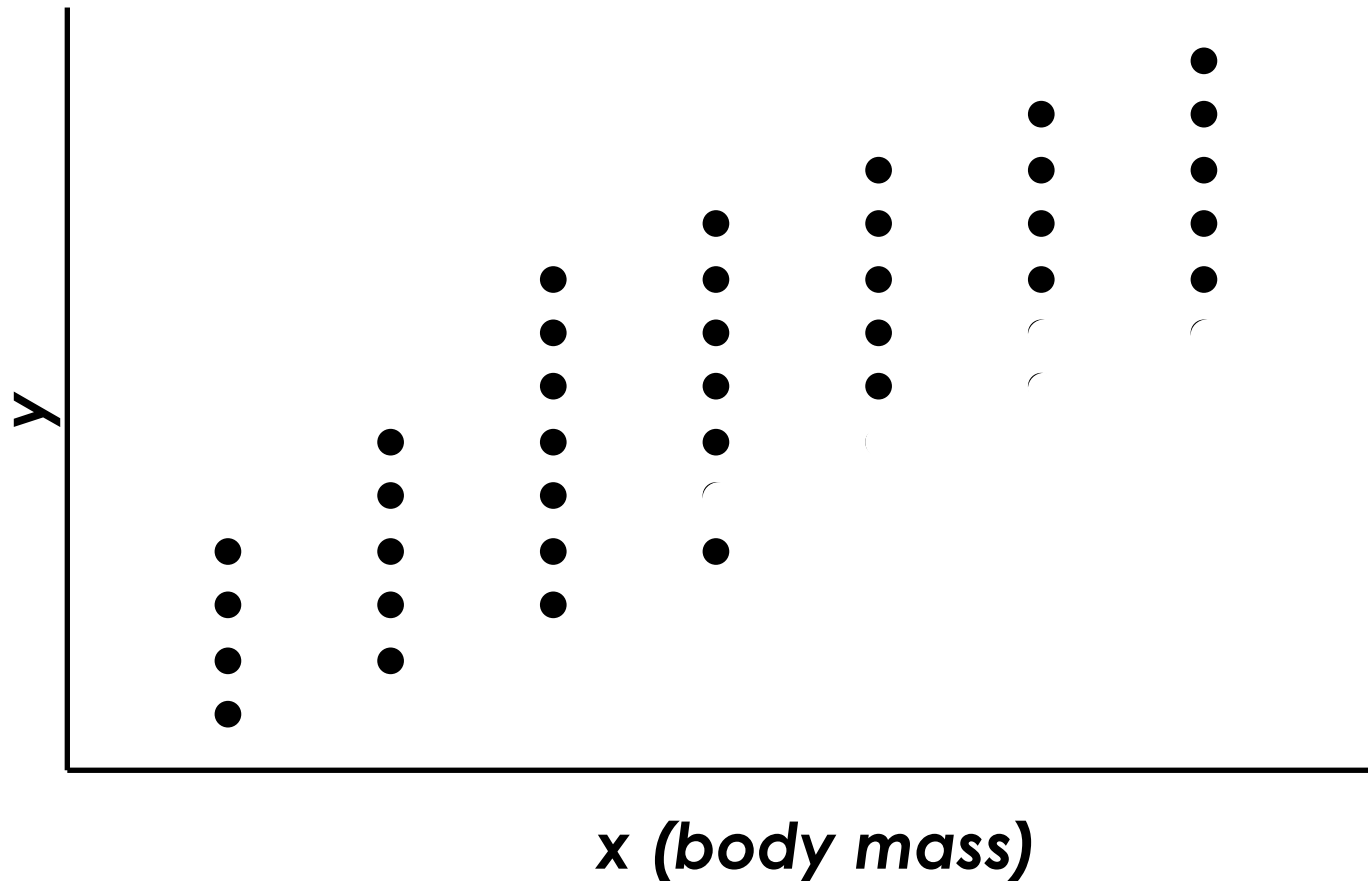


(Calder 1983)



Interpreting allometries

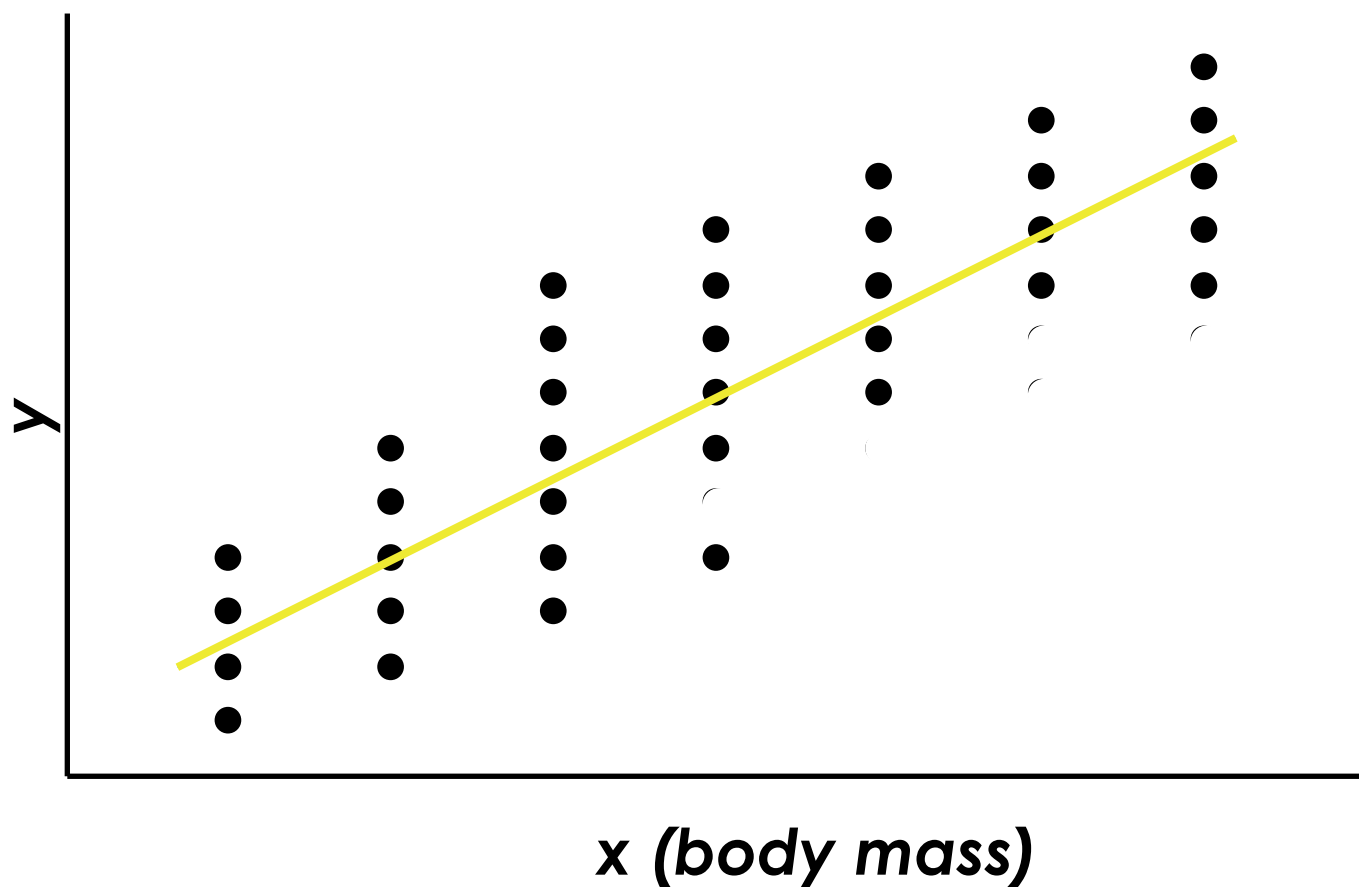
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Interpreting allometries

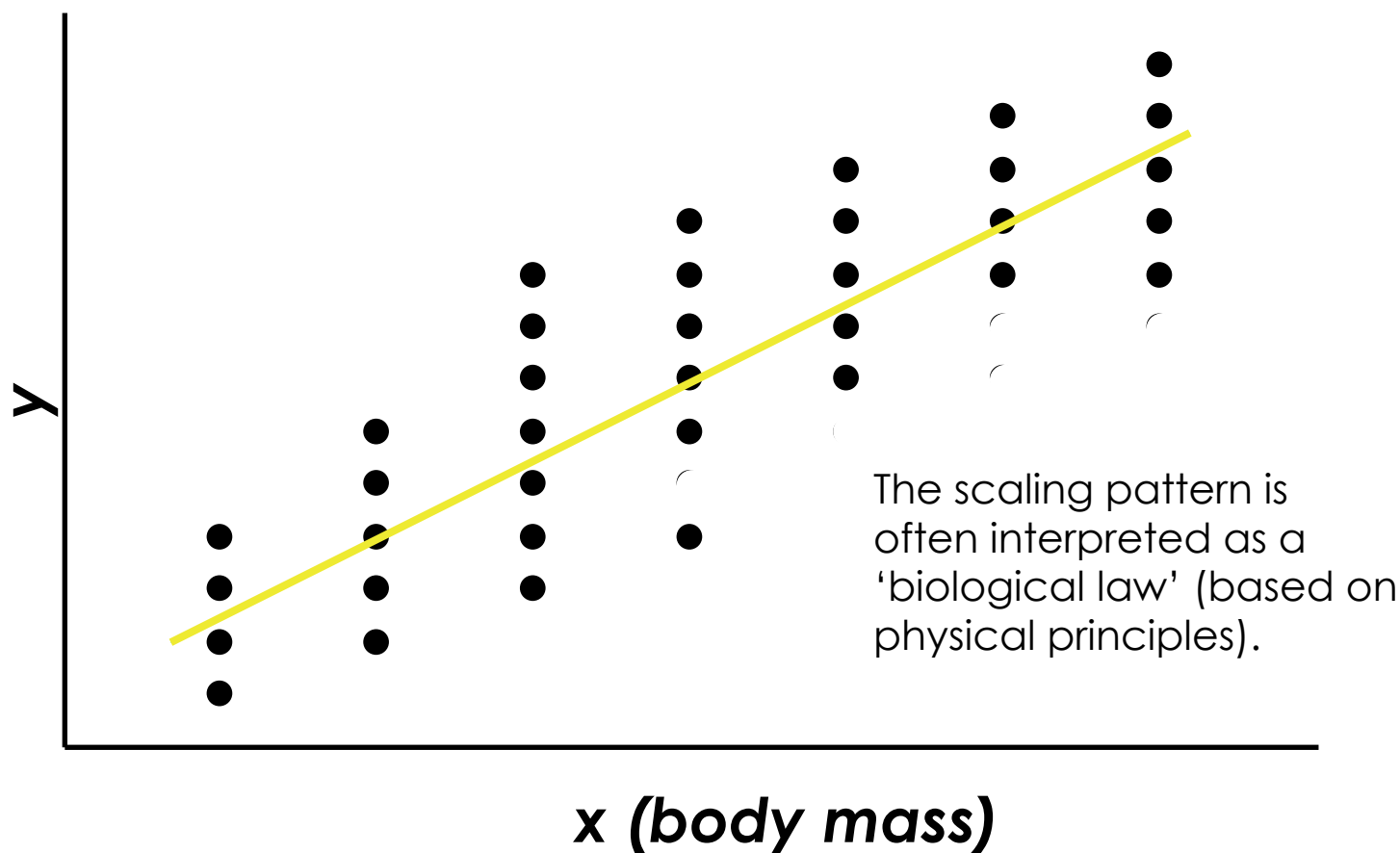
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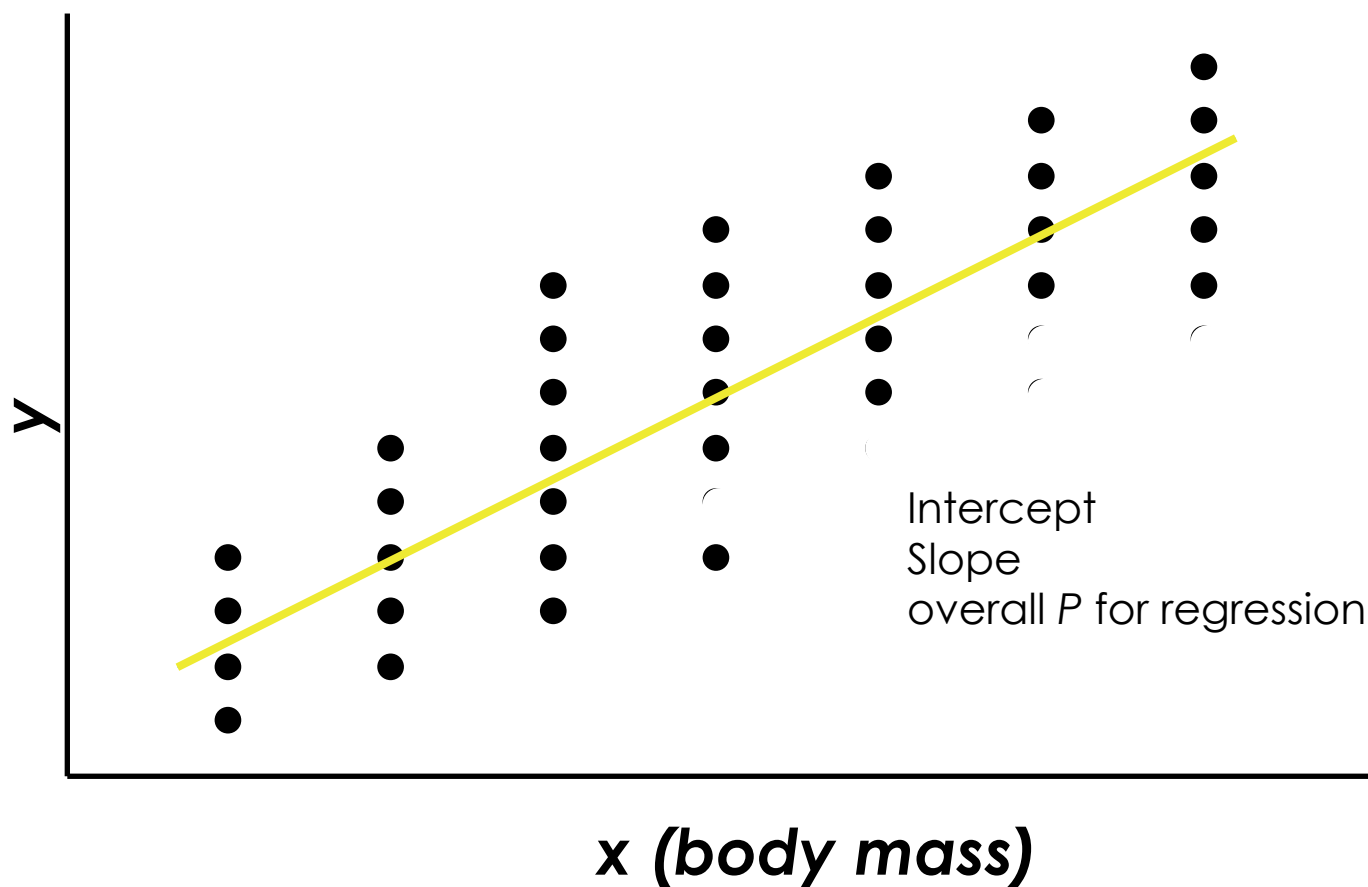
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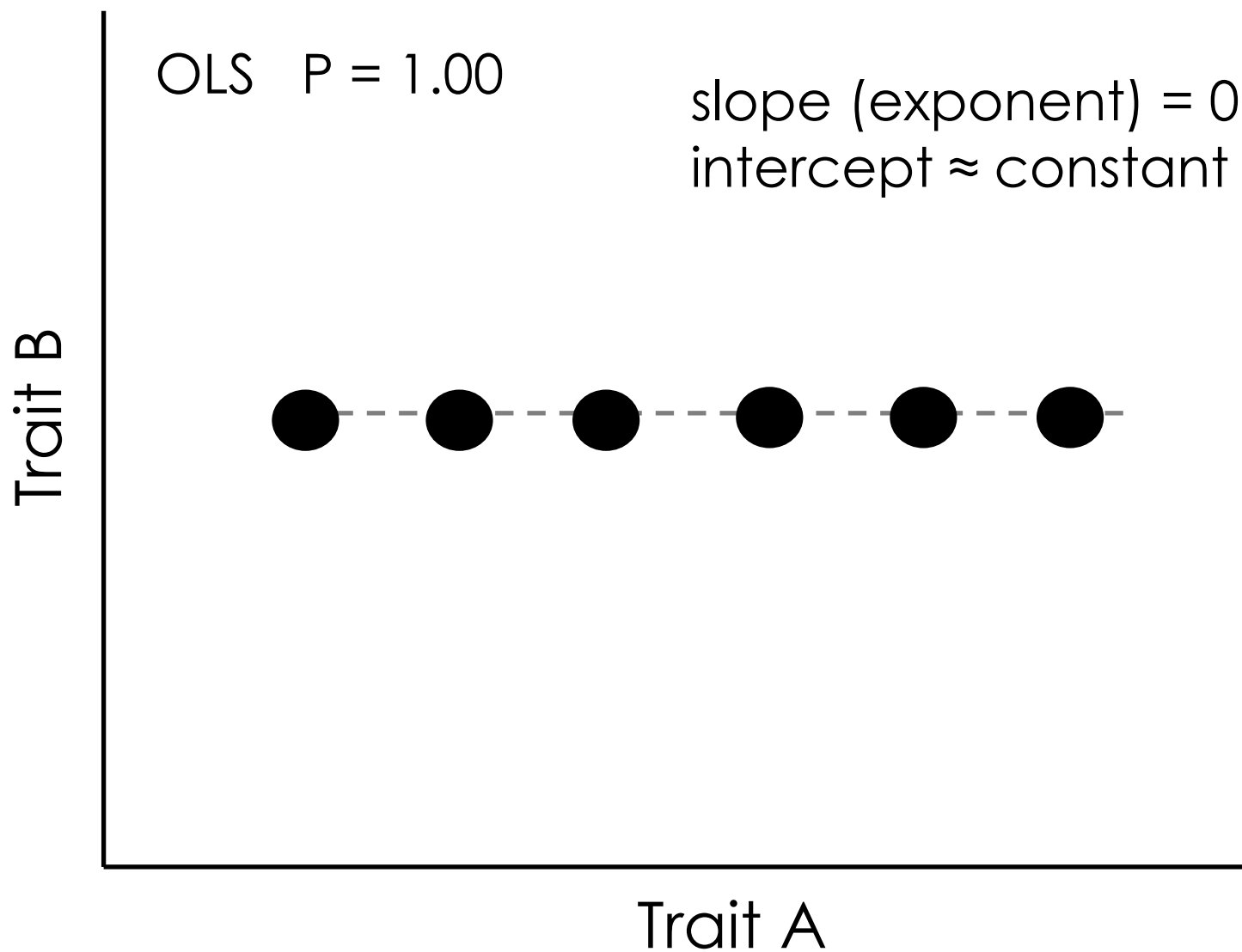
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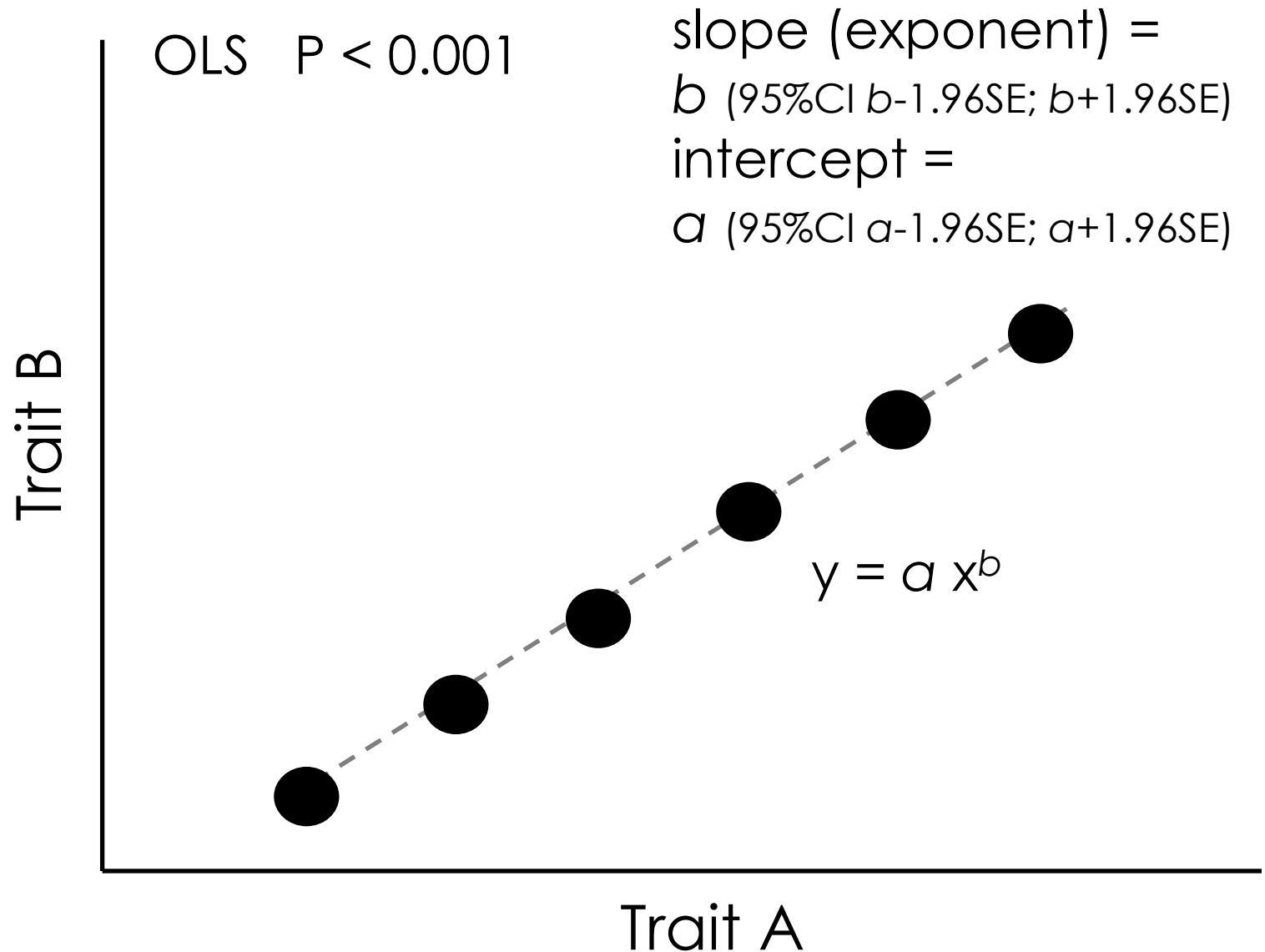


Testing for allometries





Testing for allometries





Using allometries

Using allometric relationships to extrapolate data for other species.



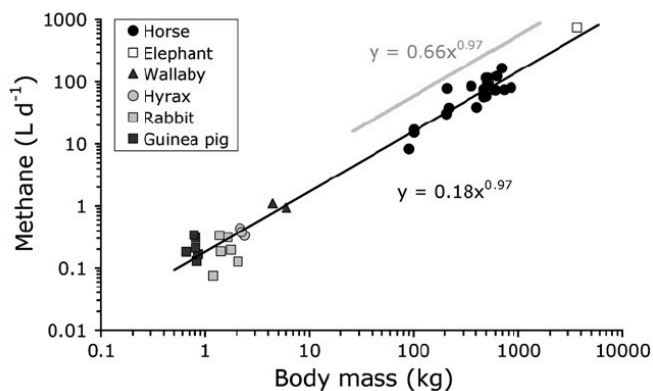
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Methane output of rabbits (*Oryctolagus cuniculus*) and guinea pigs (*Cavia porcellus*) fed a hay-only diet: Implications for the scaling of methane production with body mass in non-ruminant mammalian herbivores

Ragna Franz ^a, Carla R. Soliva ^b, Michael Kreuzer ^b, Jürgen Hummel ^c, Marcus Clauss ^{a,*}

Comparative Biochemistry and Physiology, Part A 158 (2011) 177–181





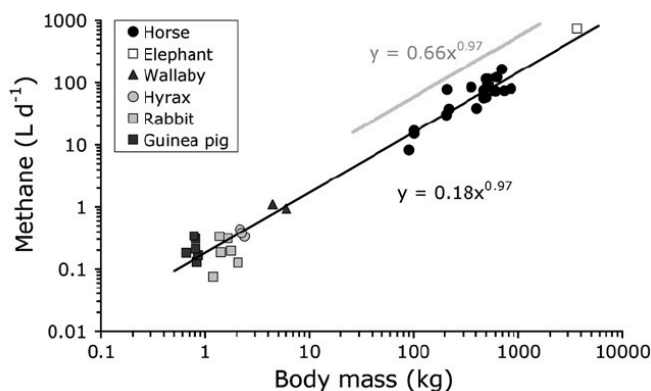
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Could methane produced by sauropod dinosaurs have helped drive Mesozoic climate warmth?

David M. Wilkinson^{1,*},
Euan G. Nisbet²,
and Graeme D. Ruxton³

Current Biology Vol 22 No 9
R292



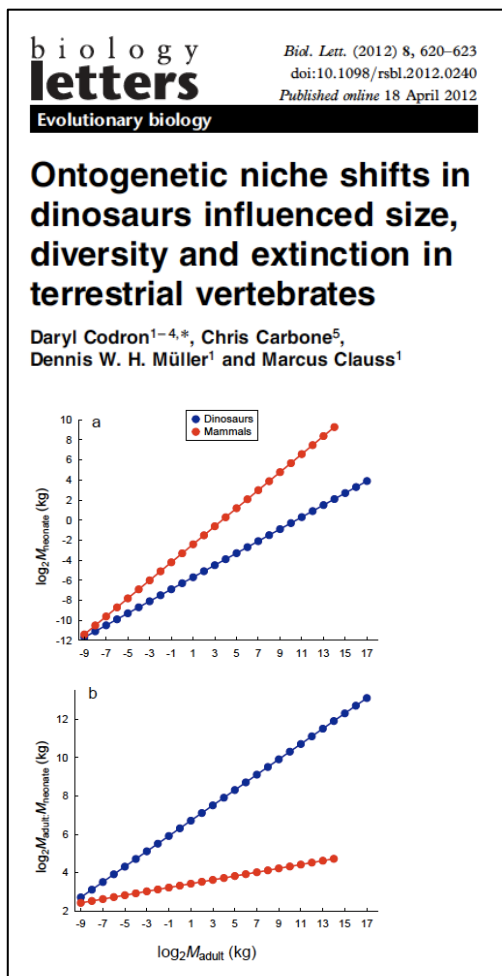
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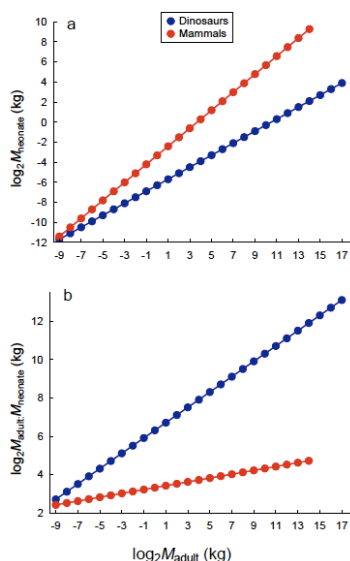
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biology
letters
Evolutionary biology

Biol. Lett. (2012) 8, 620–623
doi:10.1098/rsbl.2012.0240
Published online 18 April 2012

Ontogenetic niche shifts in dinosaurs influenced size, diversity and extinction in terrestrial vertebrates

Daryl Codron^{1–4,*}, Chris Carbone⁵,
Dennis W. H. Müller¹ and Marcus Clauss¹



Ecological modelling, size distributions and taphonomic size bias in dinosaur faunas: a comment on Codron *et al.* (2012)

Caleb Marshall Brown¹, Nicolás E. Campione¹, Henrique Corrêa Giacomini¹,
Lorna J. O'Brien¹, Matthew J. Vavrek² and David C. Evans^{1,2}

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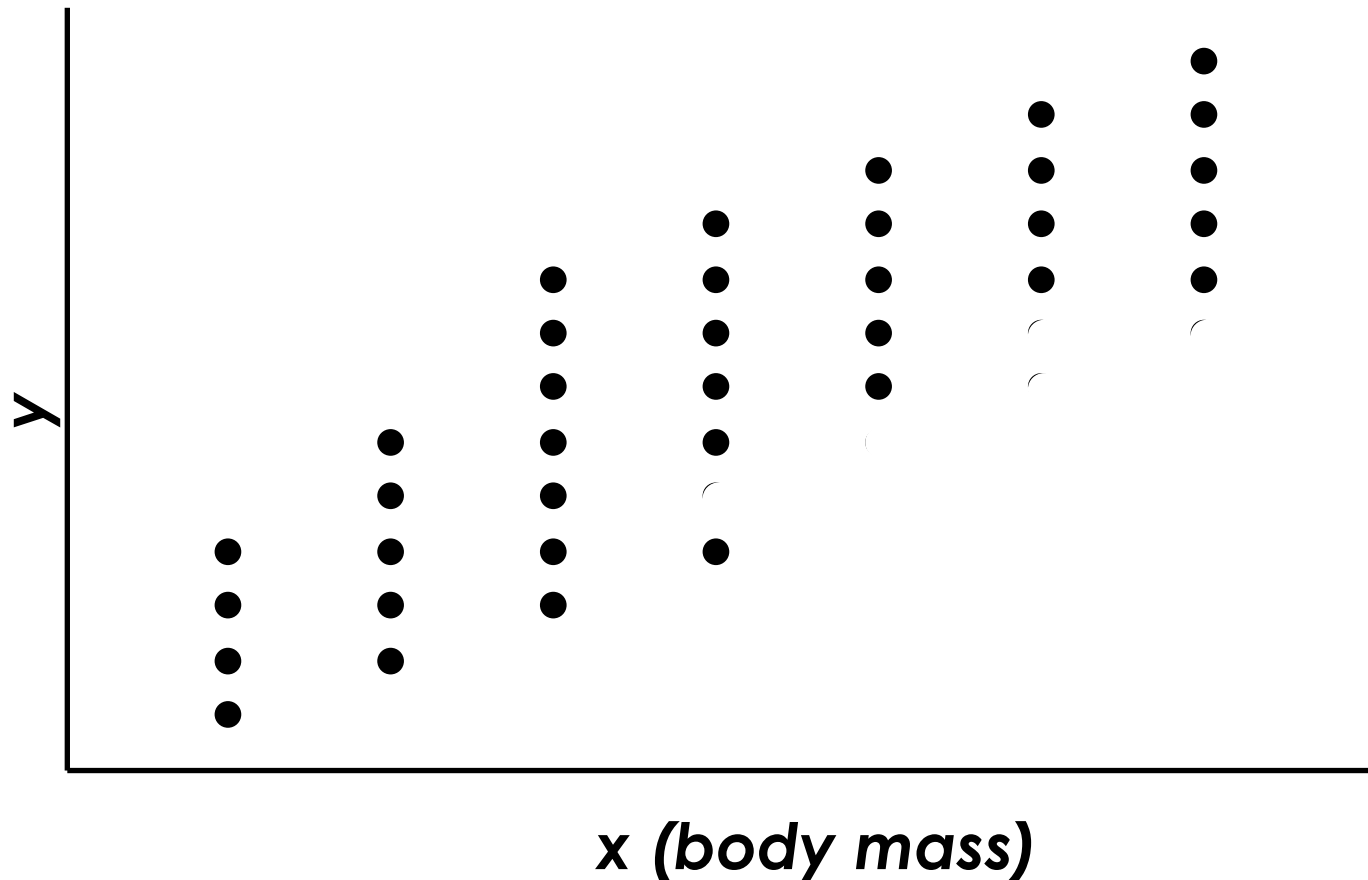


Phylogenetic statistics



Comparative statistics

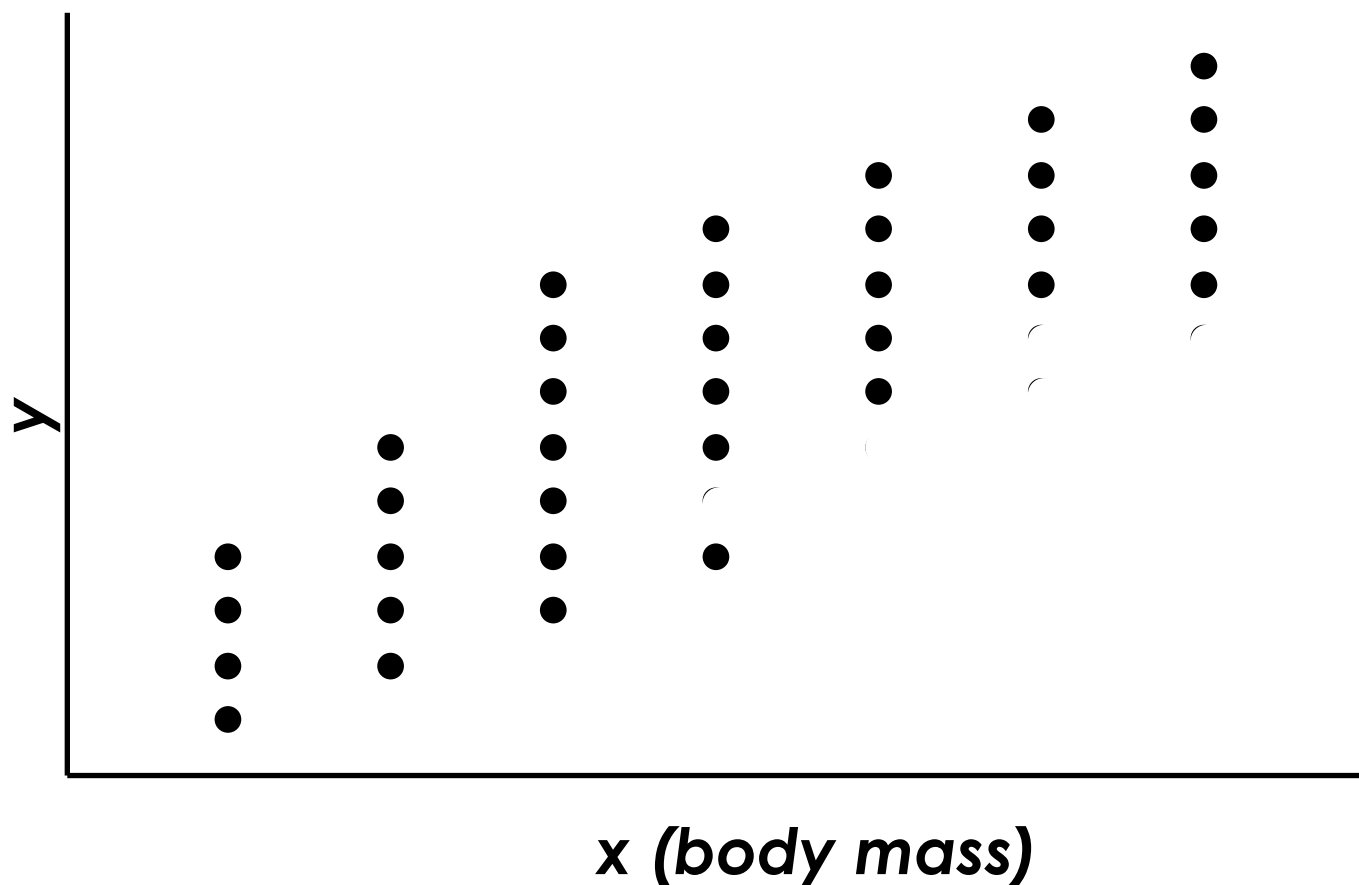
Conventional regression analysis assumes independence of data points.





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Comparative statistics

Conventional regression analysis assumes independence of data points.
But this is violated by phylogenetic relationships.

Therefore, we perform allometric analyses also with accounting for phylogeny, using PGLS (Phylogenetic Generalized Least Squares).

Results mostly did not differ from conventional statistics in a relevant way, but the intensive use of comparative statistics (also with additional examples) led to formulation of some concepts new to ourselves.



Comparative statistics

Three (of many) important test statistics:



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Pagel's lambda λ : if 95%CI includes 0, then there is no phylogenetic structure in the dataset.

Does not decide whether the relationship is significant or not, but whether phylogenetic statistics need to be used or not.

Assumes Brownian motion; other measures of phylogenetic structure assuming other evolutionary scenarios exist.



Comparative statistics - errors



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Type II error: you overlook a relationship where there is one (evident when you account for the phylogenetic structure of the data)

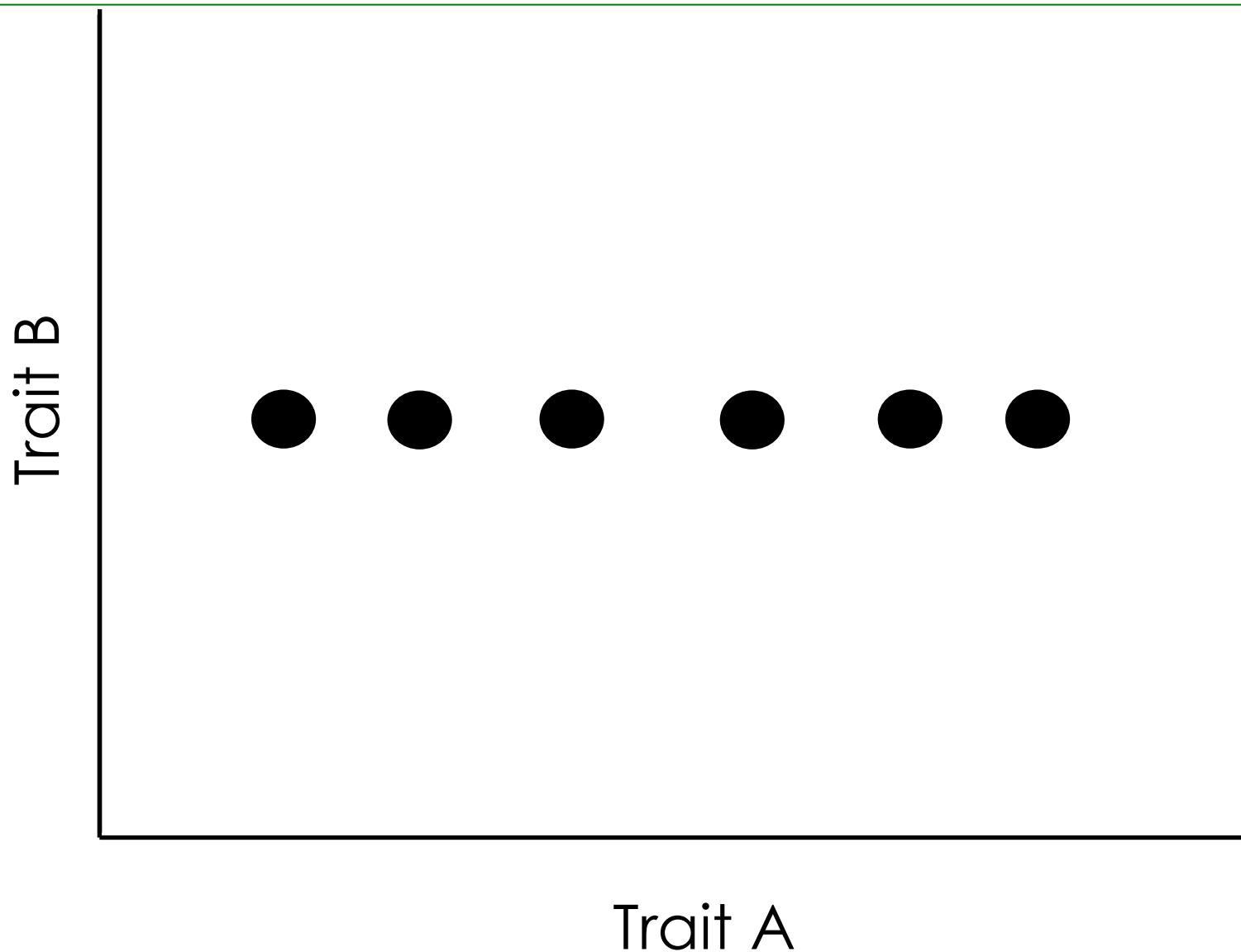
Just an error: you estimate a different parameter (e.g., allometric slope) depending on whether you account for phylogeny or not



Page1's lambda(λ) examples



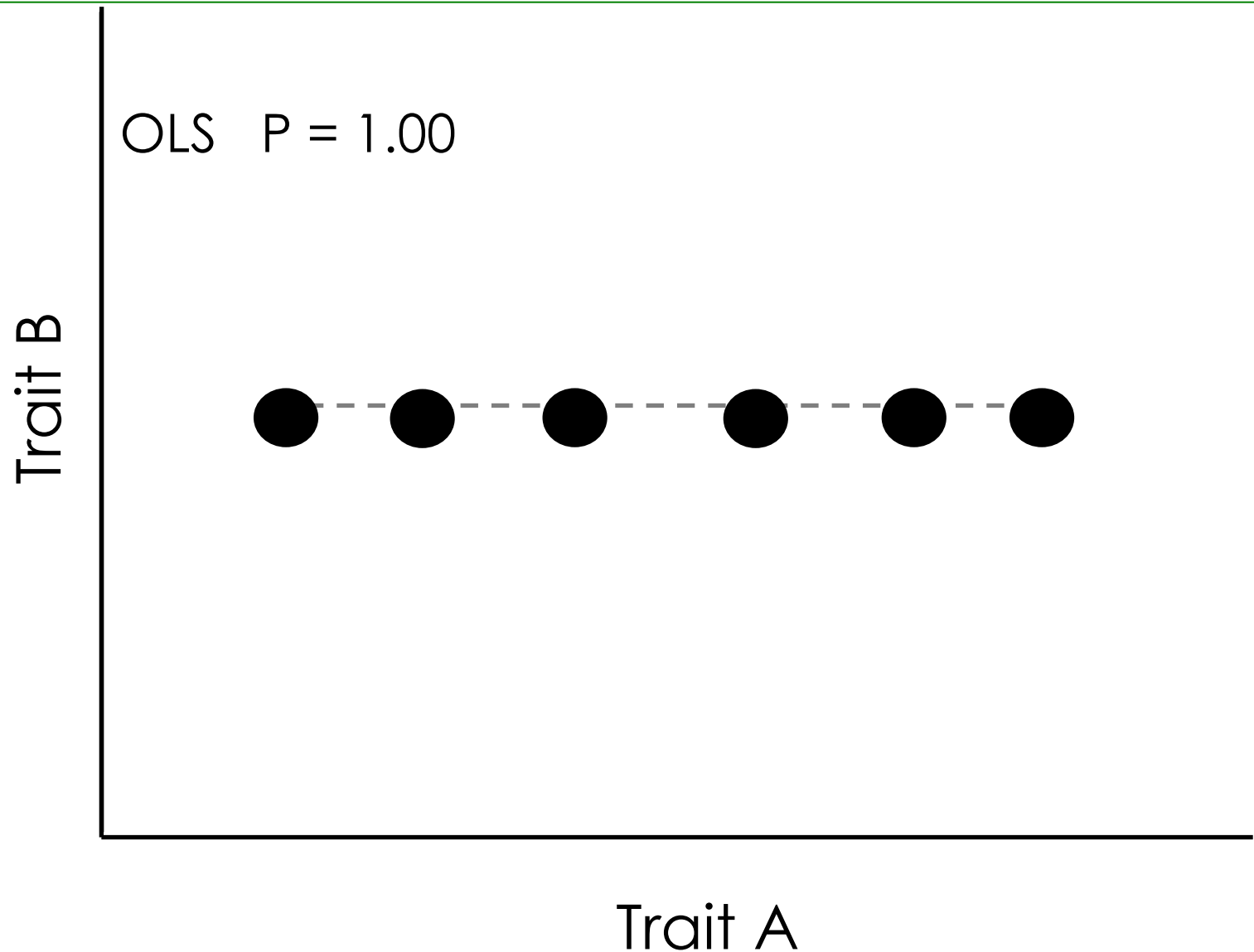
Accounting for phylogeny



from Dittmann et al. (2015)

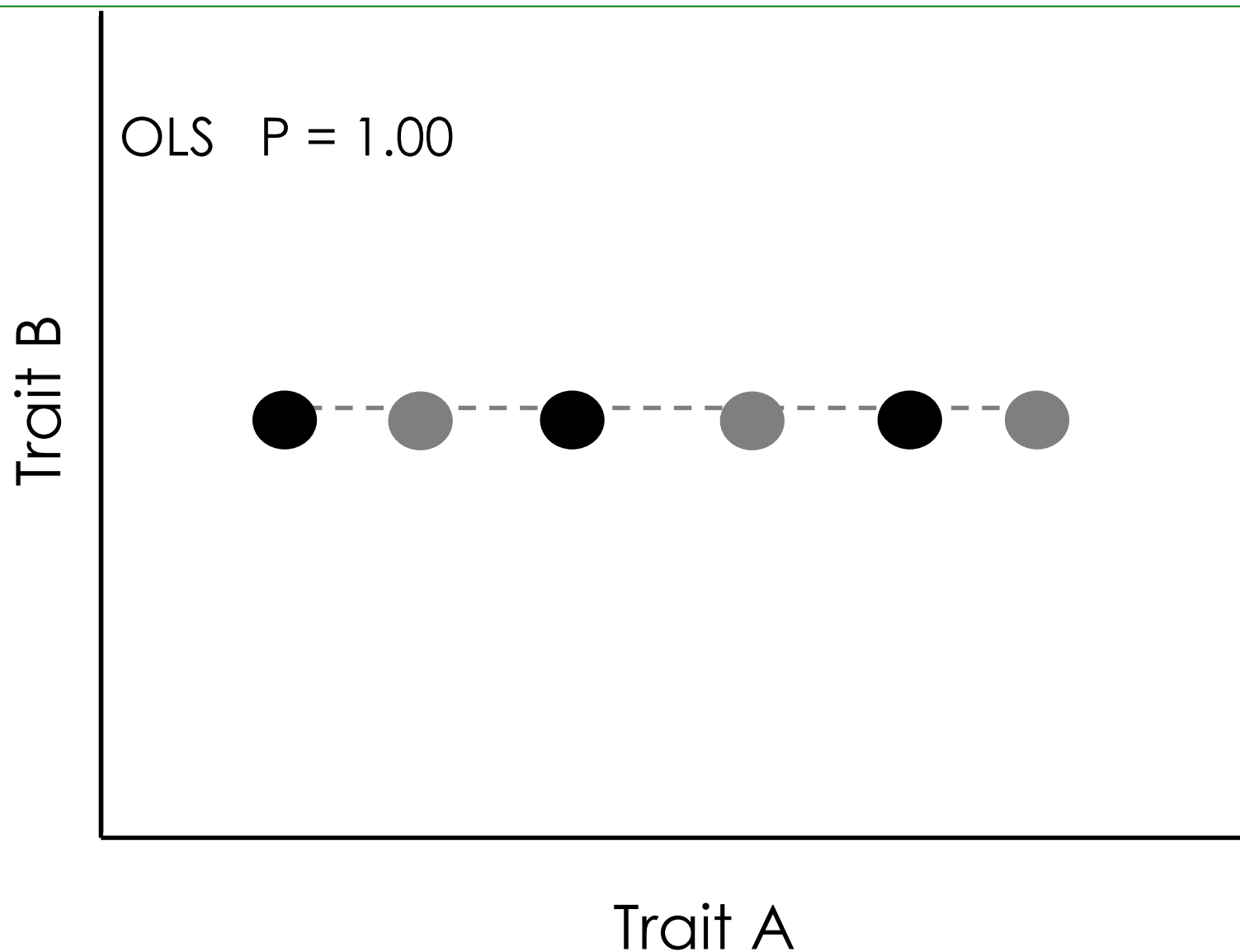


Accounting for phylogeny



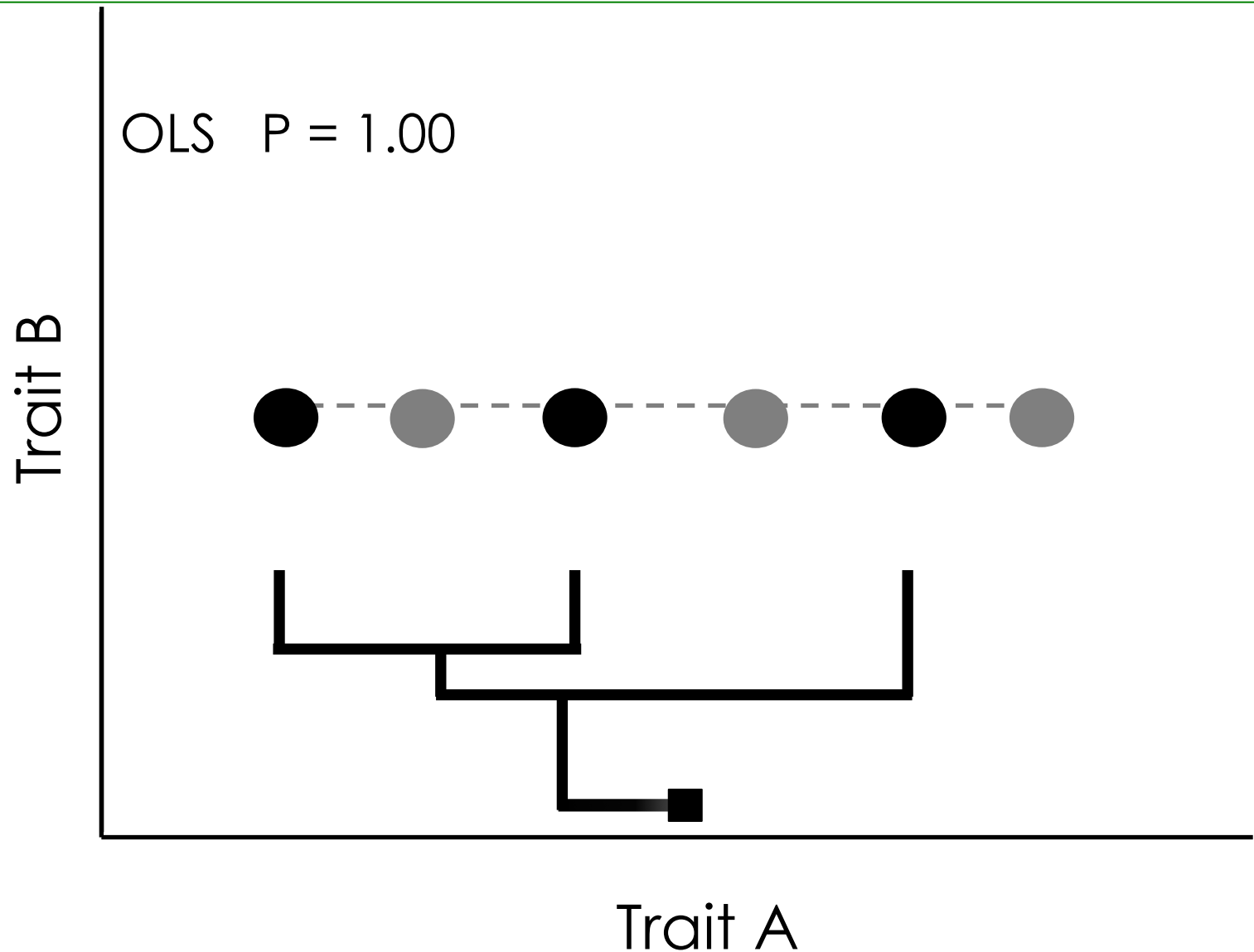


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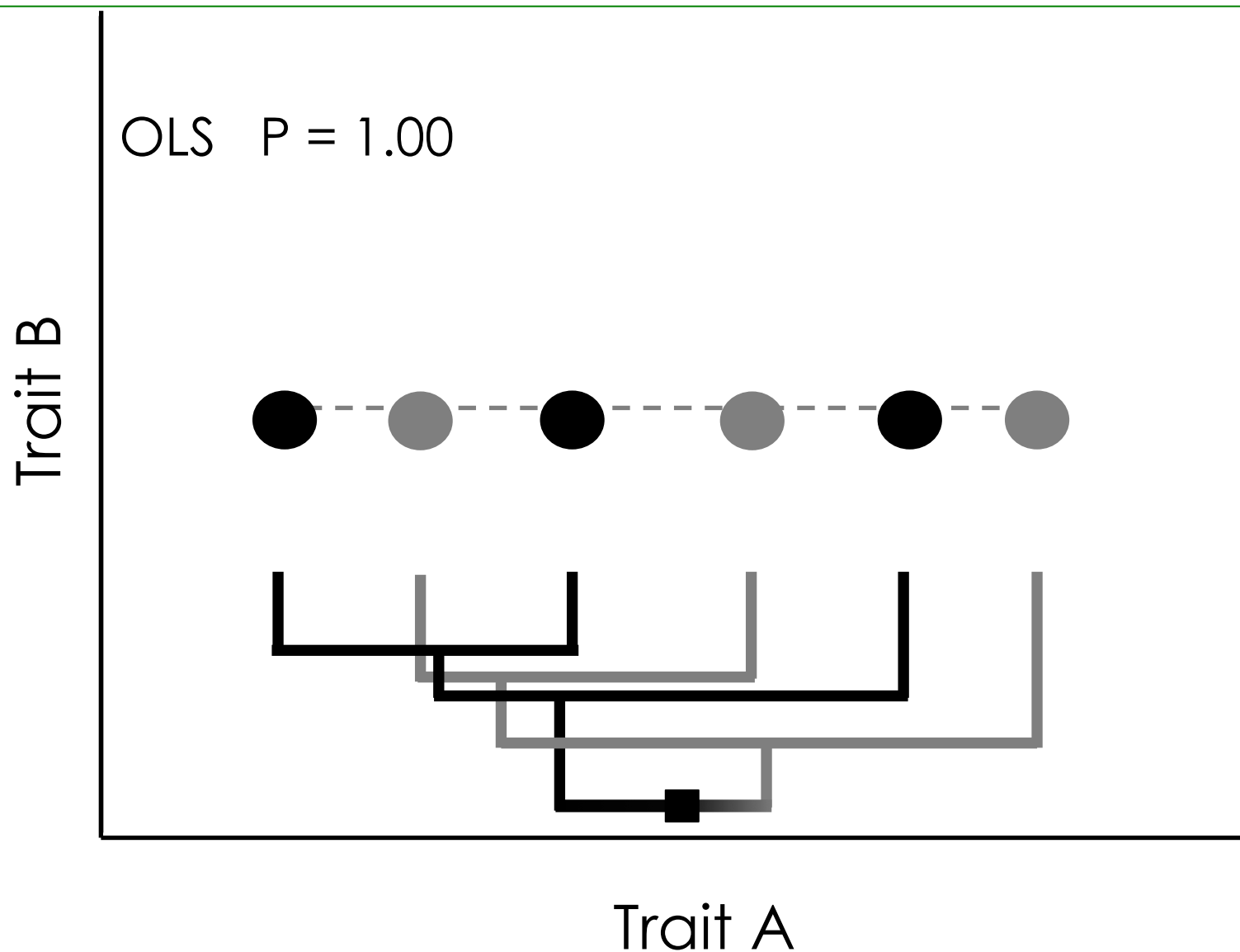


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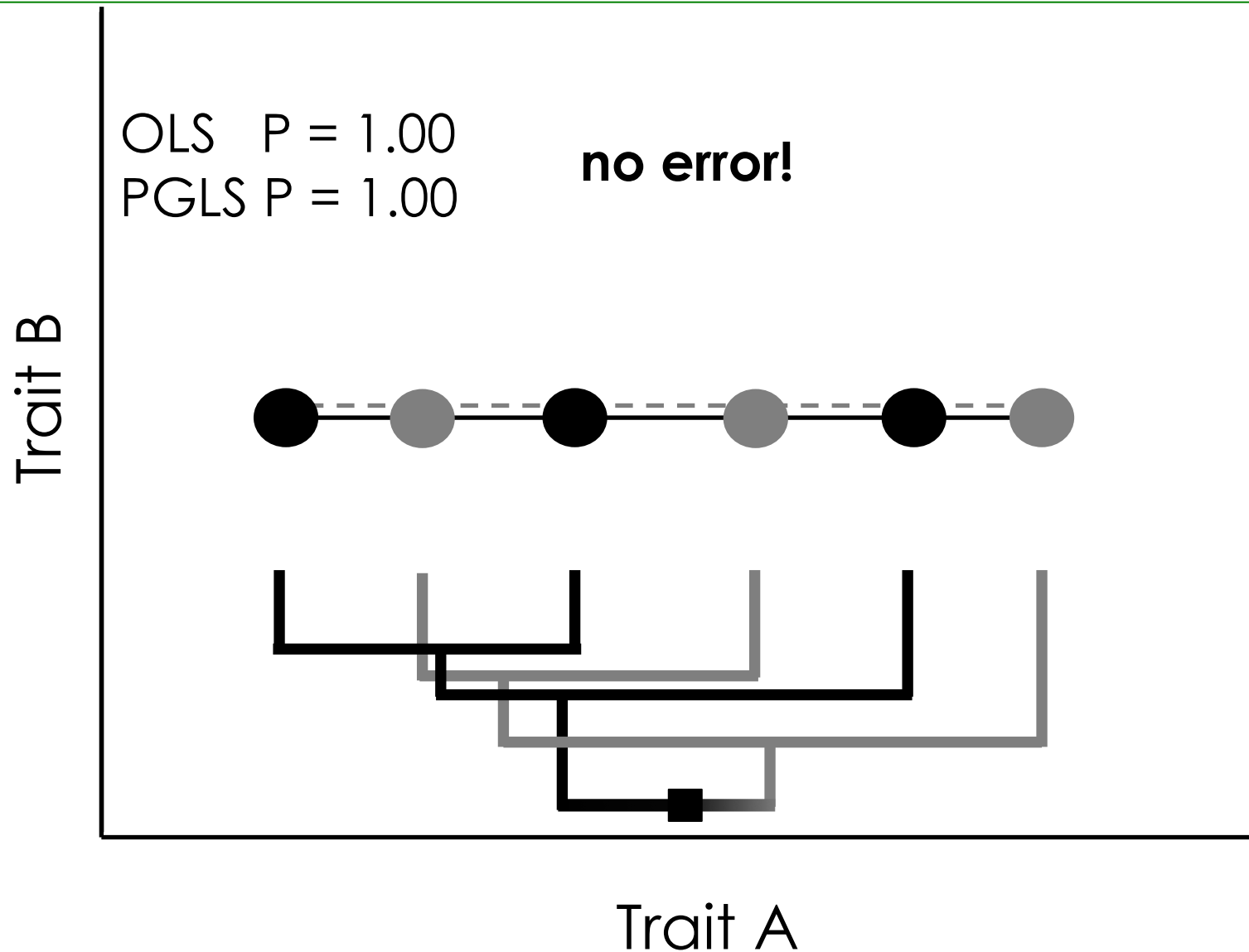


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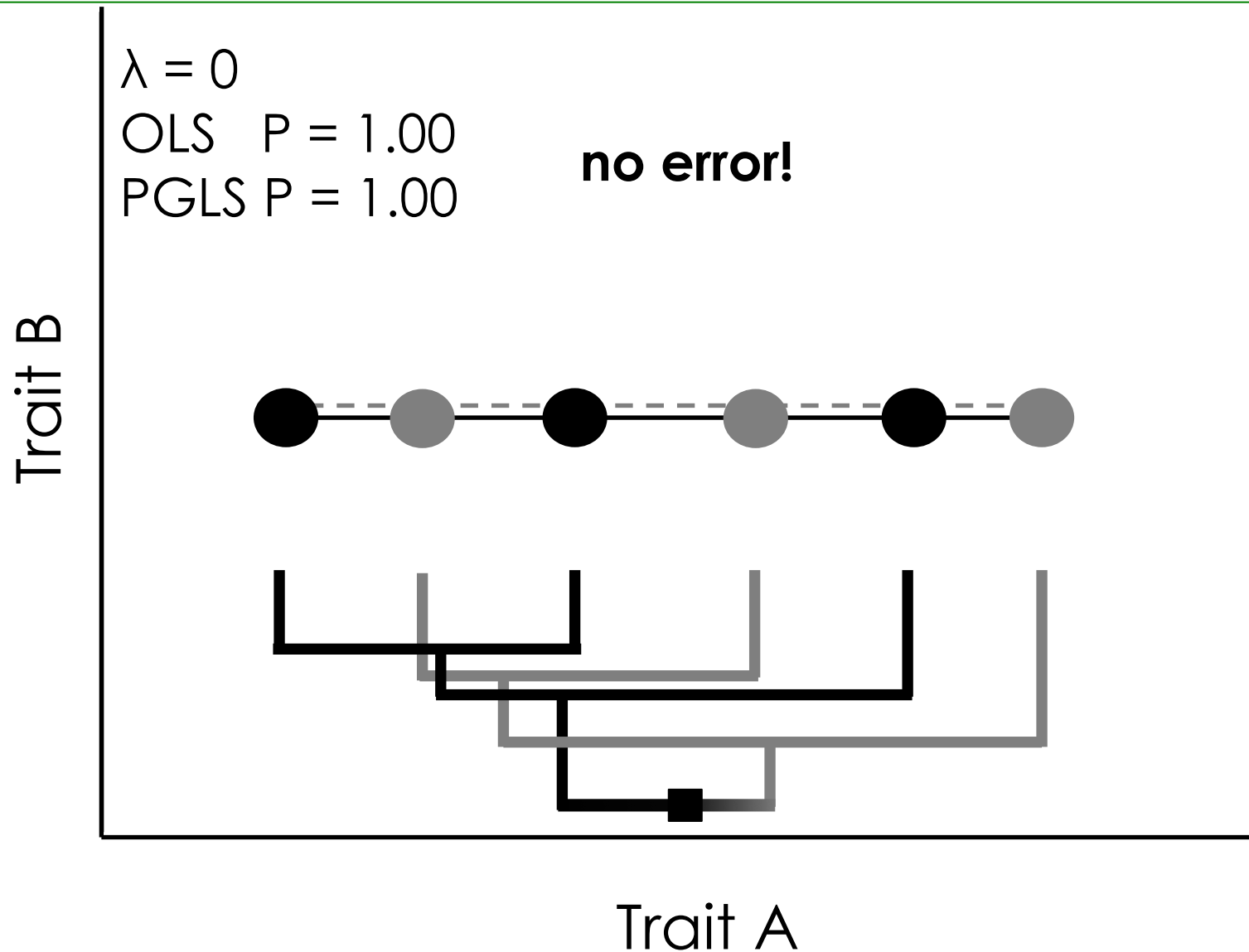


Accounting for phylogeny





Accounting for phylogeny



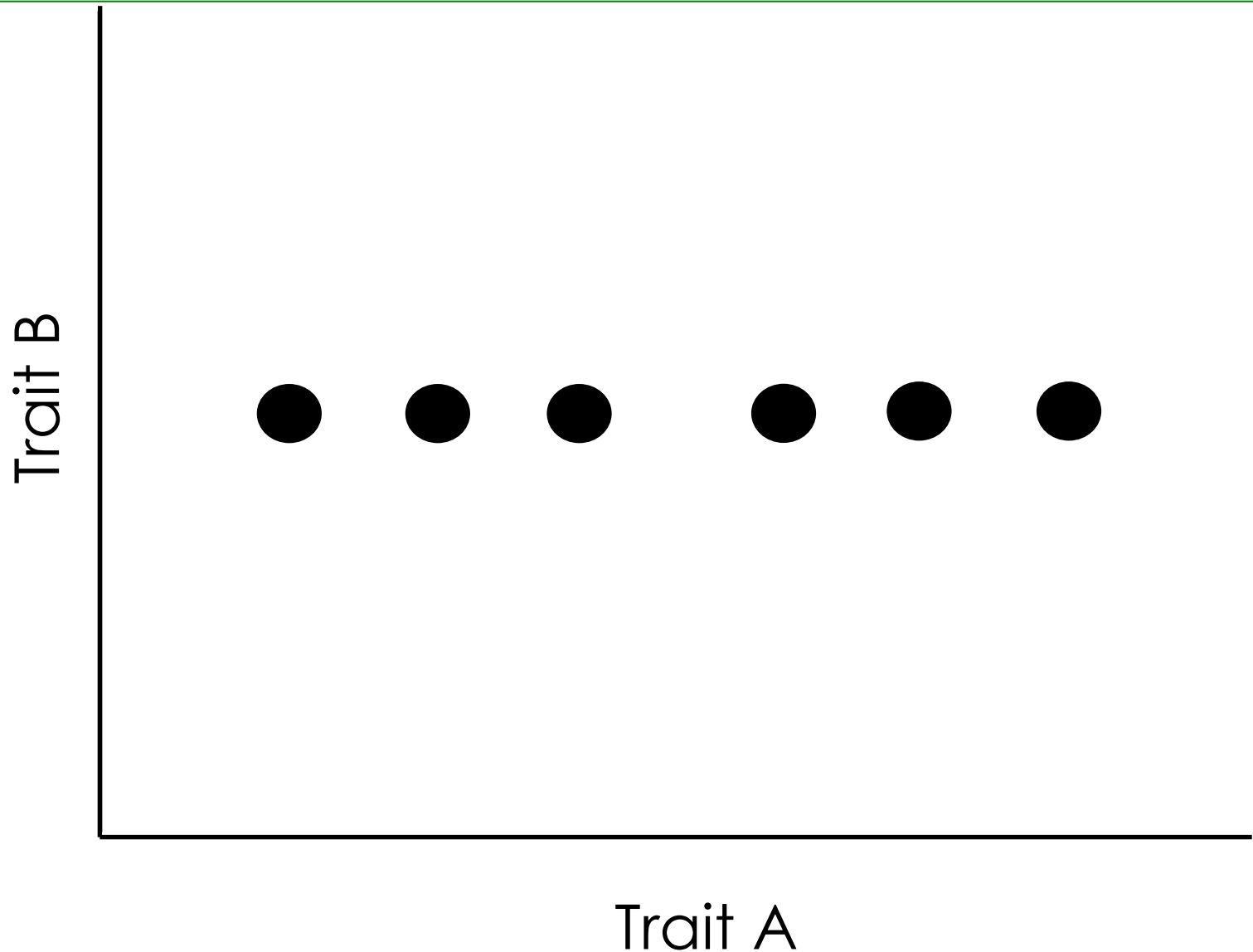


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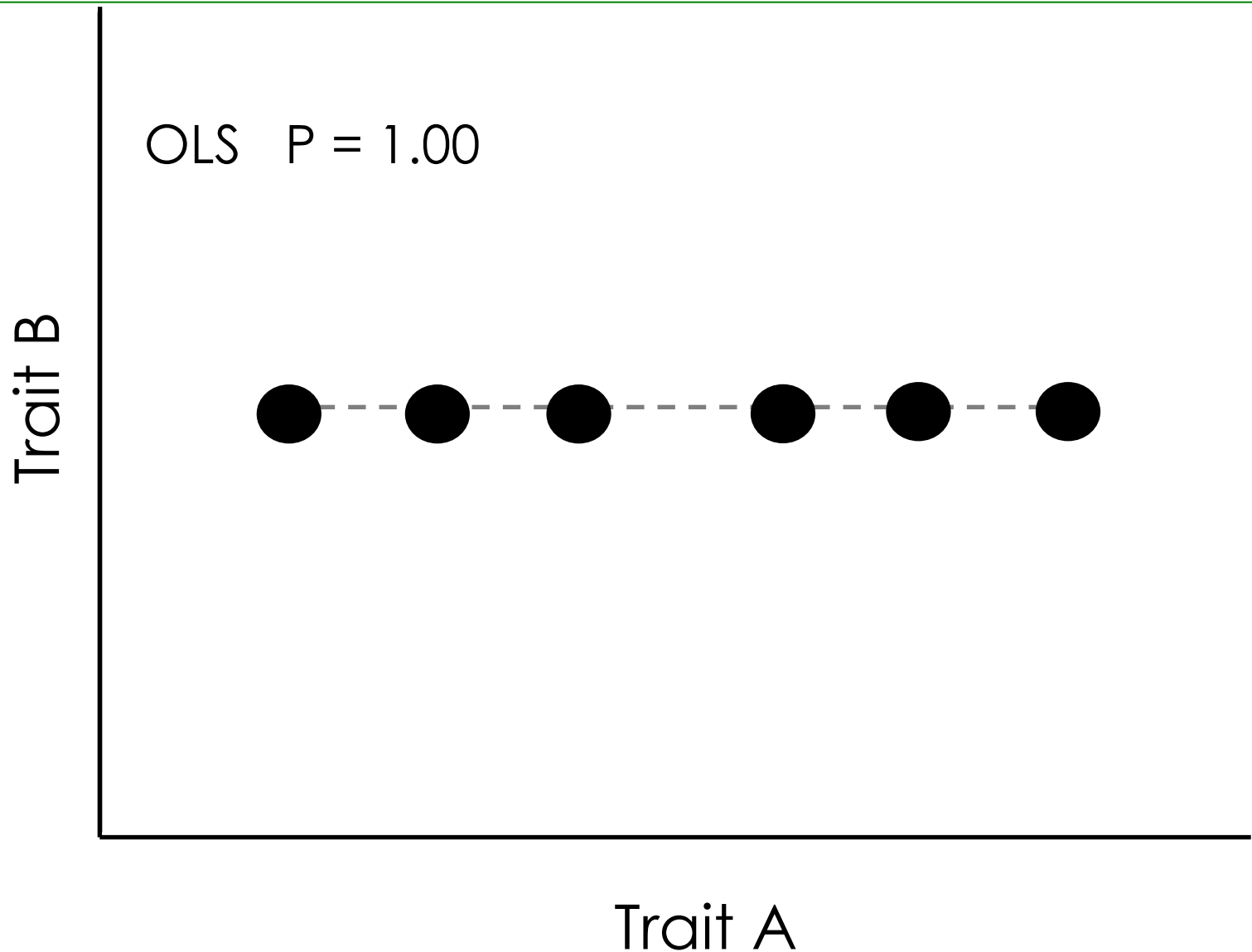
Accounting for phylogeny



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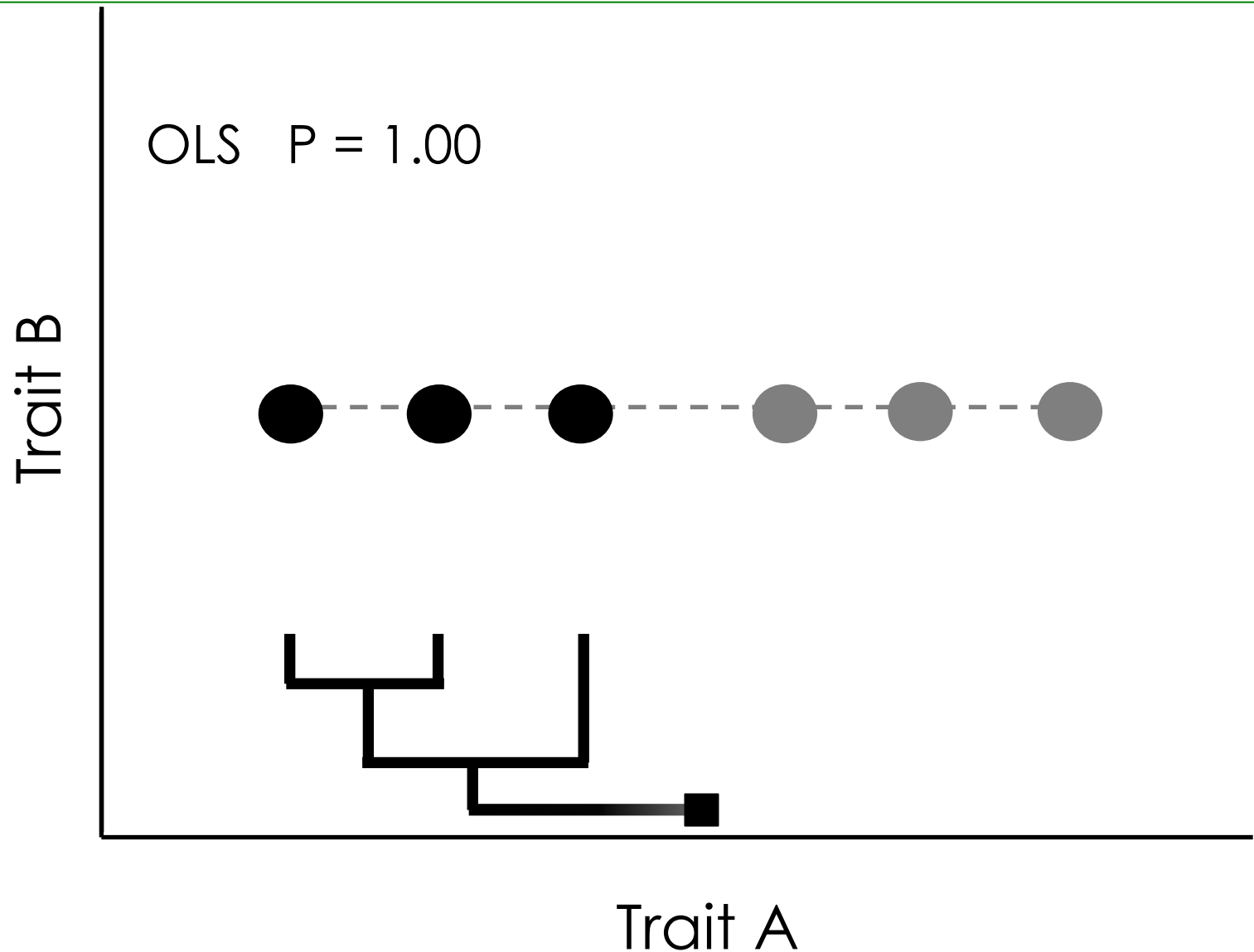


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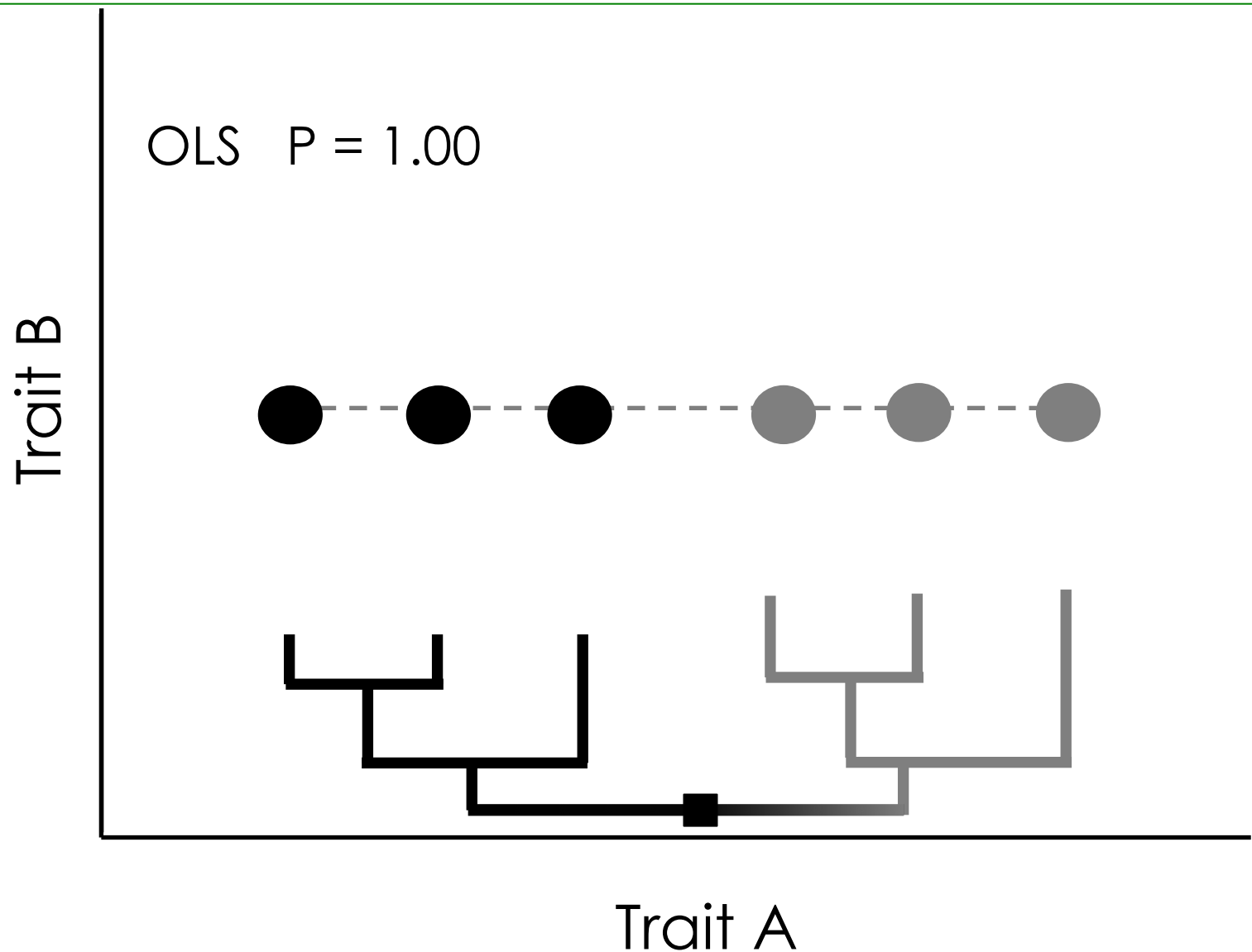


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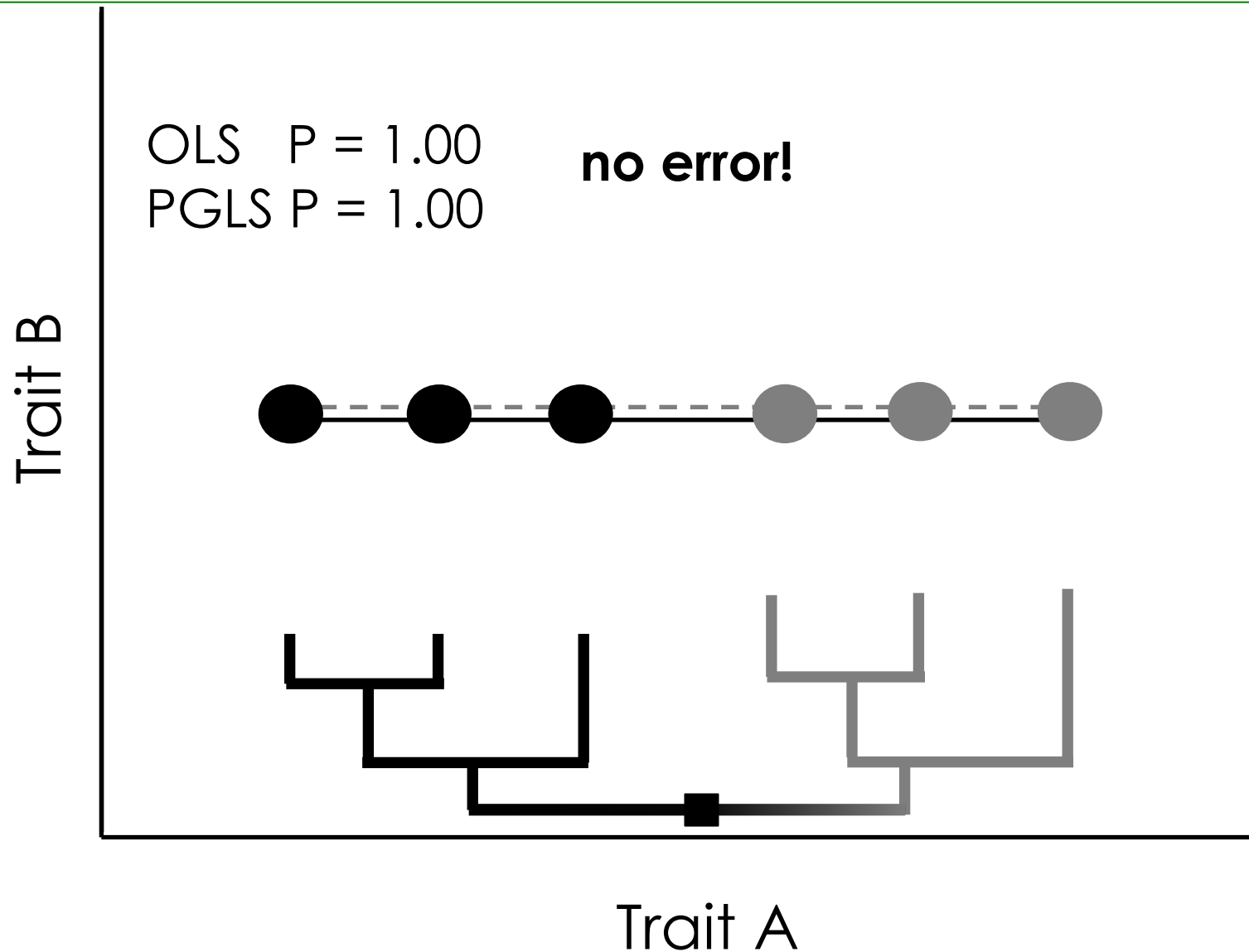


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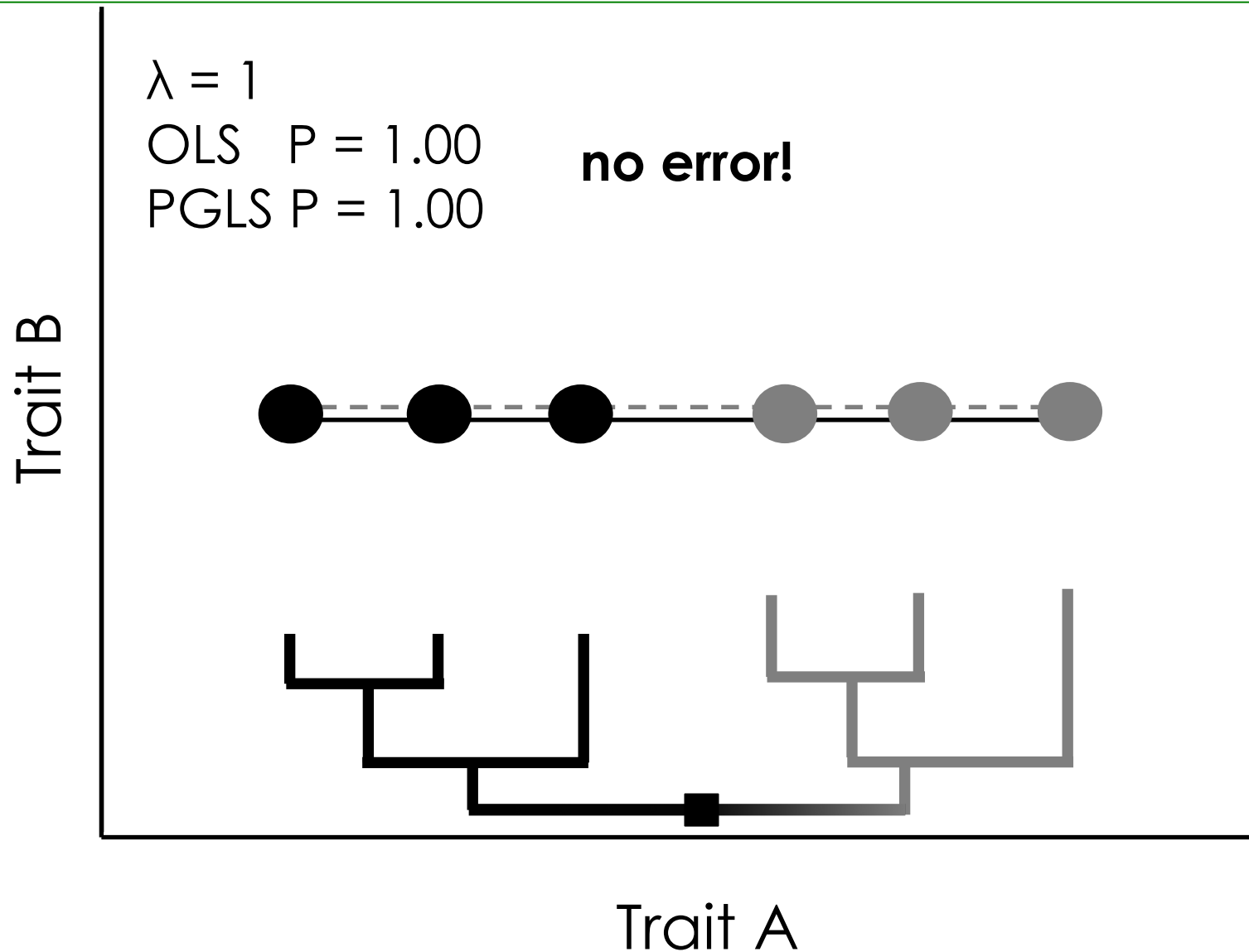


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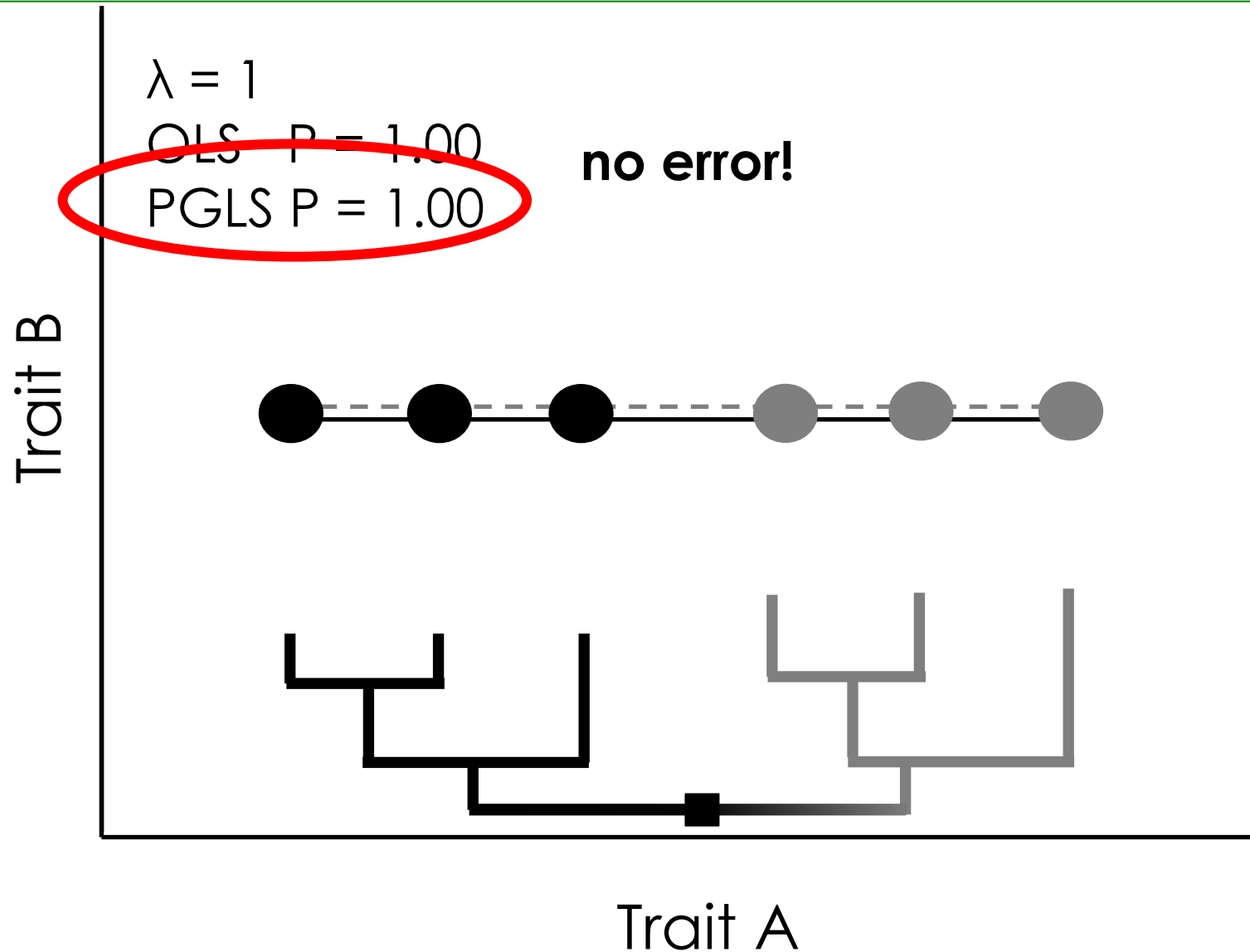


Accounting for phylogeny



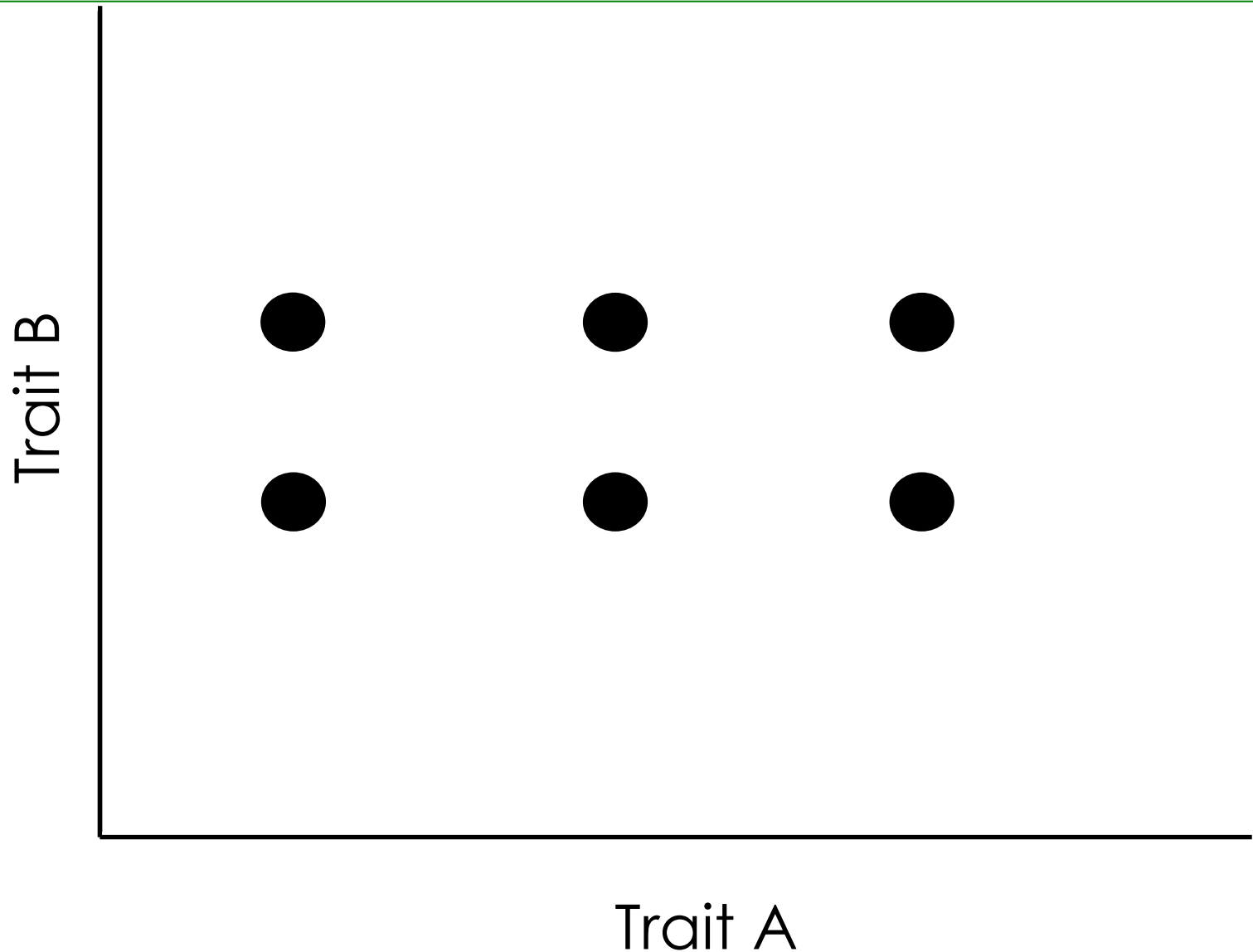


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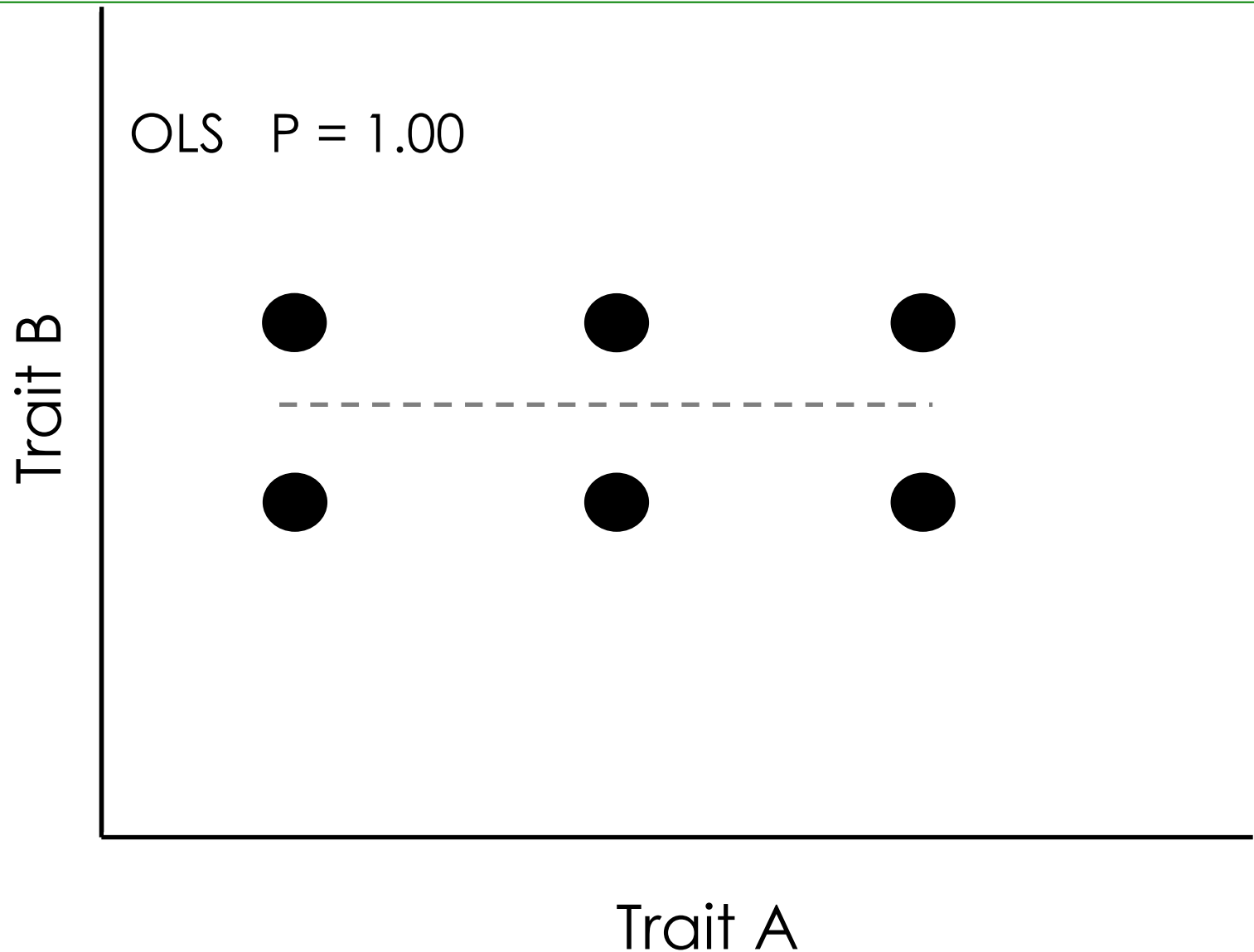
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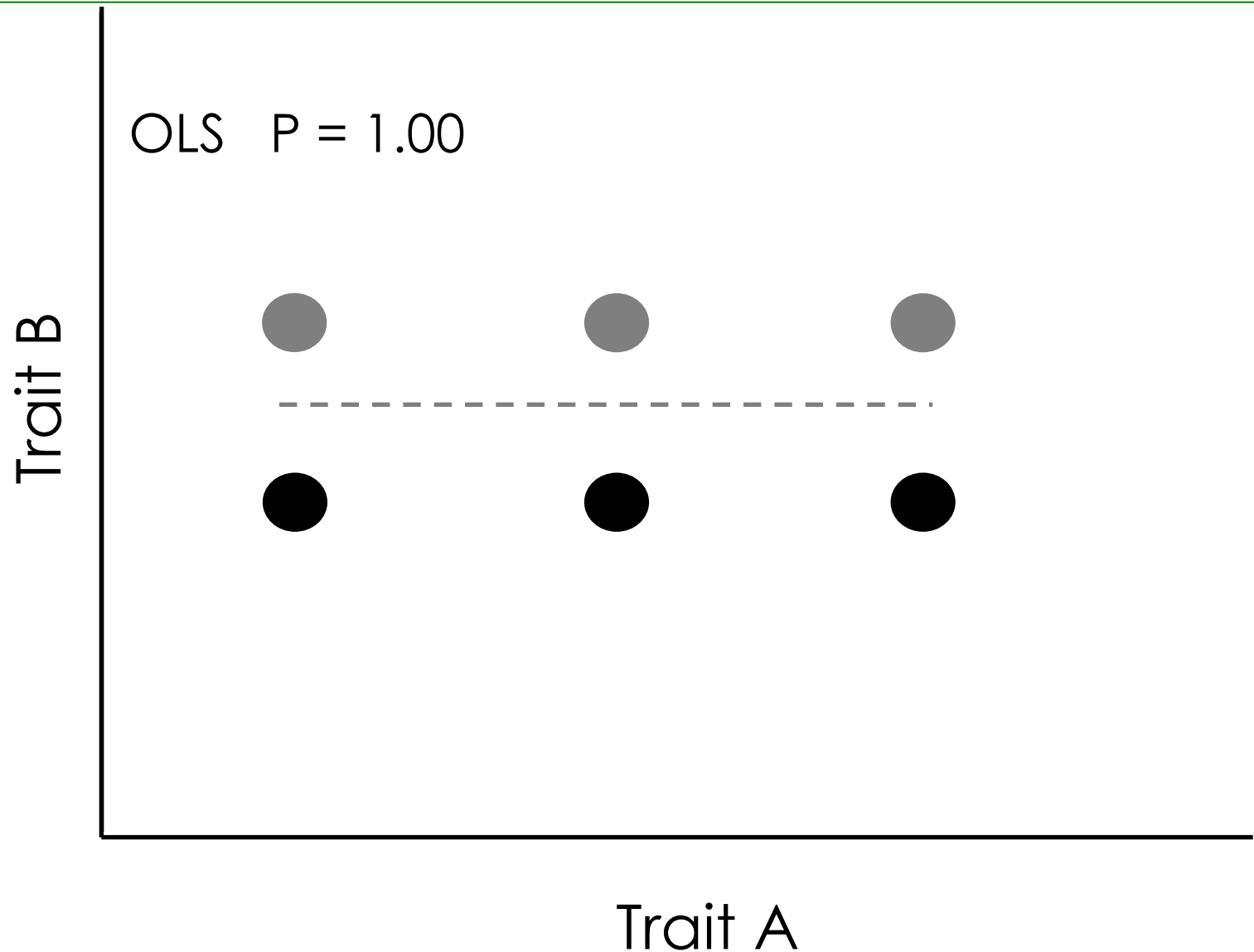


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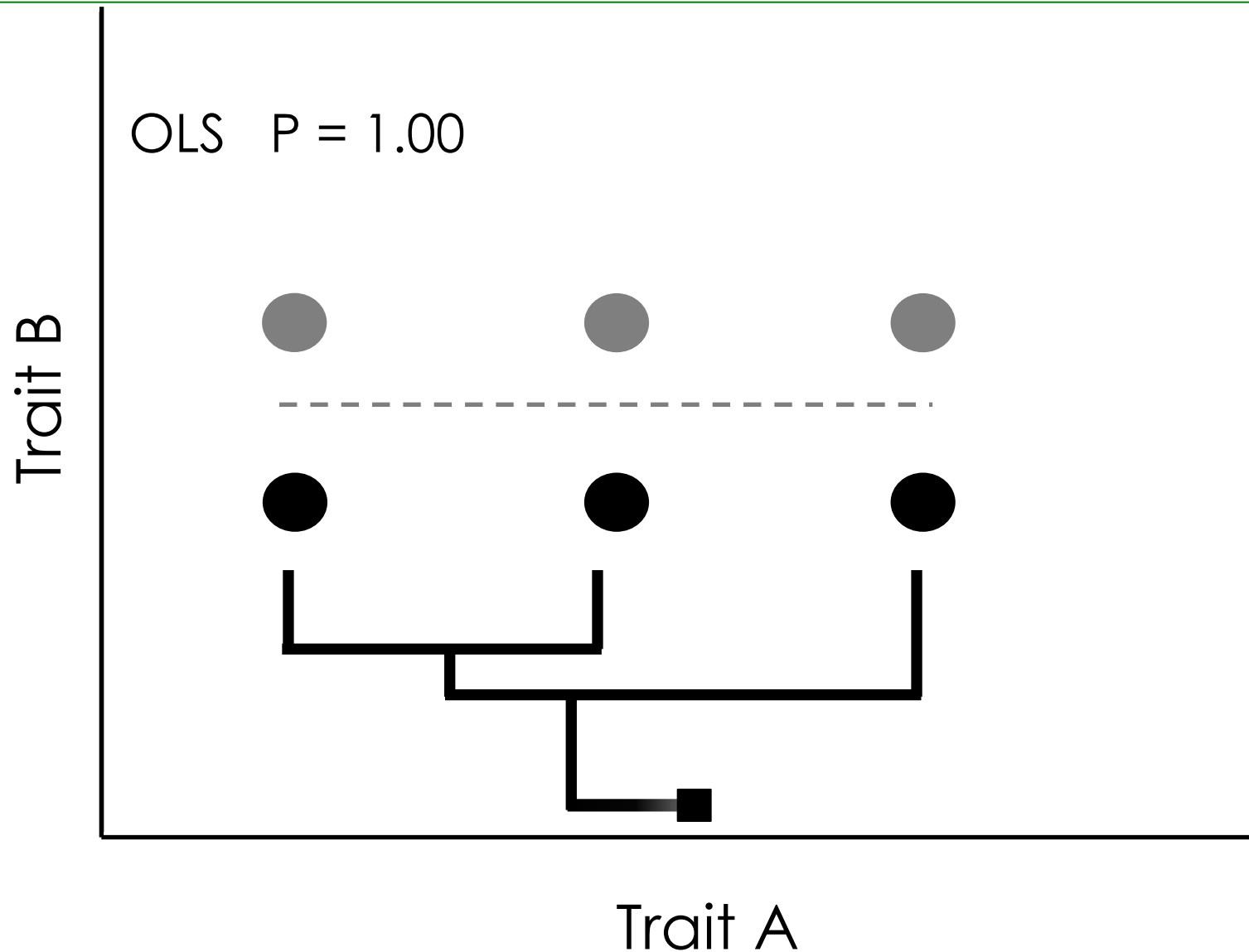


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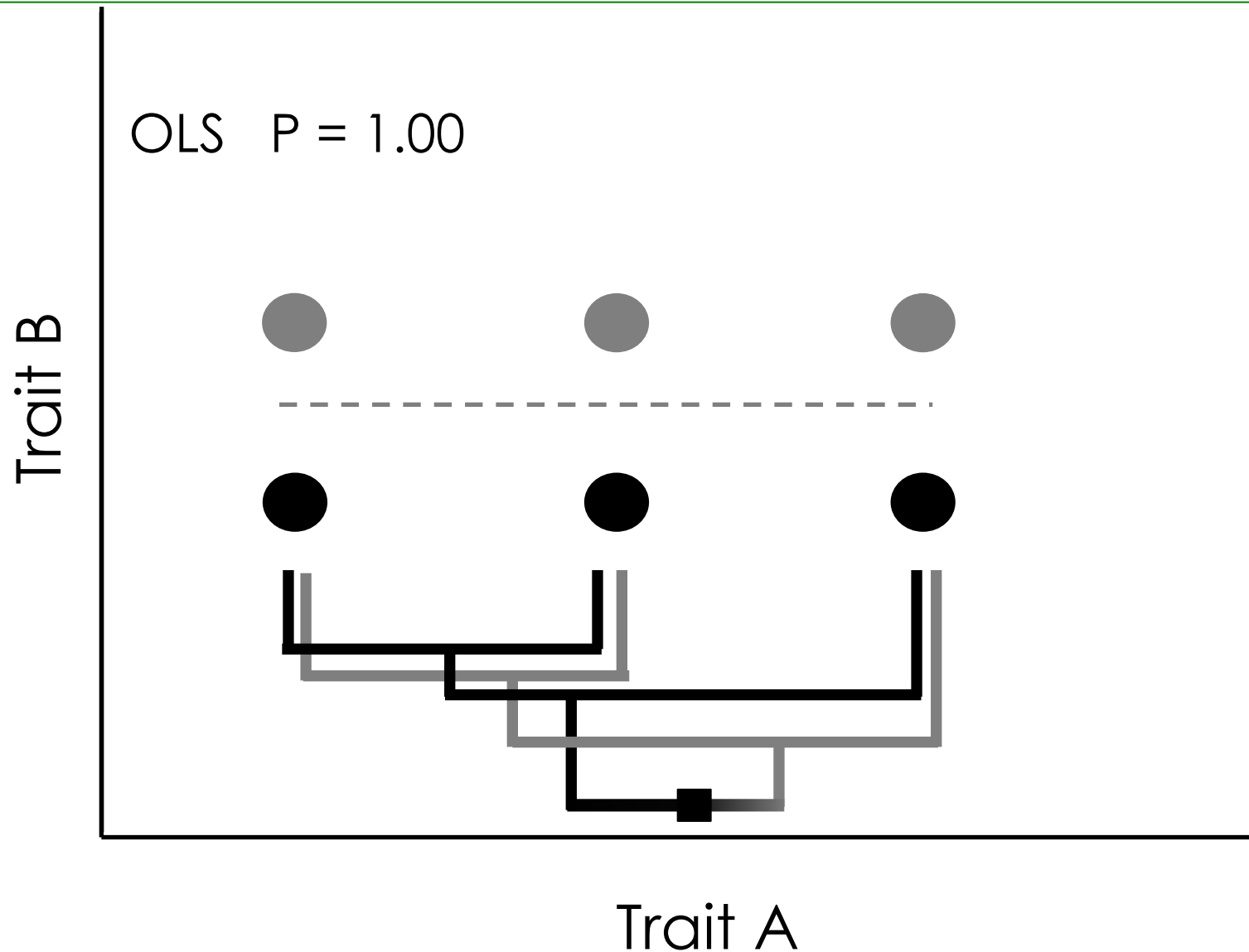


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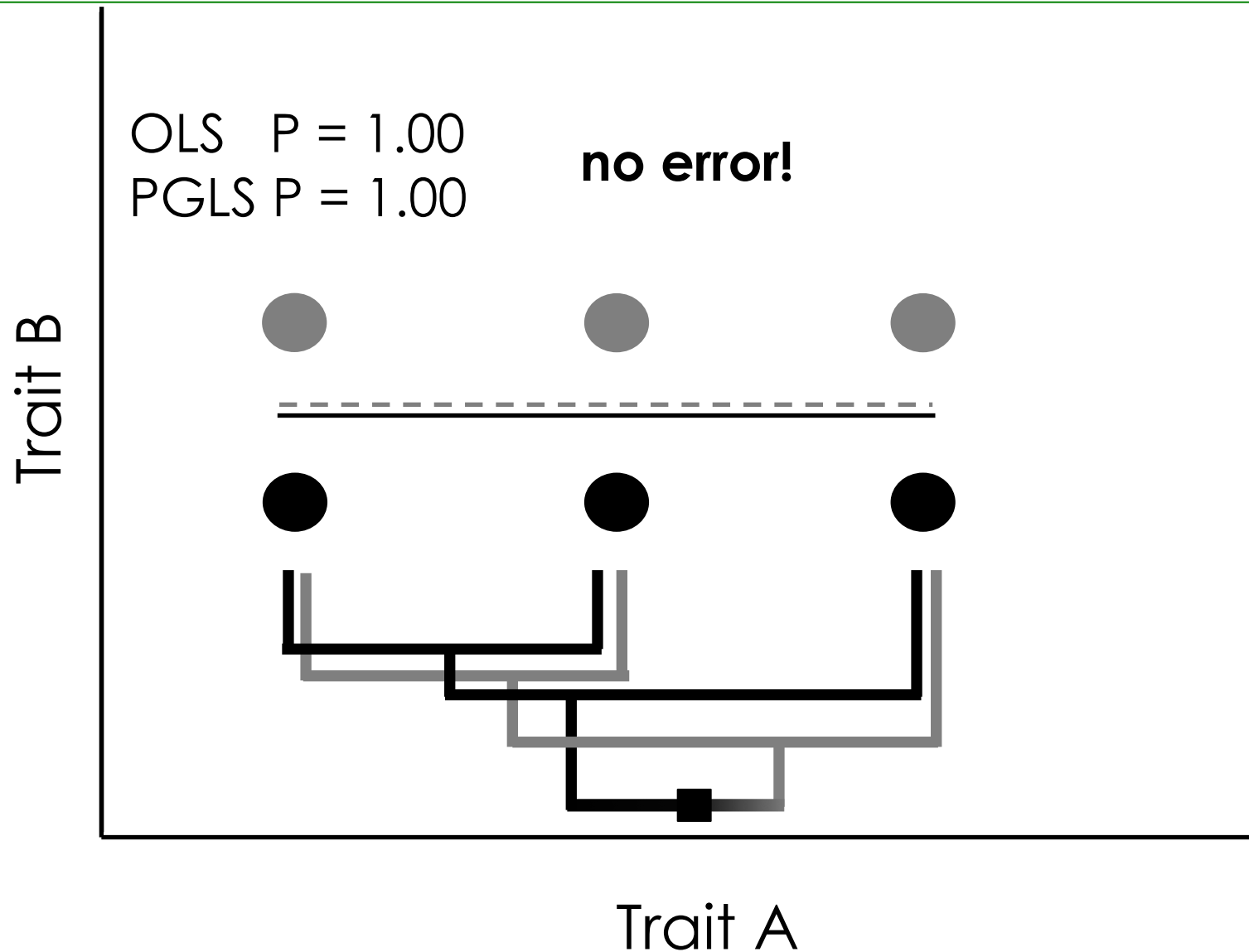


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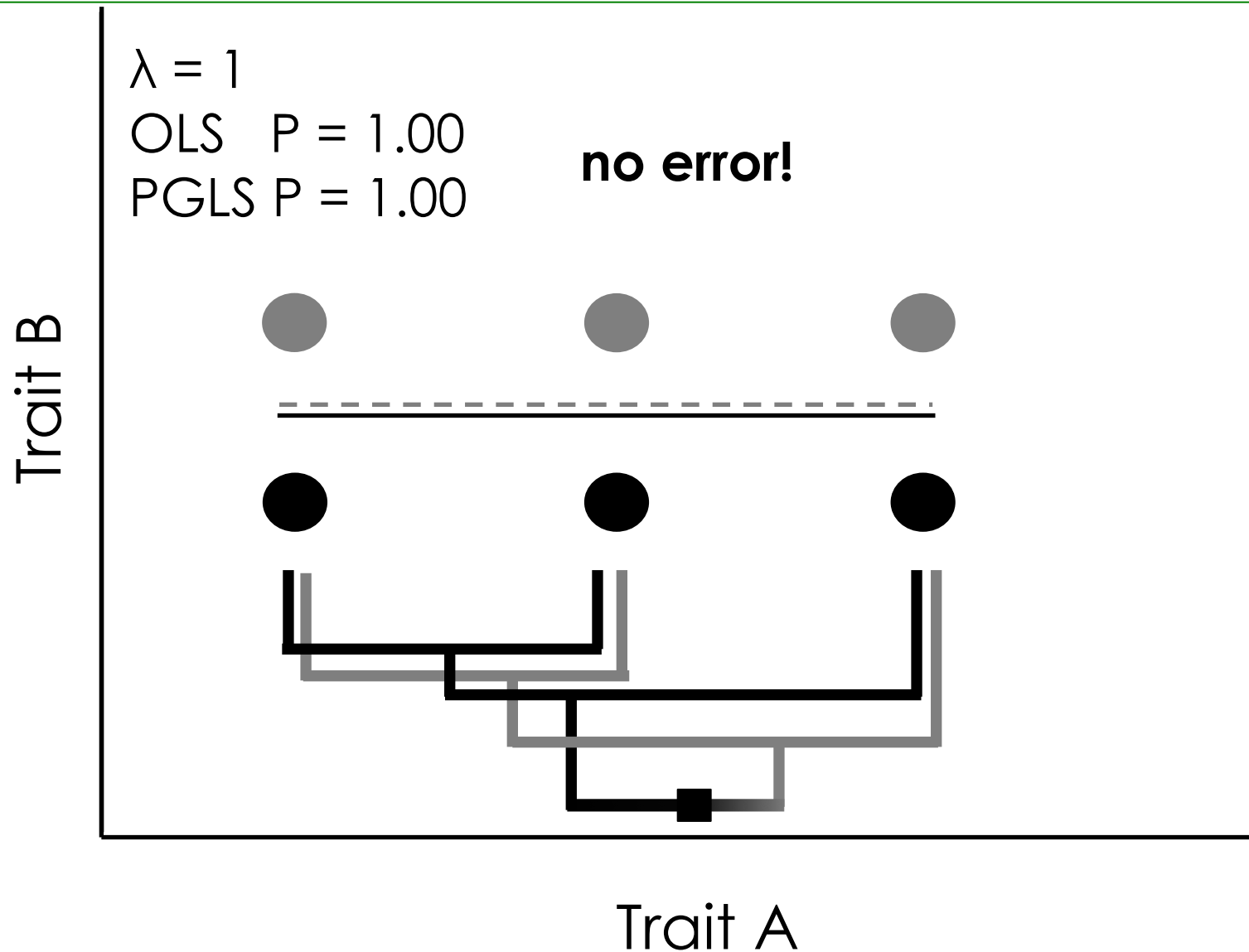


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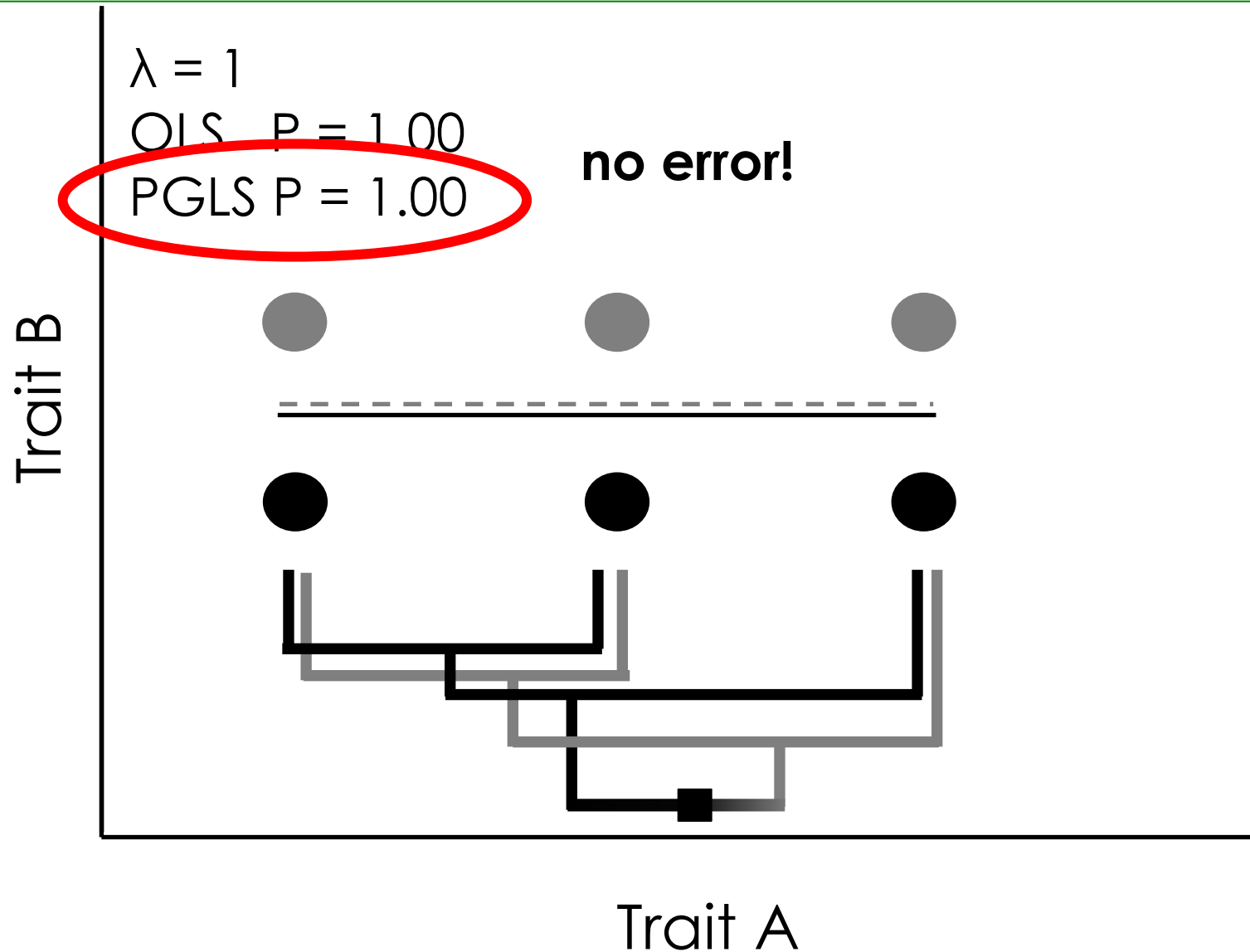


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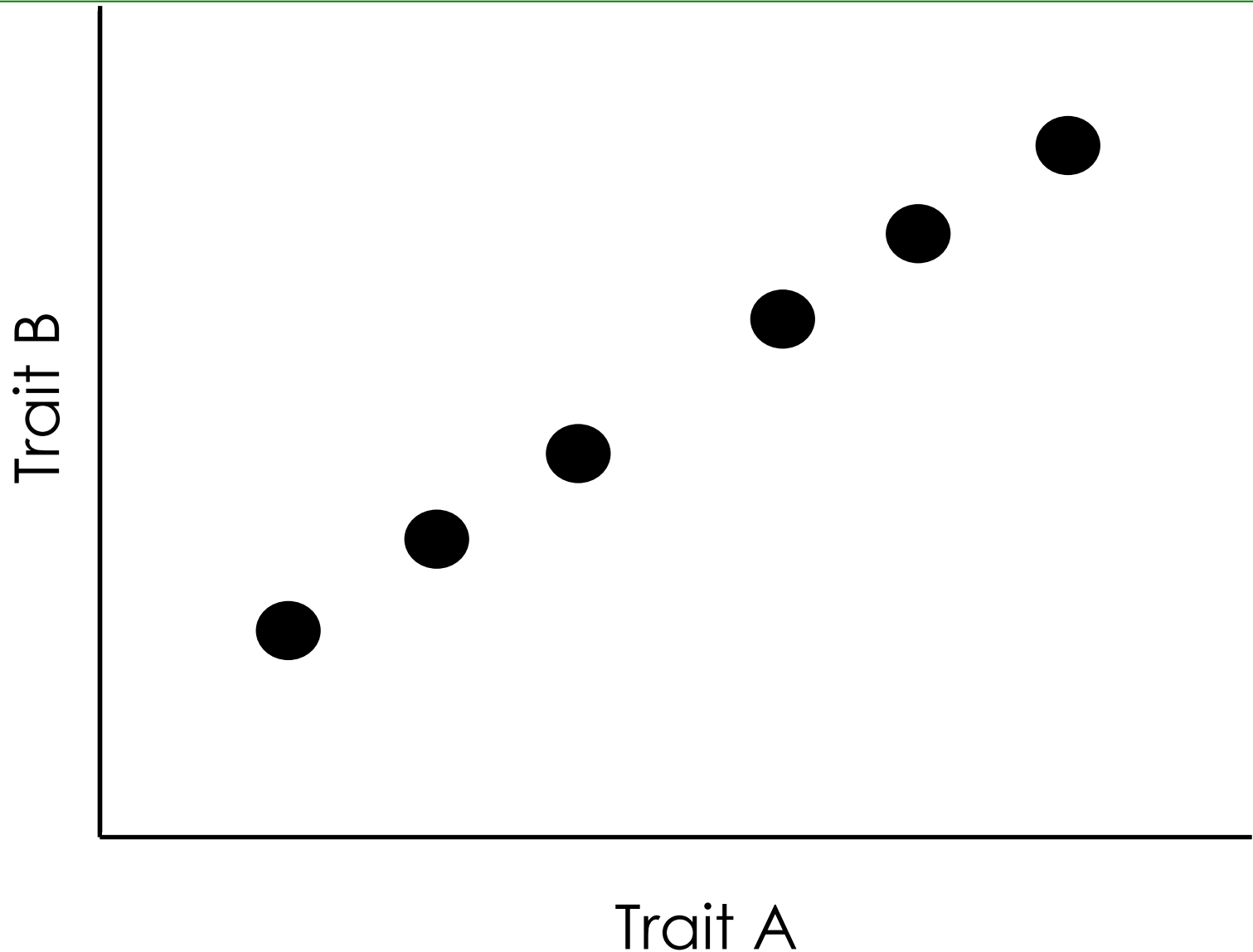


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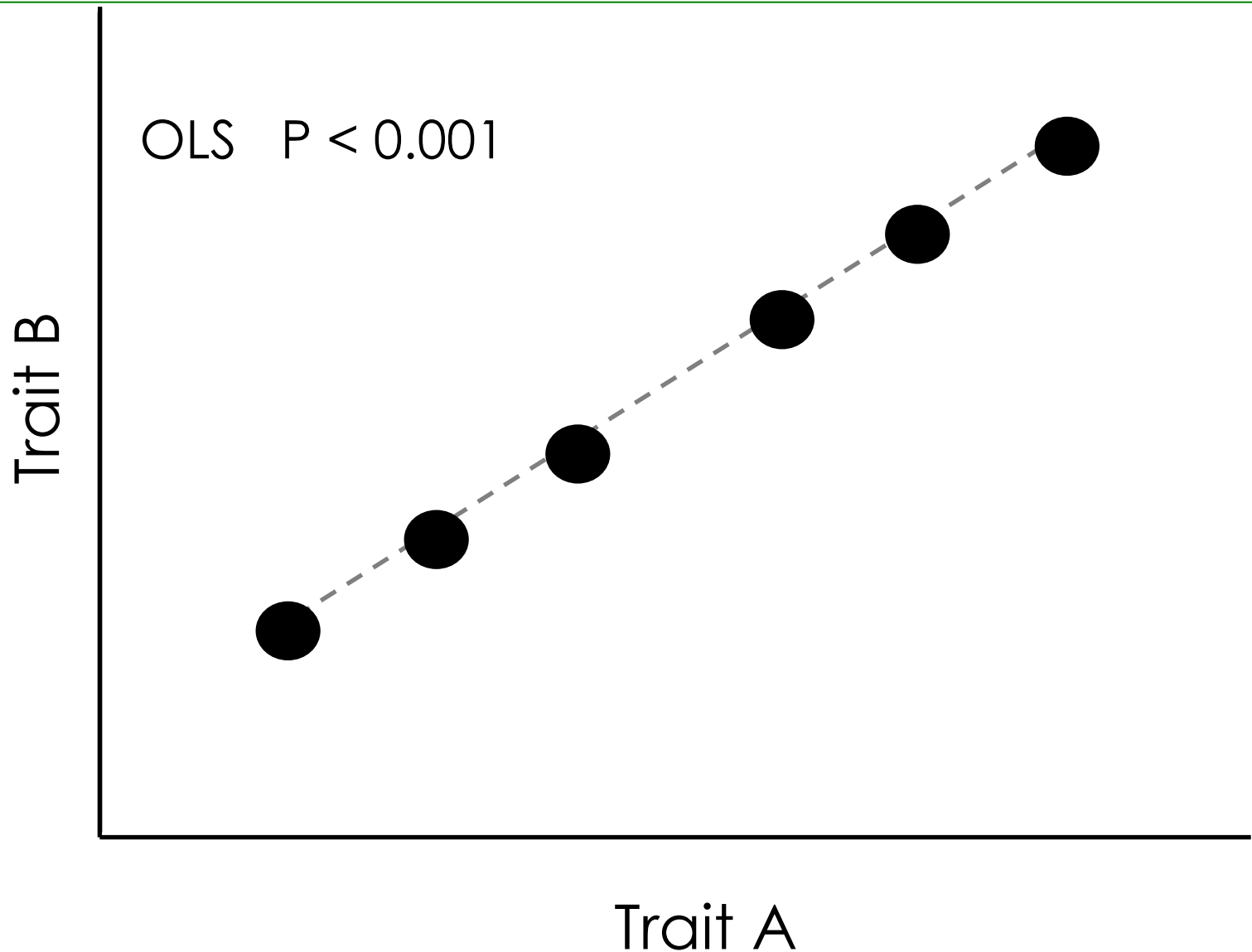
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from Dittmann et al. (2015)



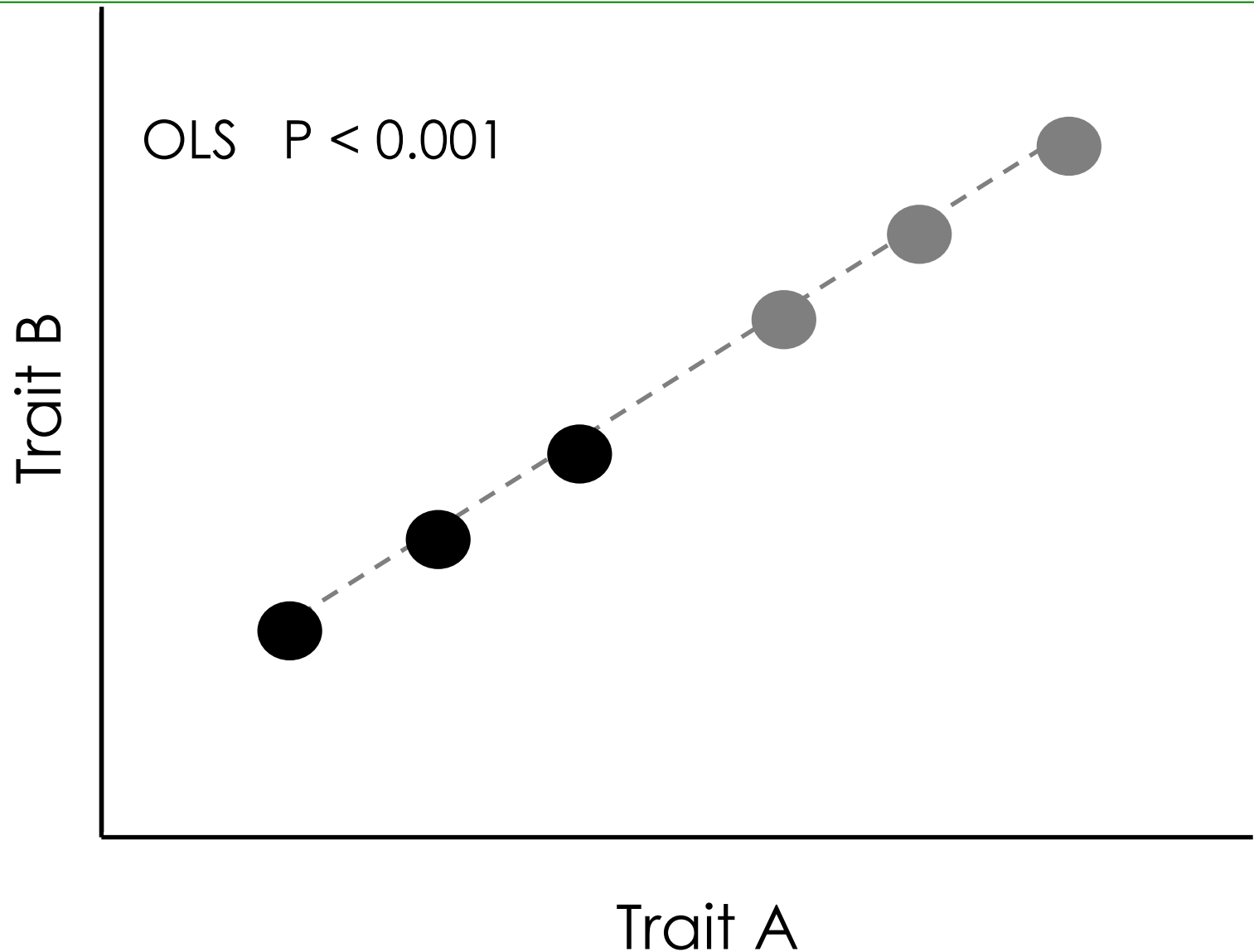
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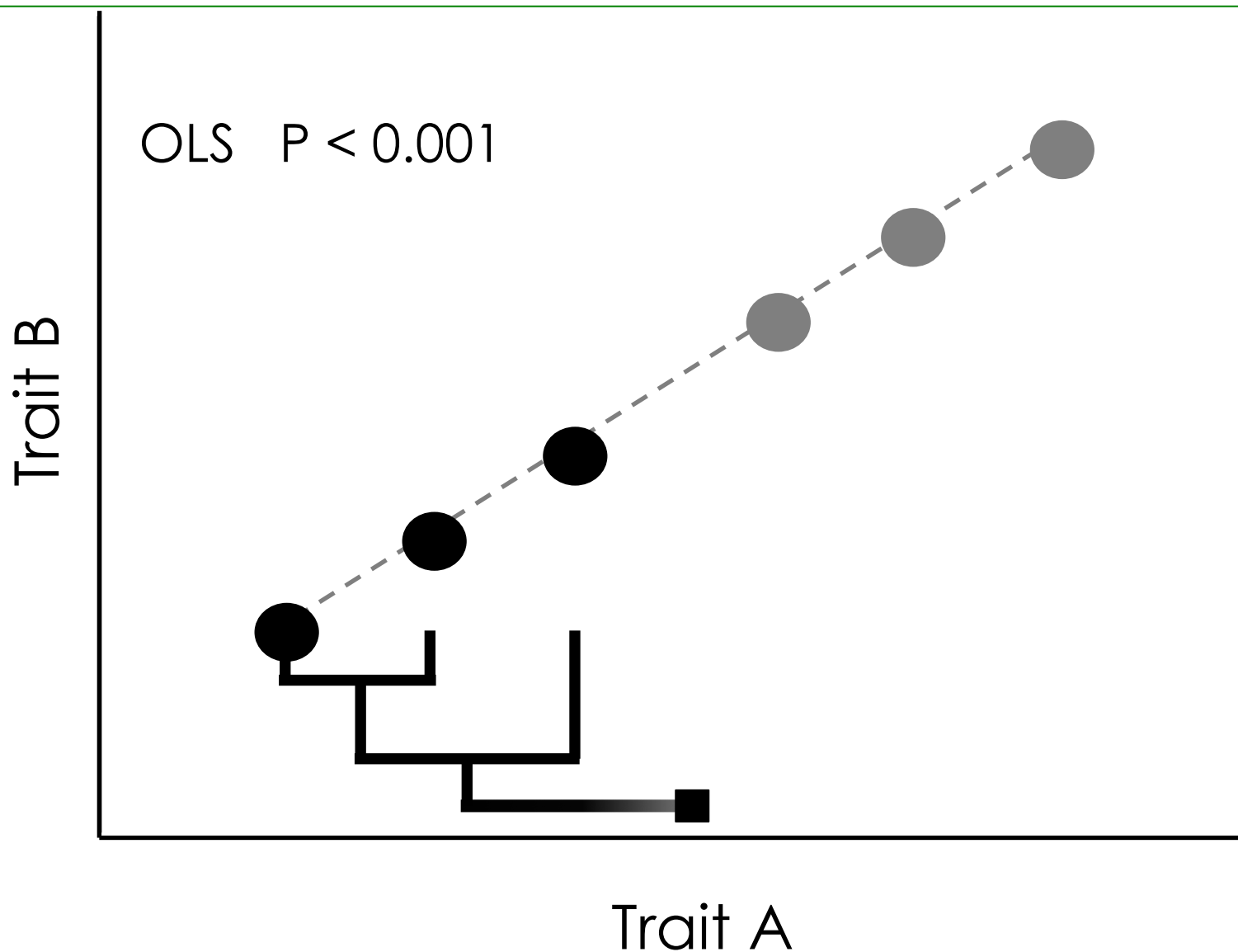
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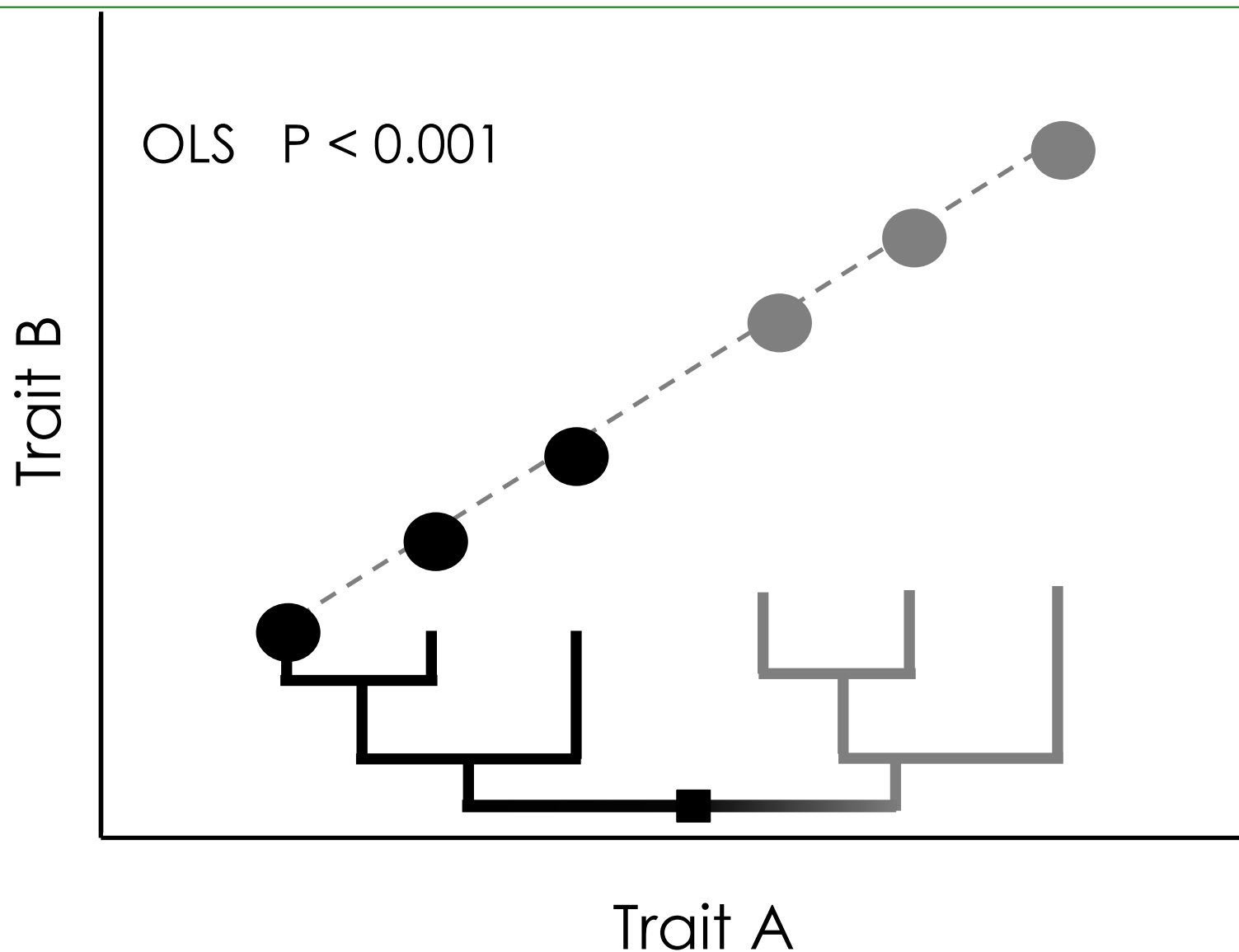
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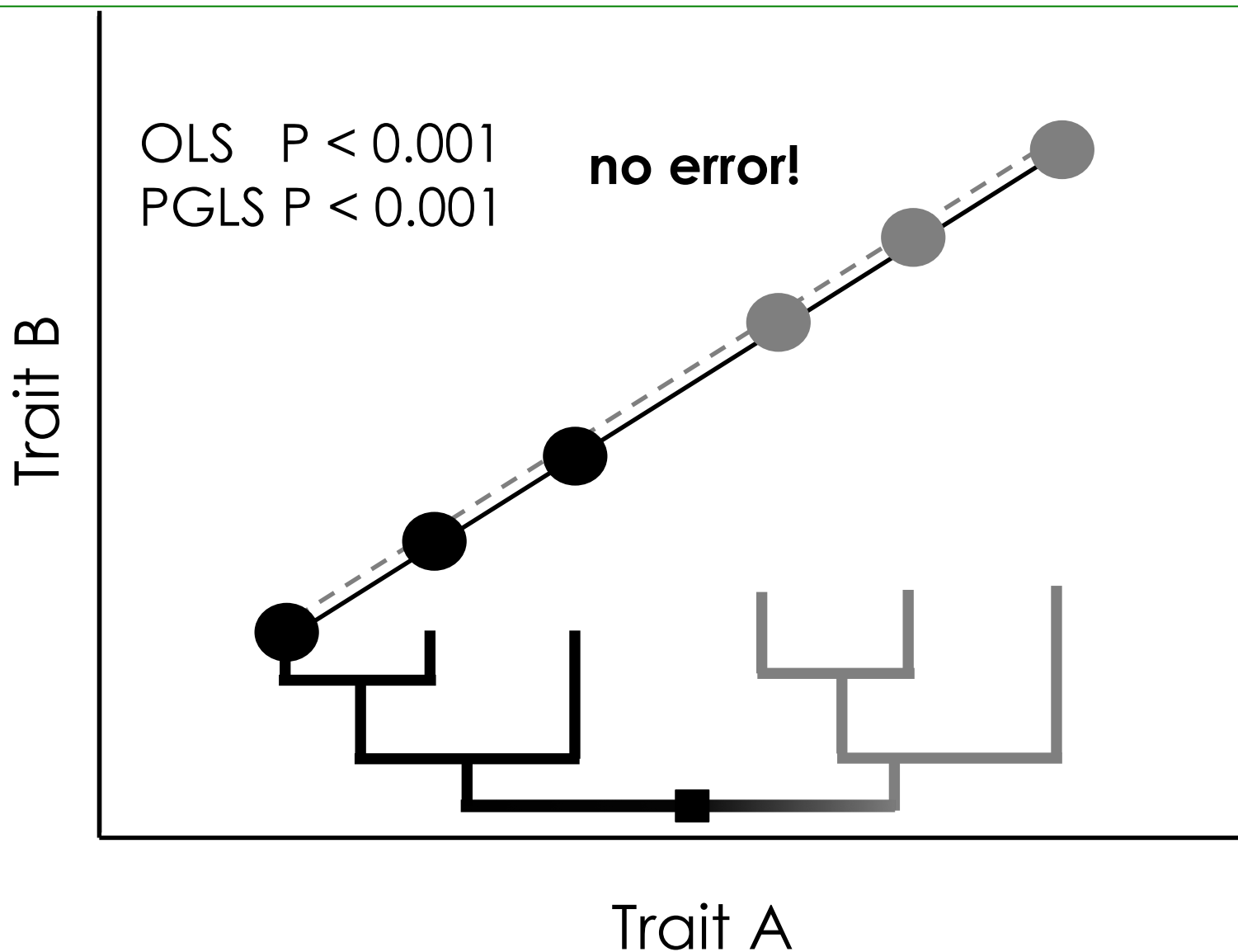


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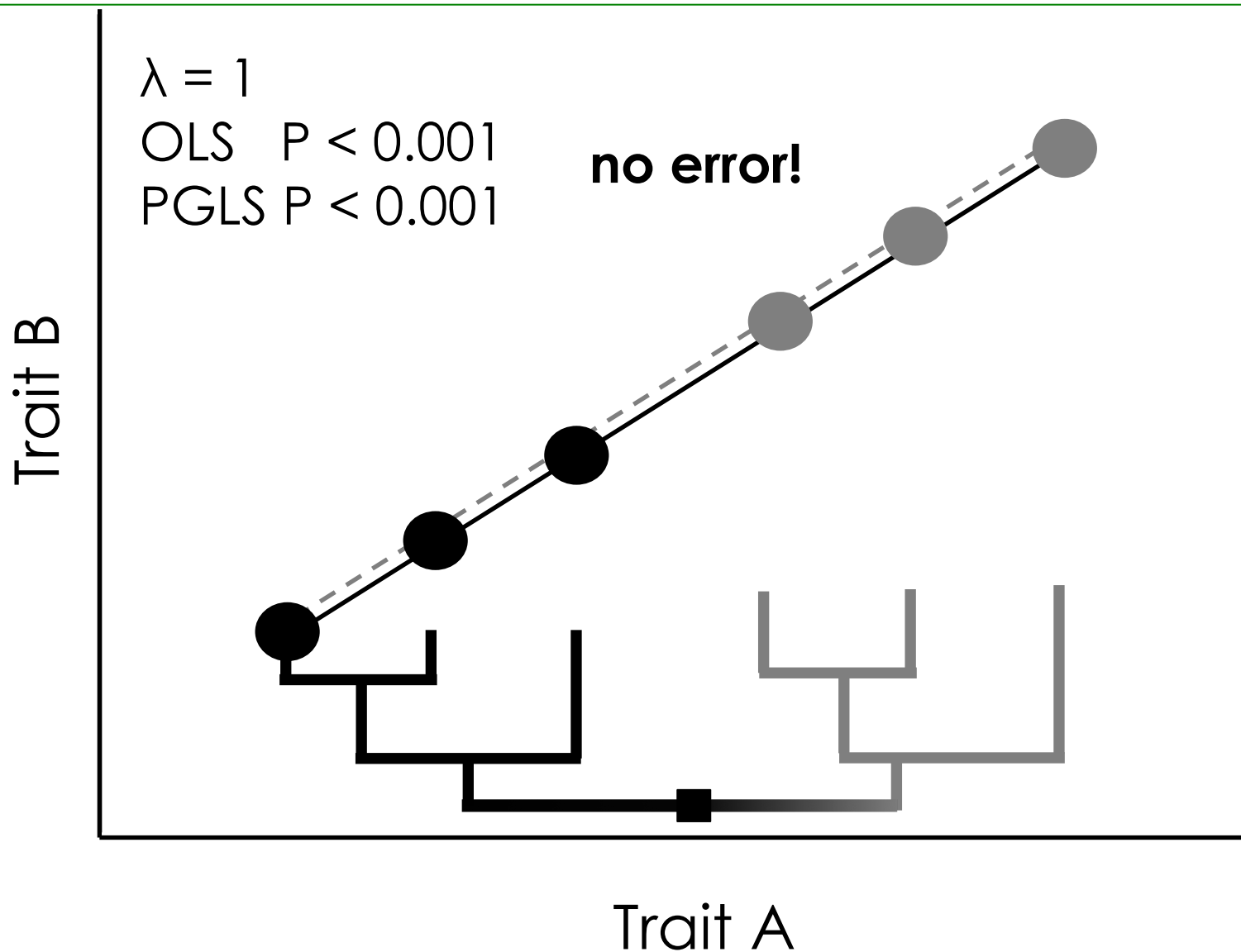


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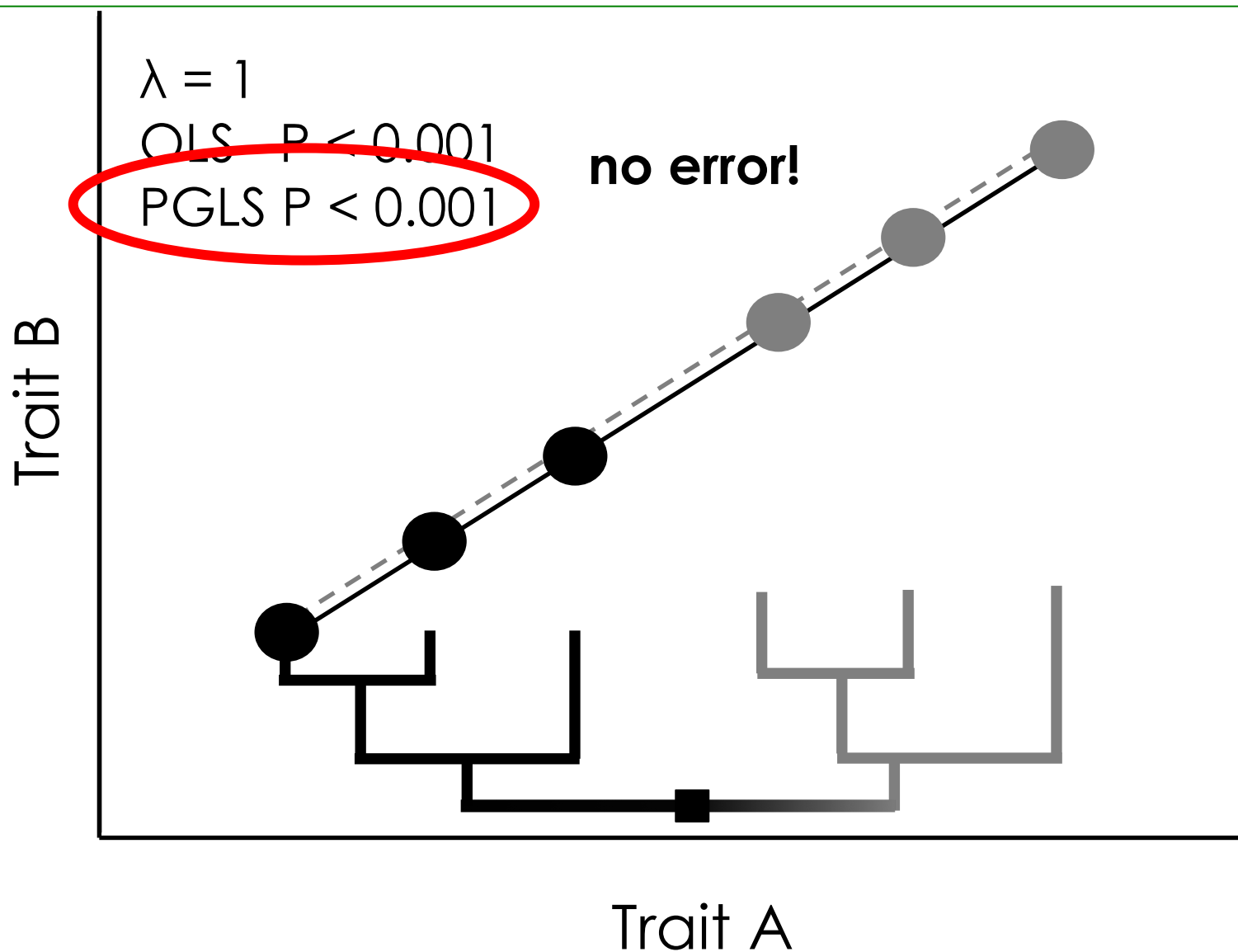


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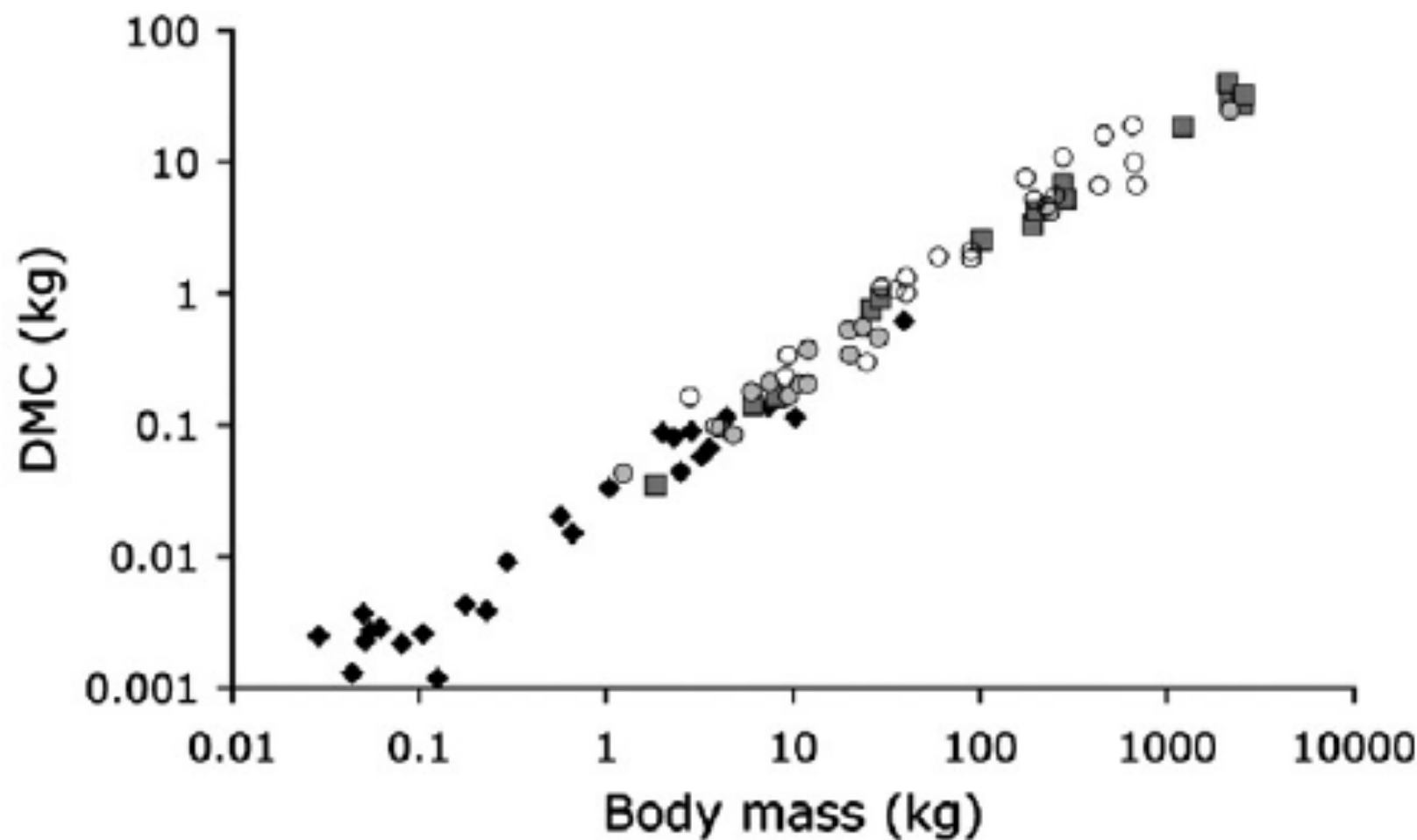


Accounting for phylogeny





Example I: gut contents





Accounting for phylogeny

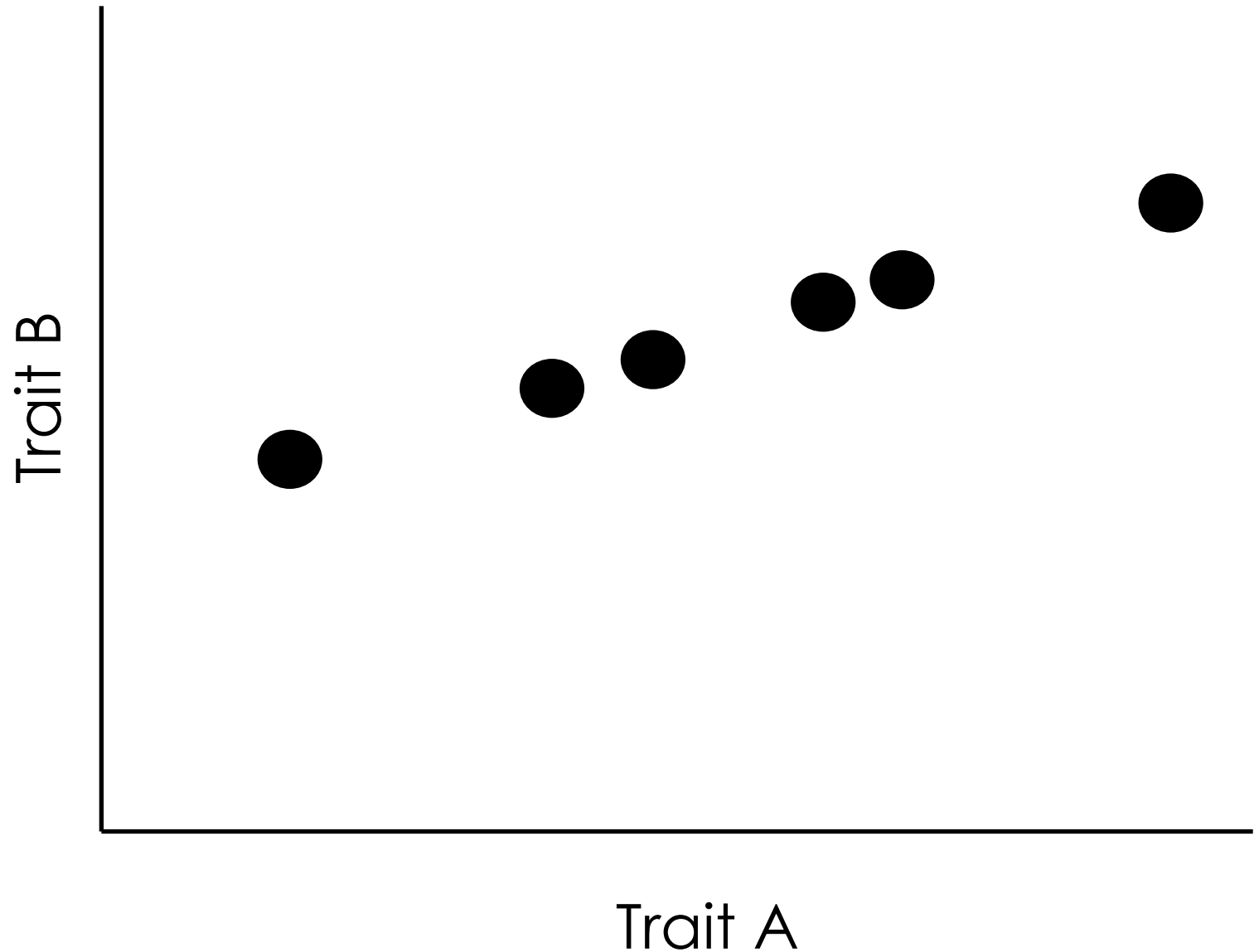
Trait B

Trait A

from Clauss et al. (2013)



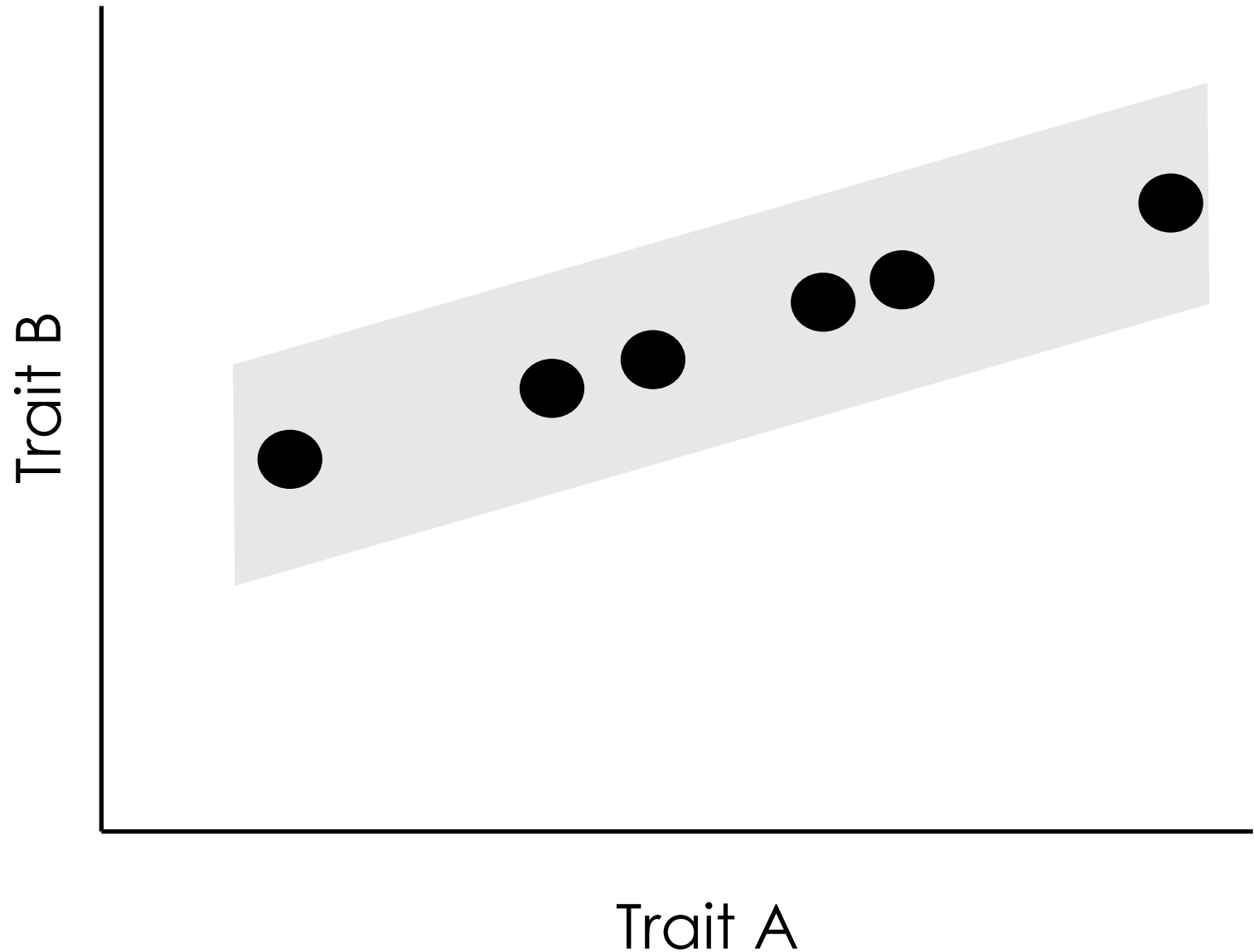
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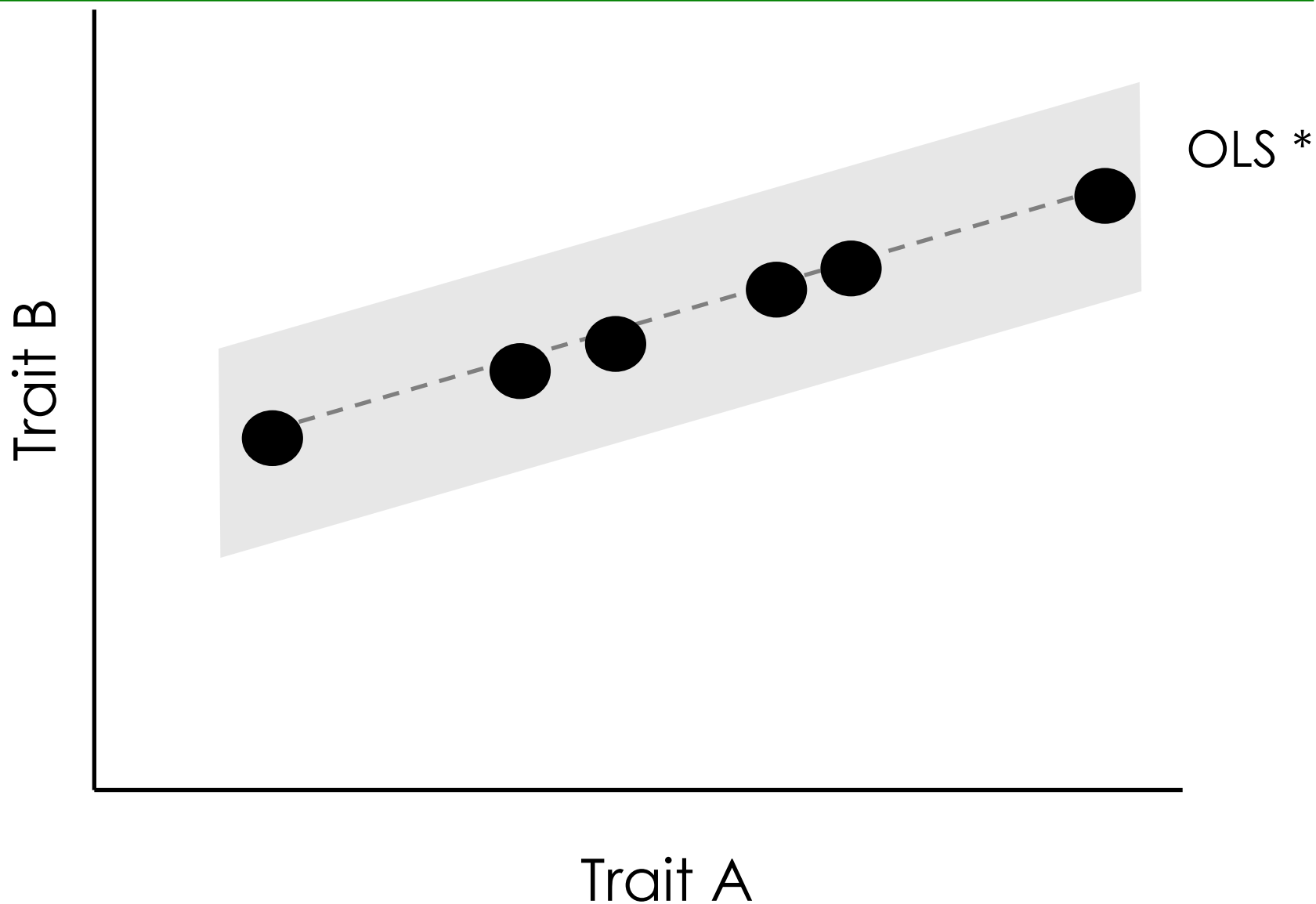
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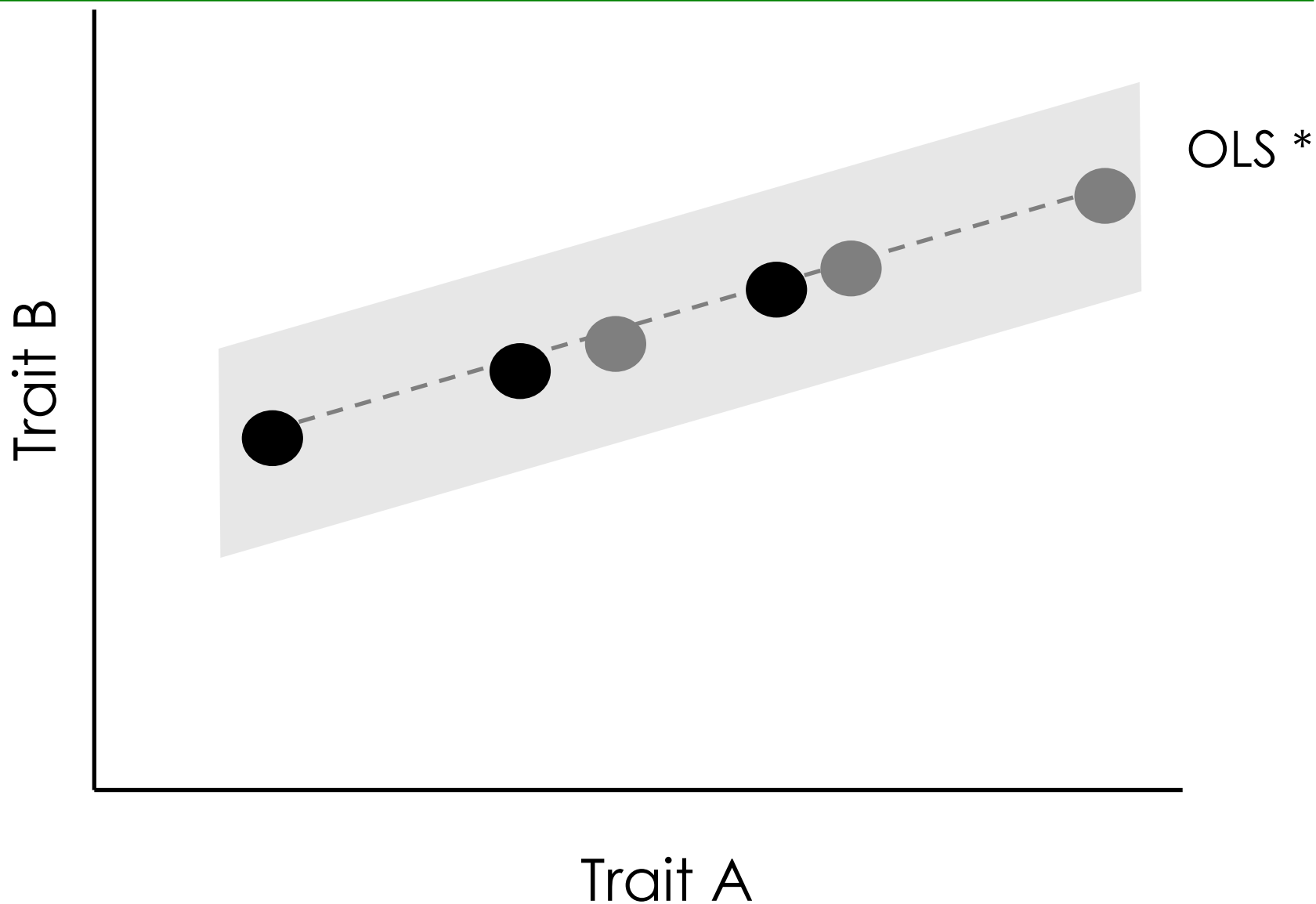
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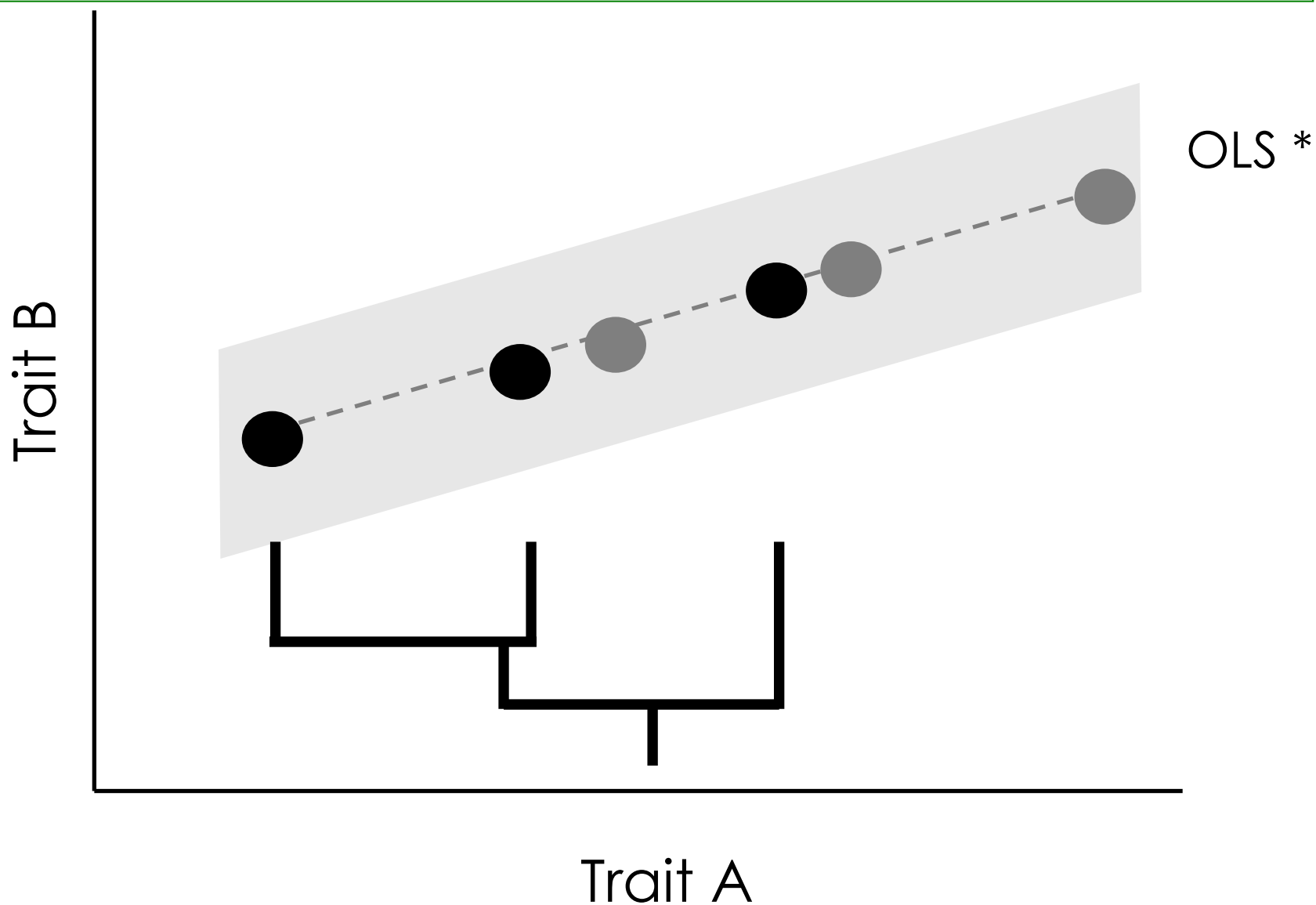
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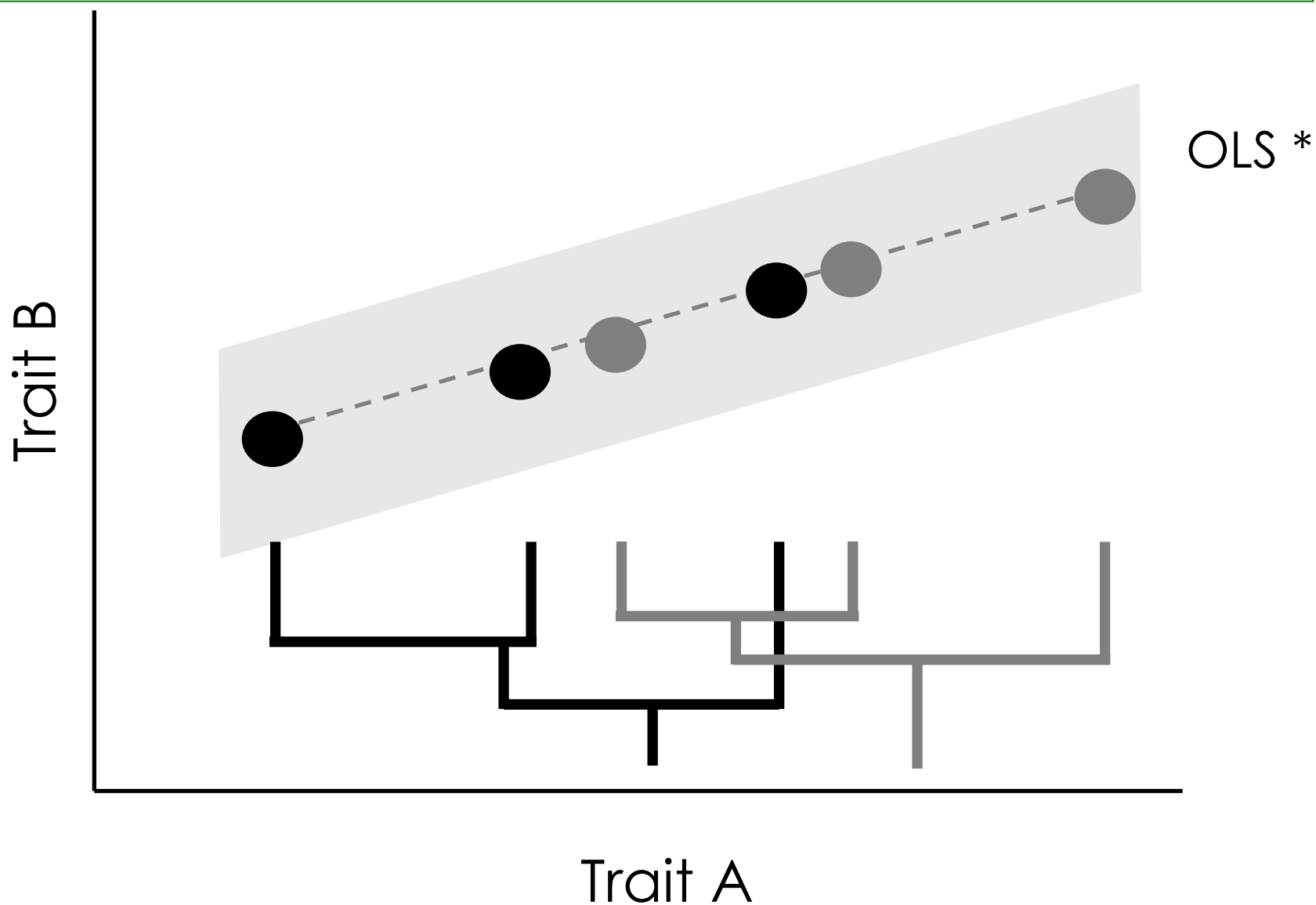
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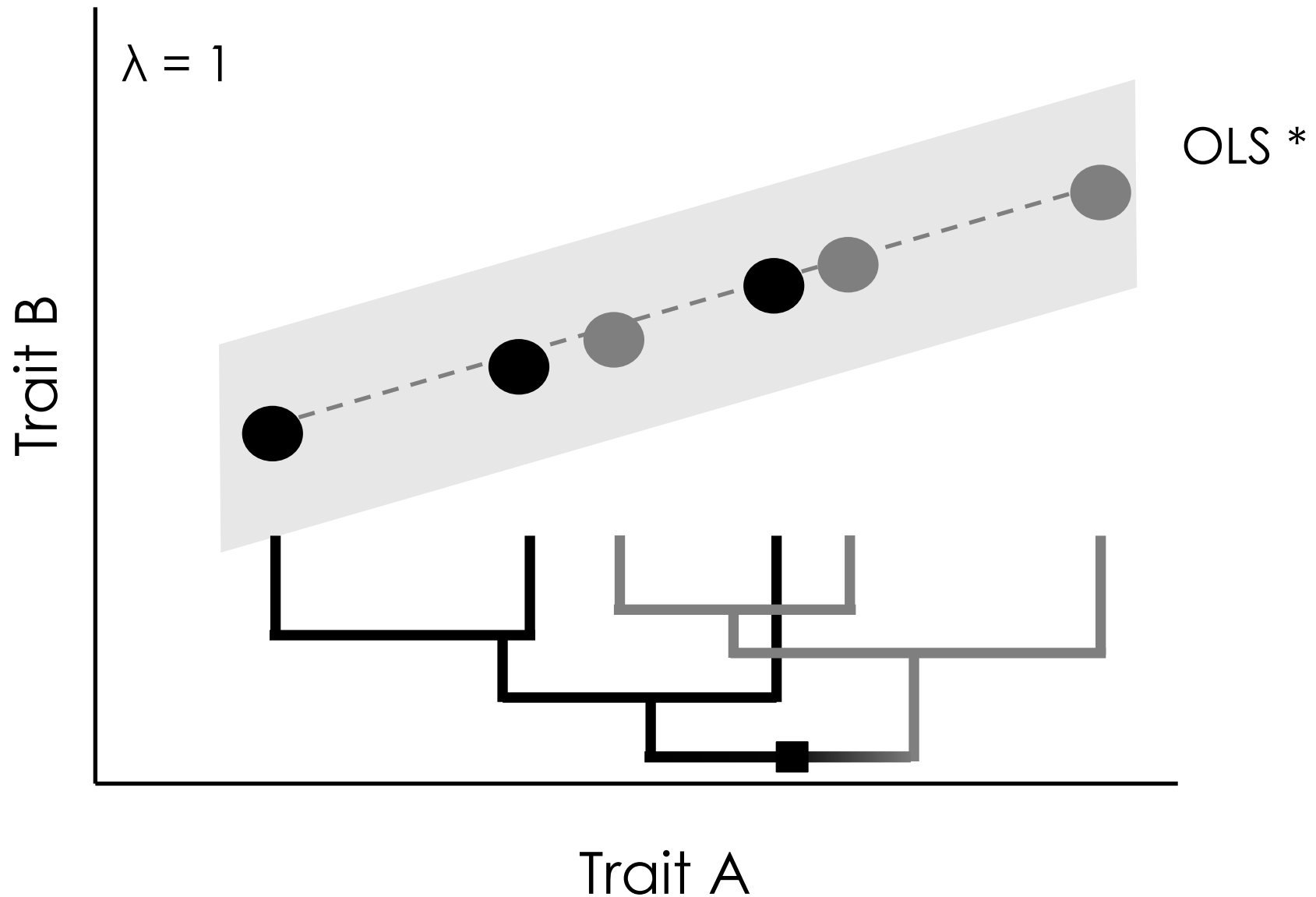
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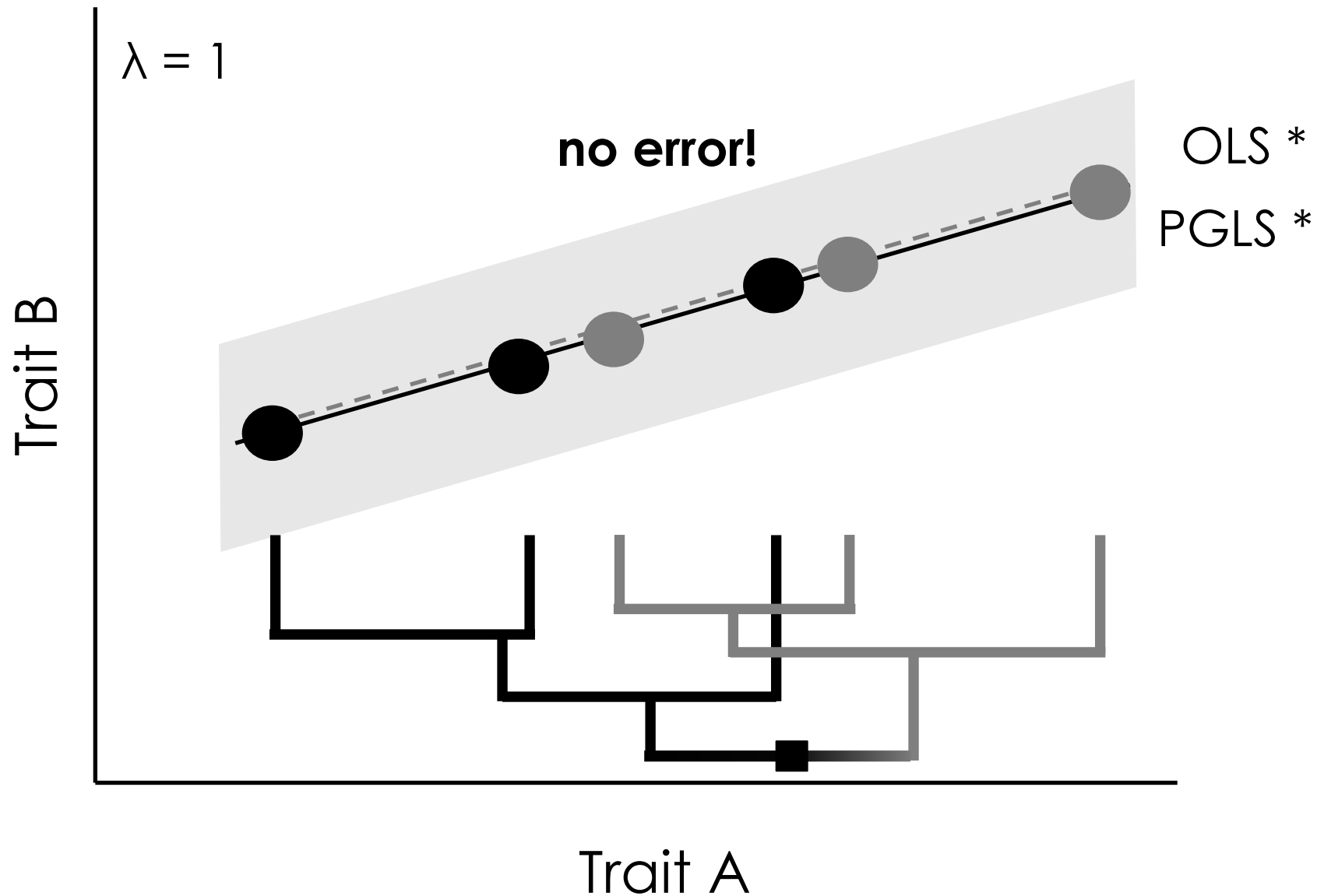


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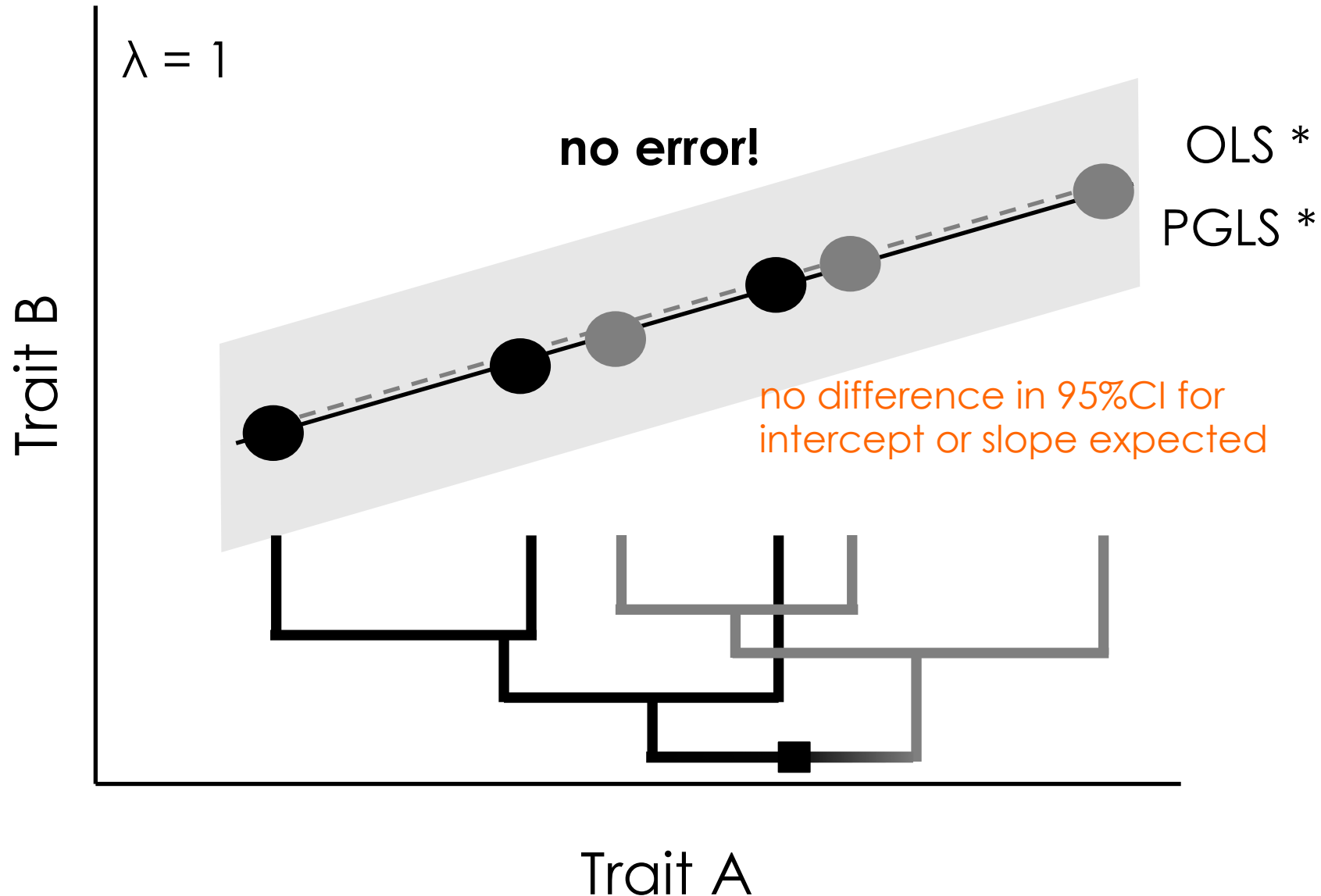


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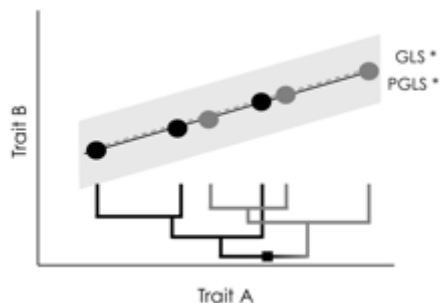


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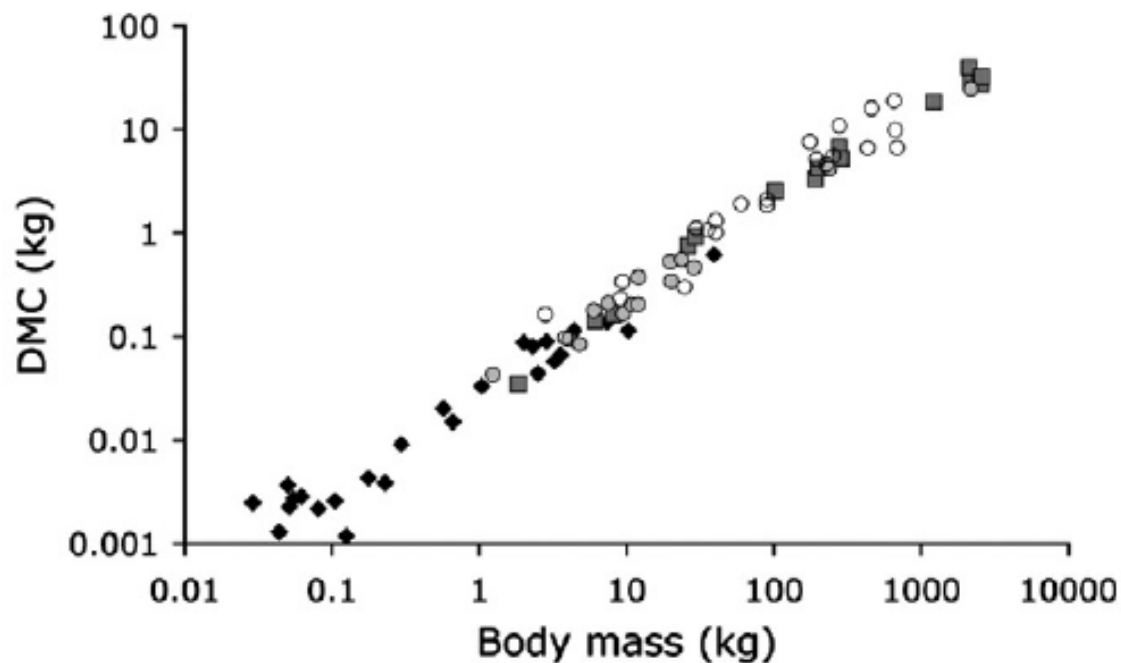


Example I: gut contents



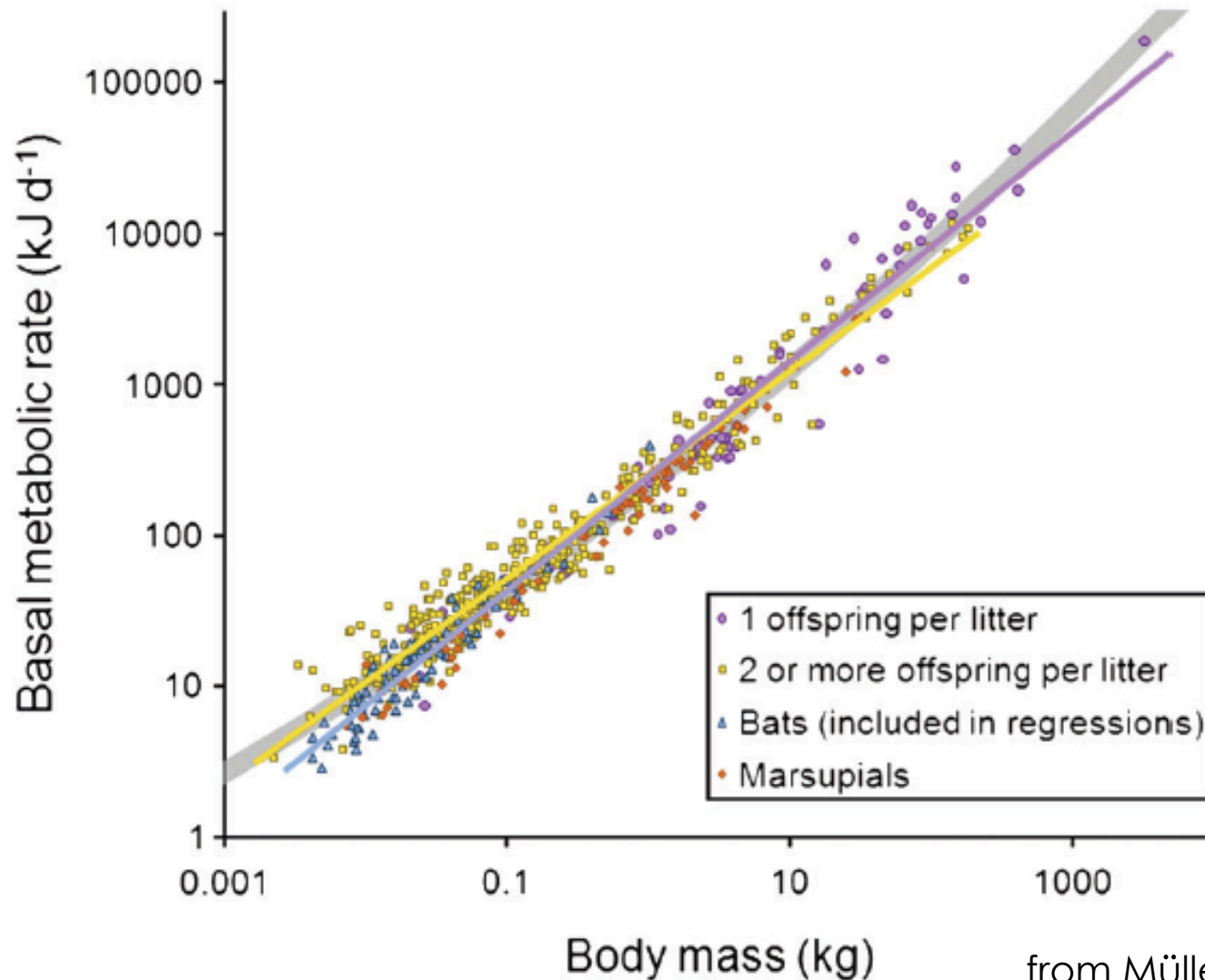
OLS: 0.03 (0.025-0.032) $BM^{0.93}$ (0.90-0.96)

PGLS: 0.03 (0.010-0.075) $BM^{0.92}$ (0.85-0.98)





Example II: basal metabolic rate



from Müller et al. (2012)



Accounting for phylogeny

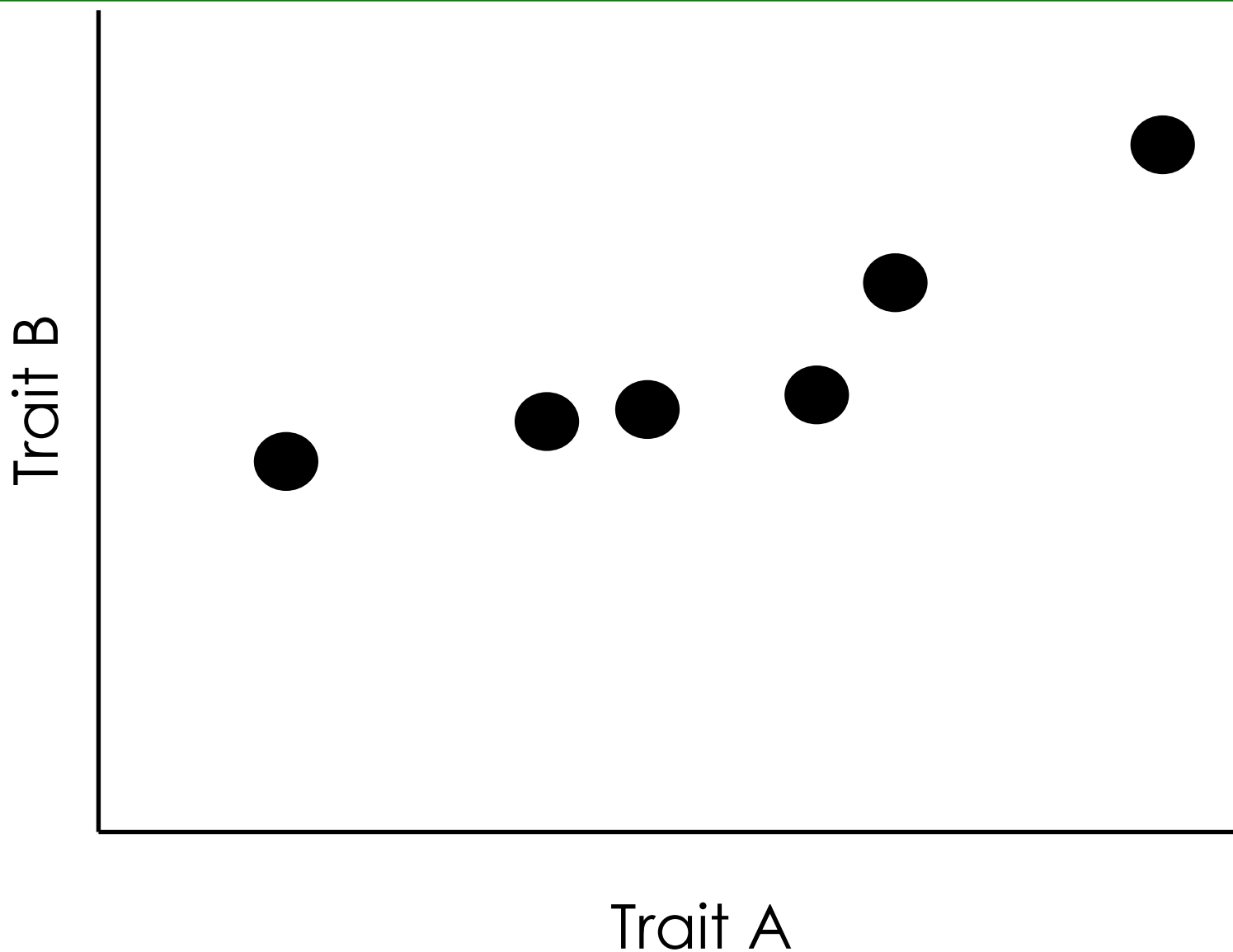
Trait B

Trait A

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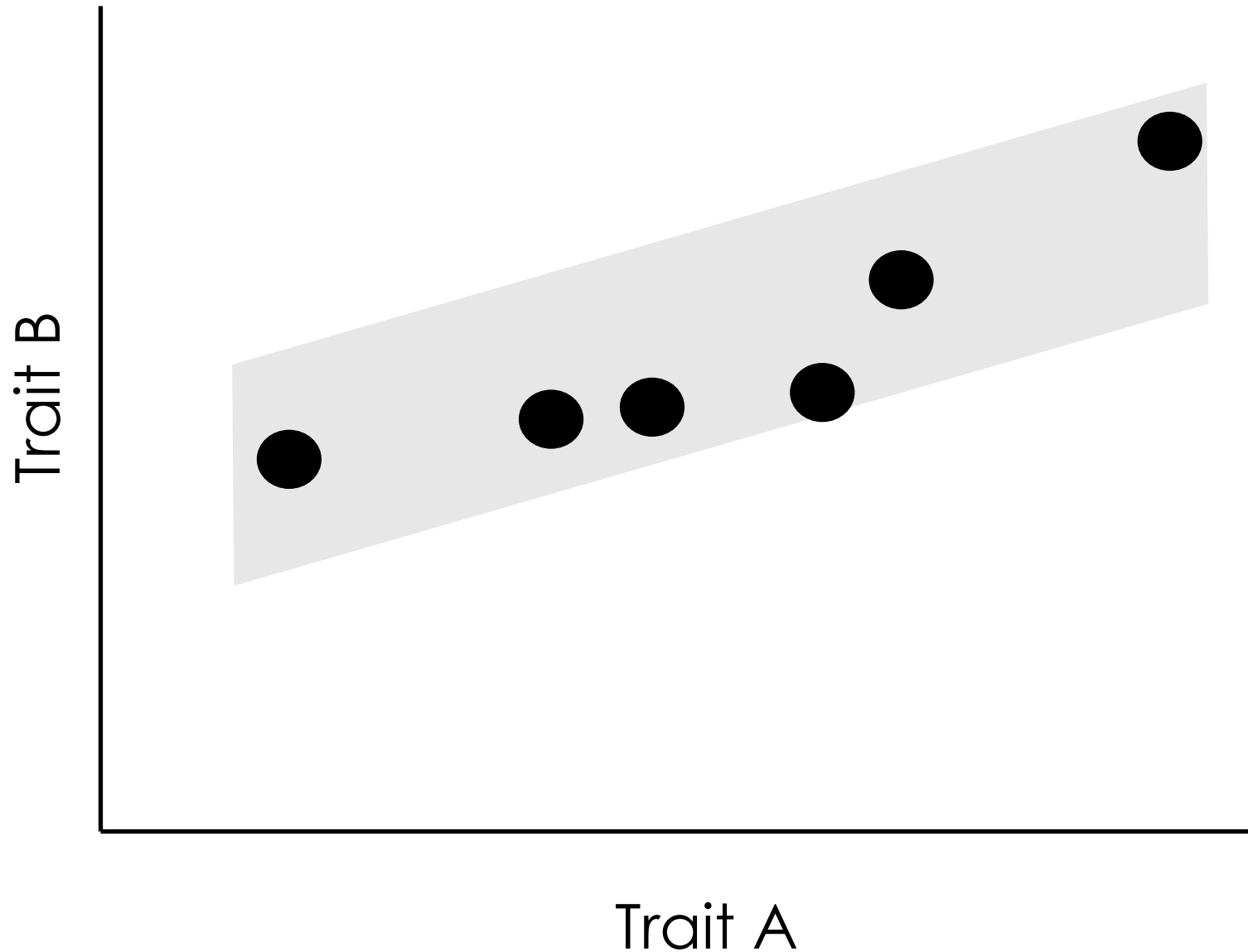
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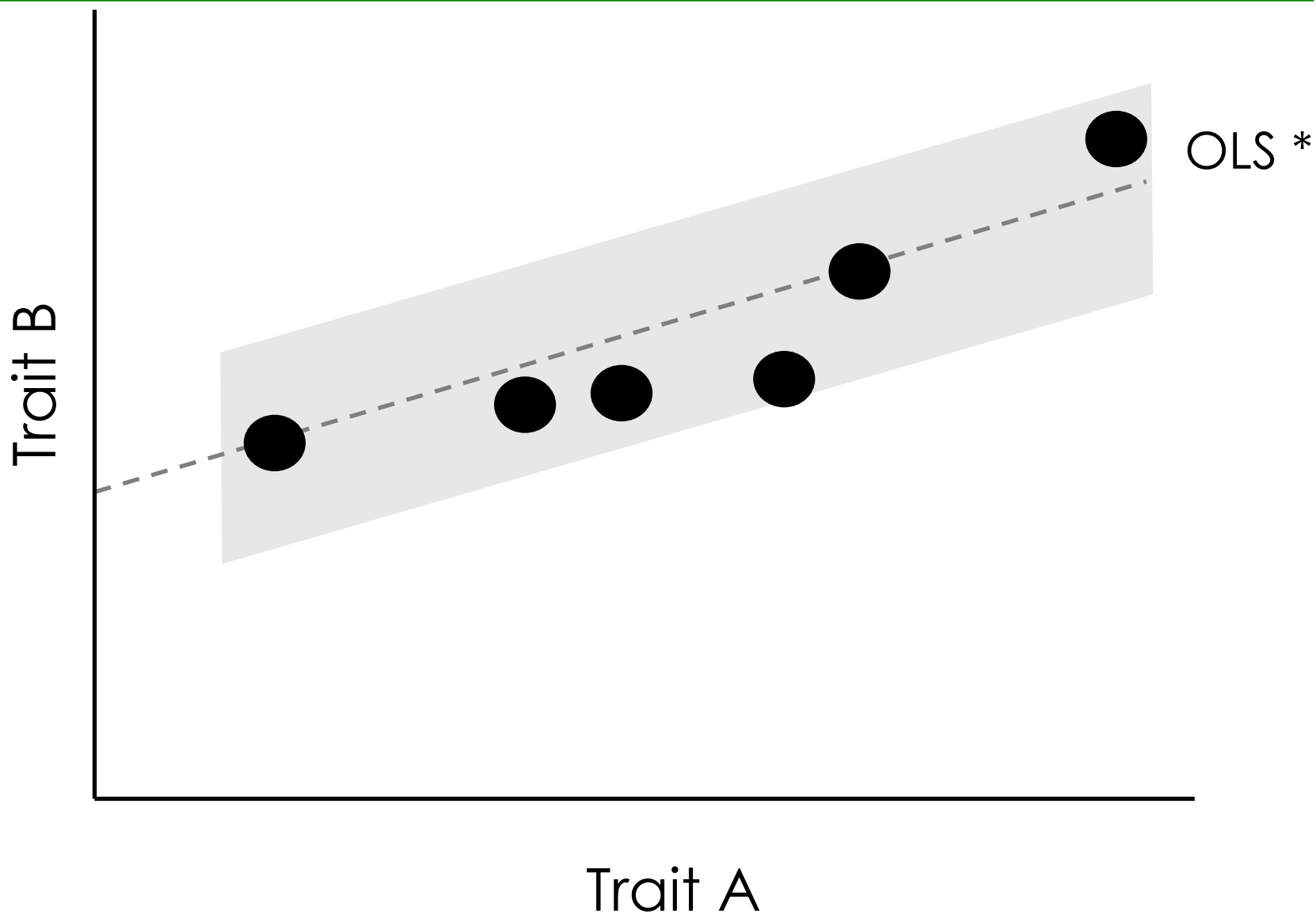
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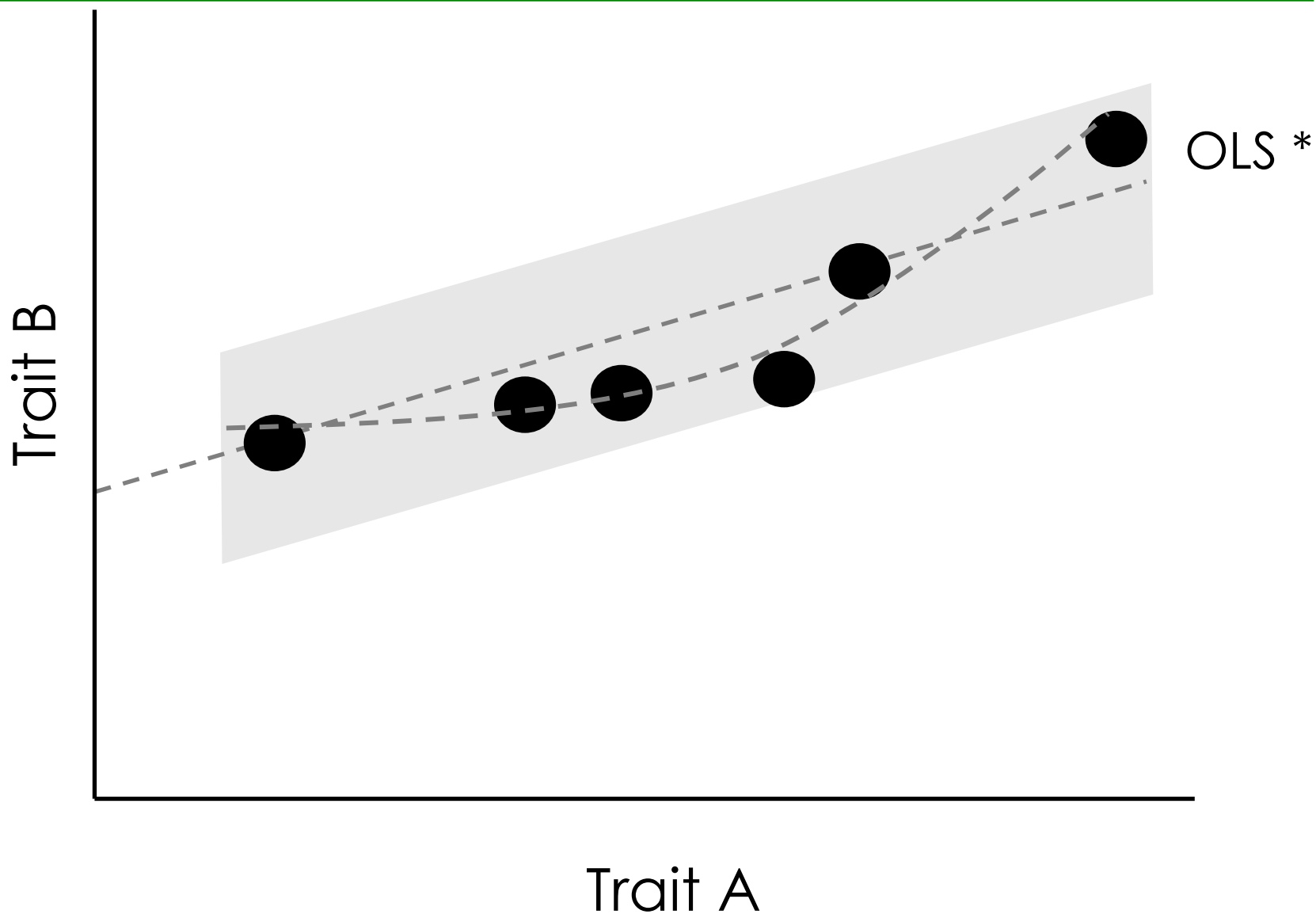
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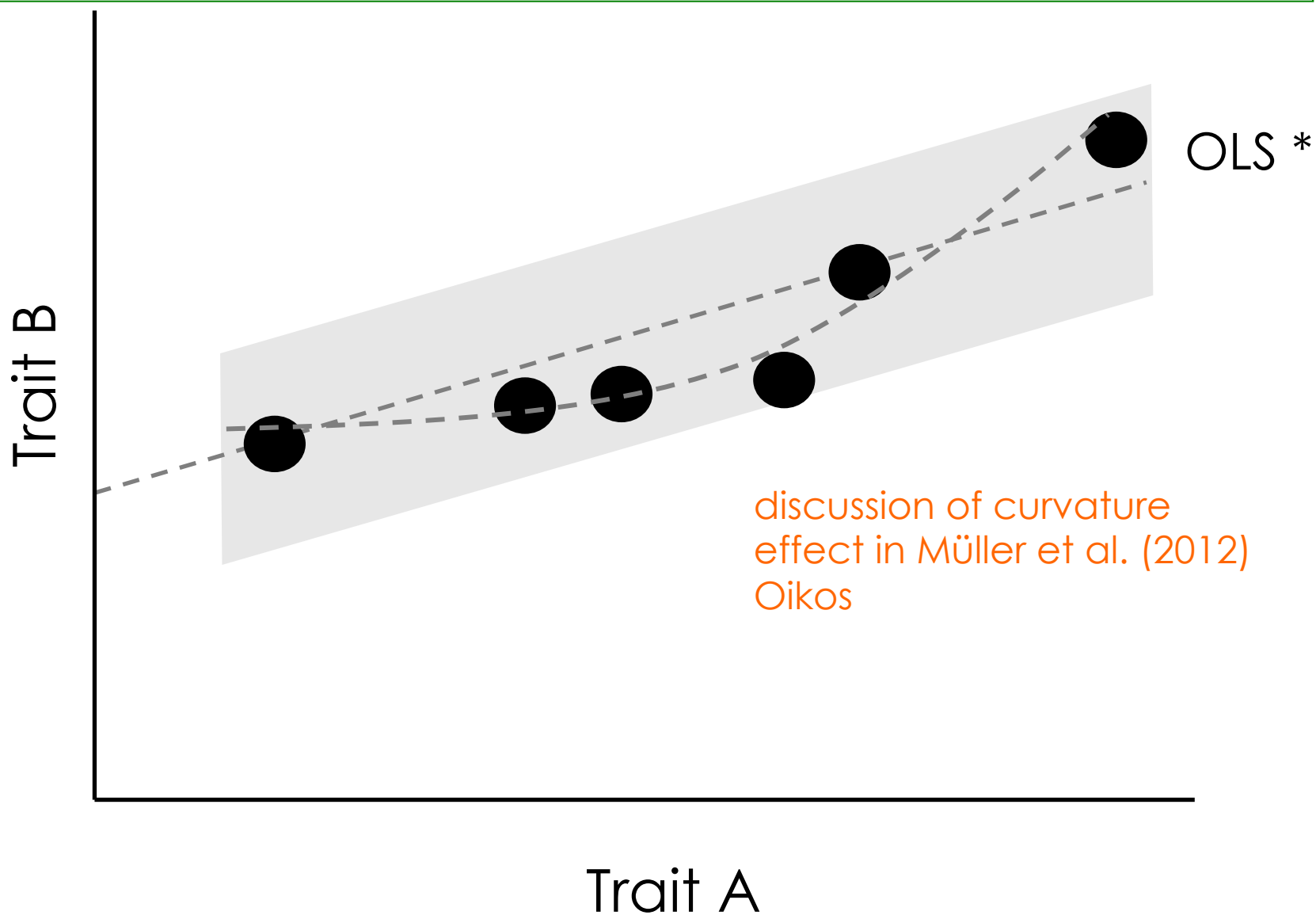
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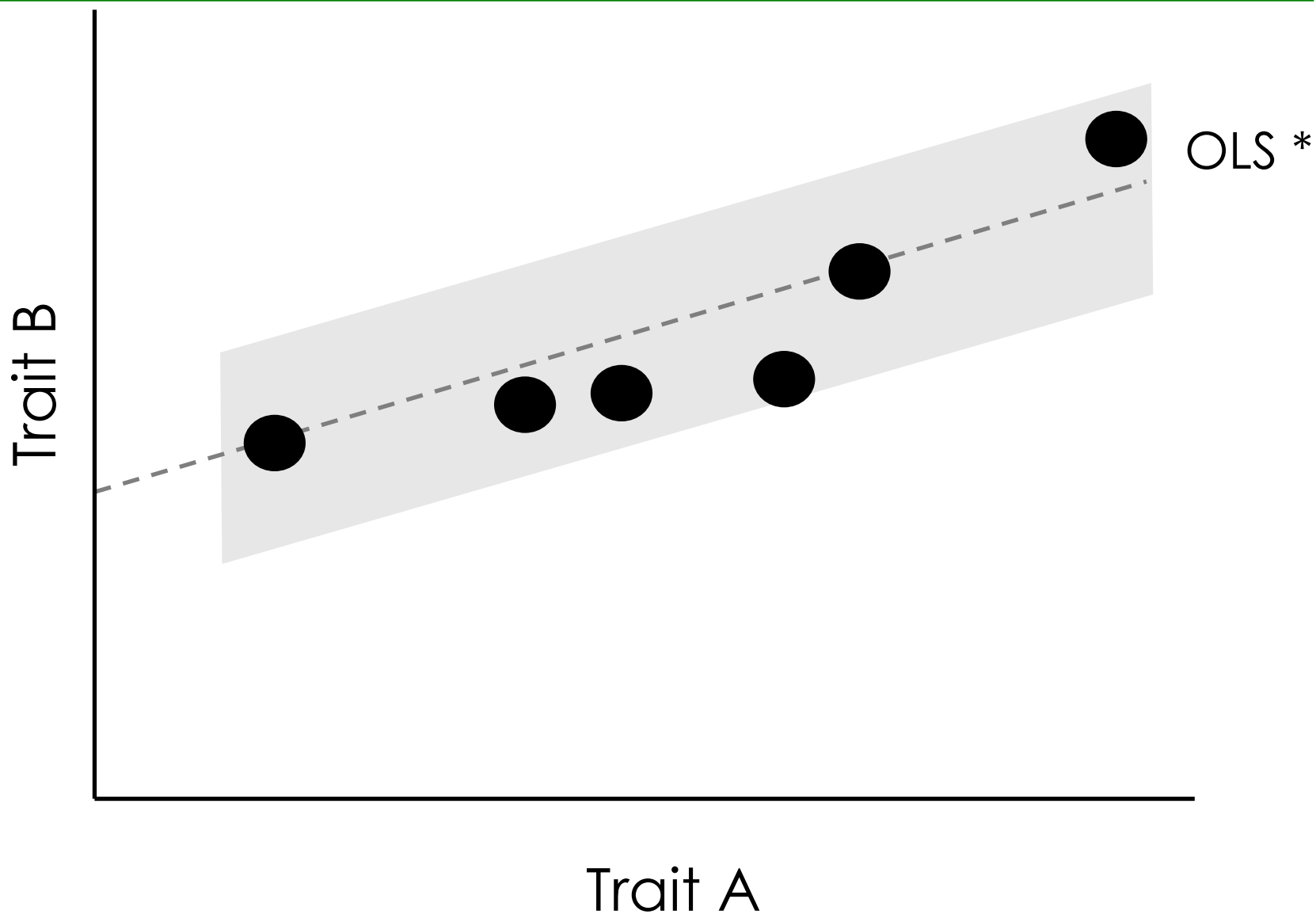
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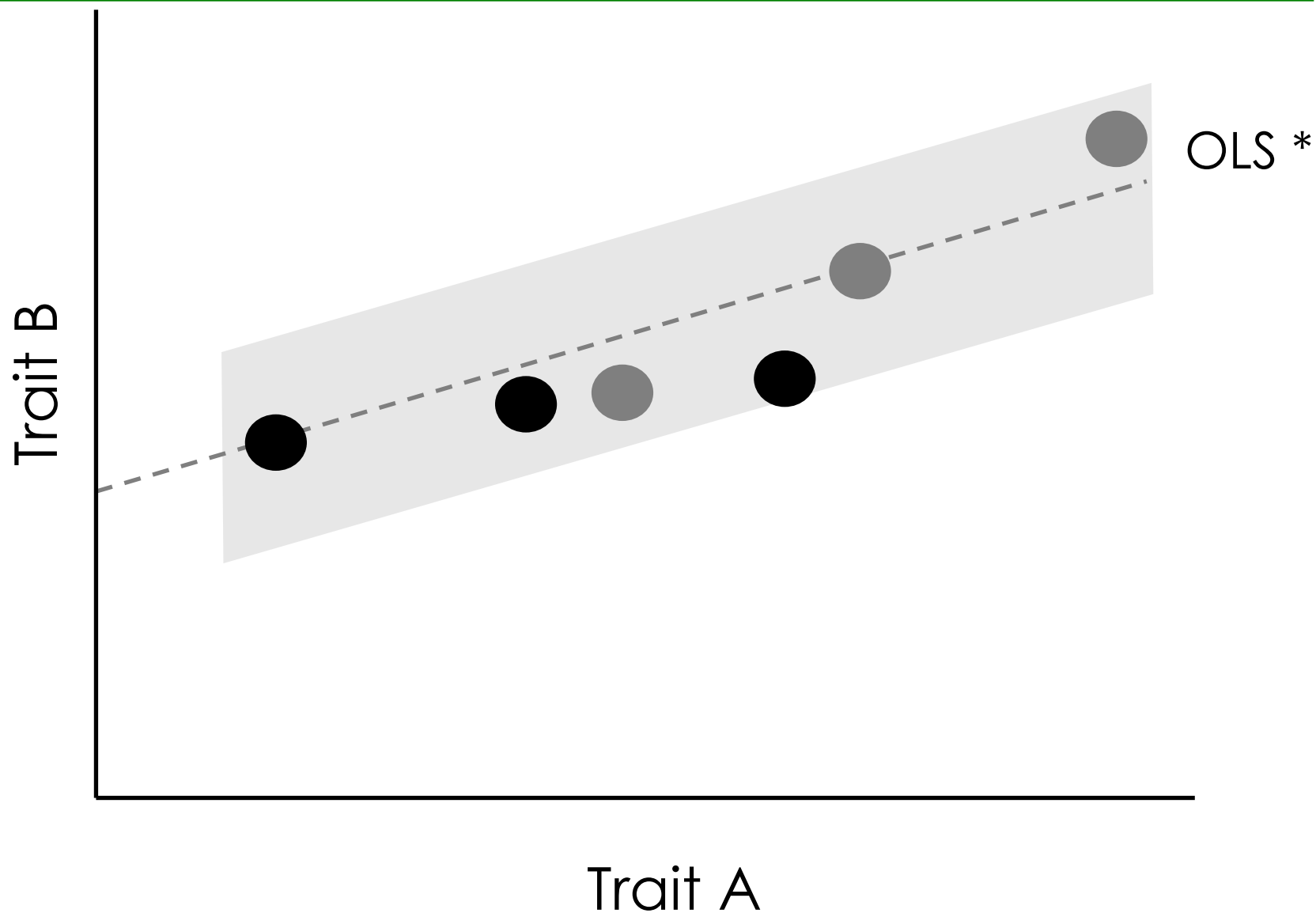
Accounting for phylogeny



from Clauss et al. (2013)



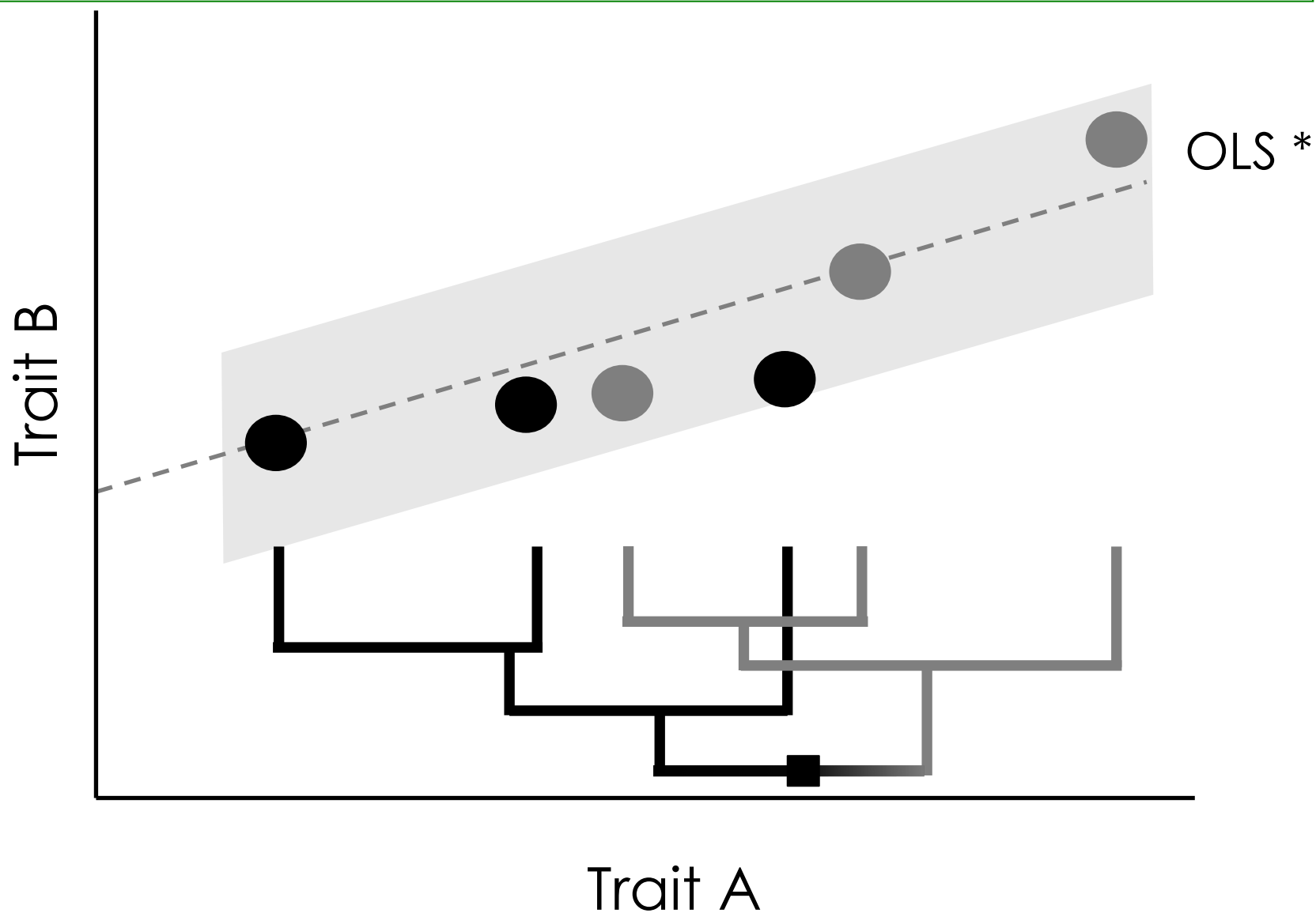
Accounting for phylogeny



from Clauss et al. (2013)



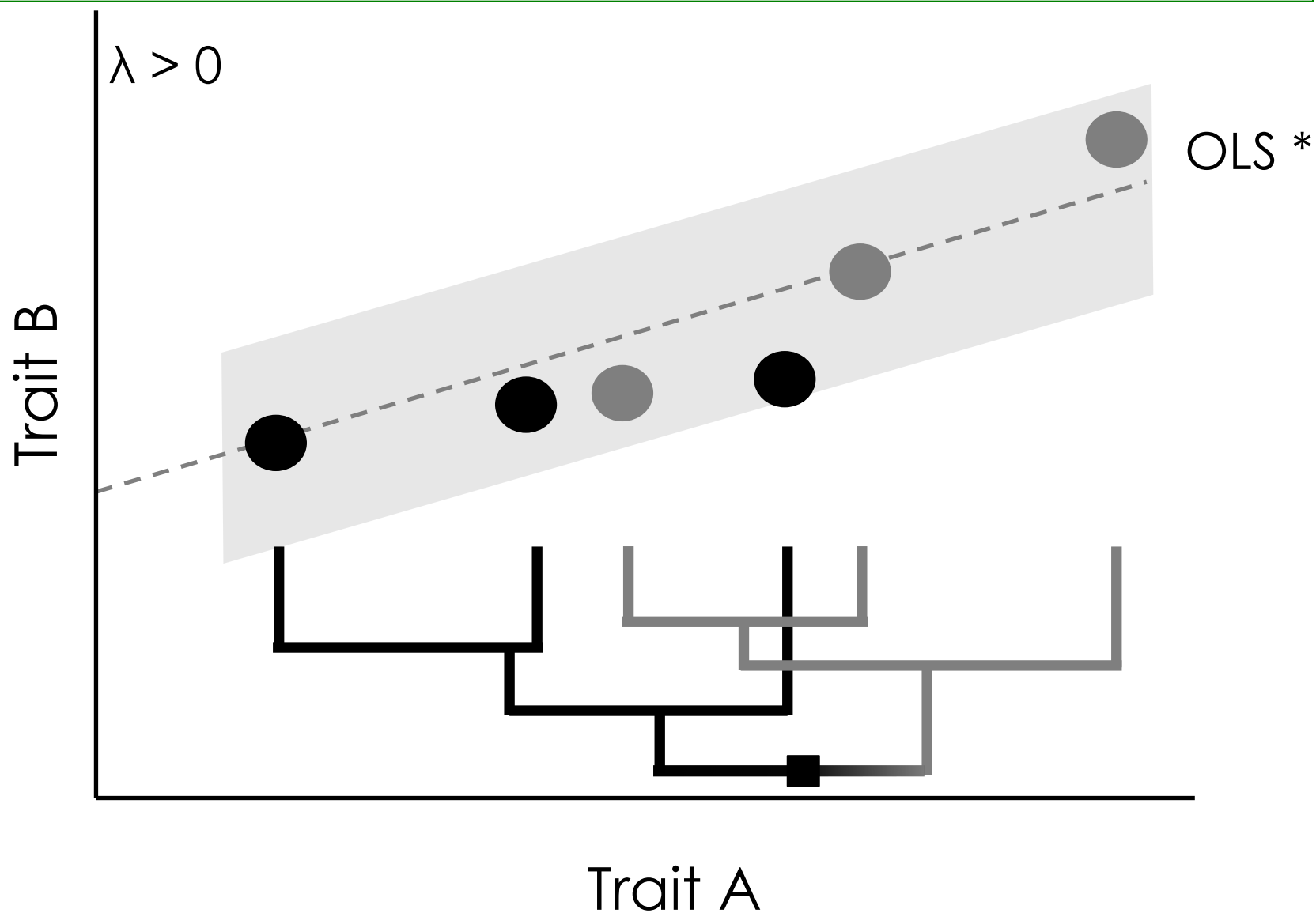
Accounting for phylogeny



from Clauss et al. (2013)



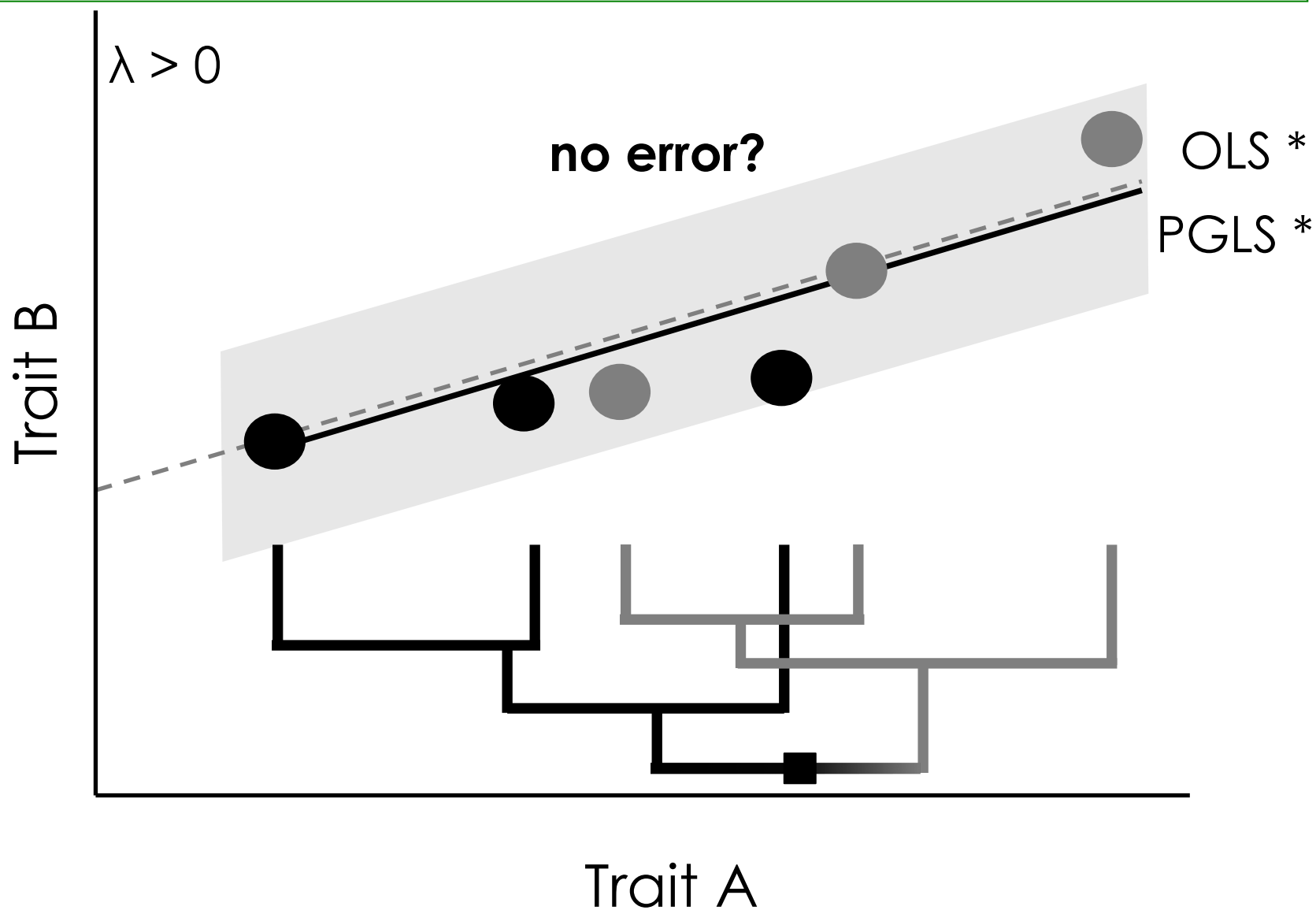
Accounting for phylogeny



from Clauss et al. (2013)

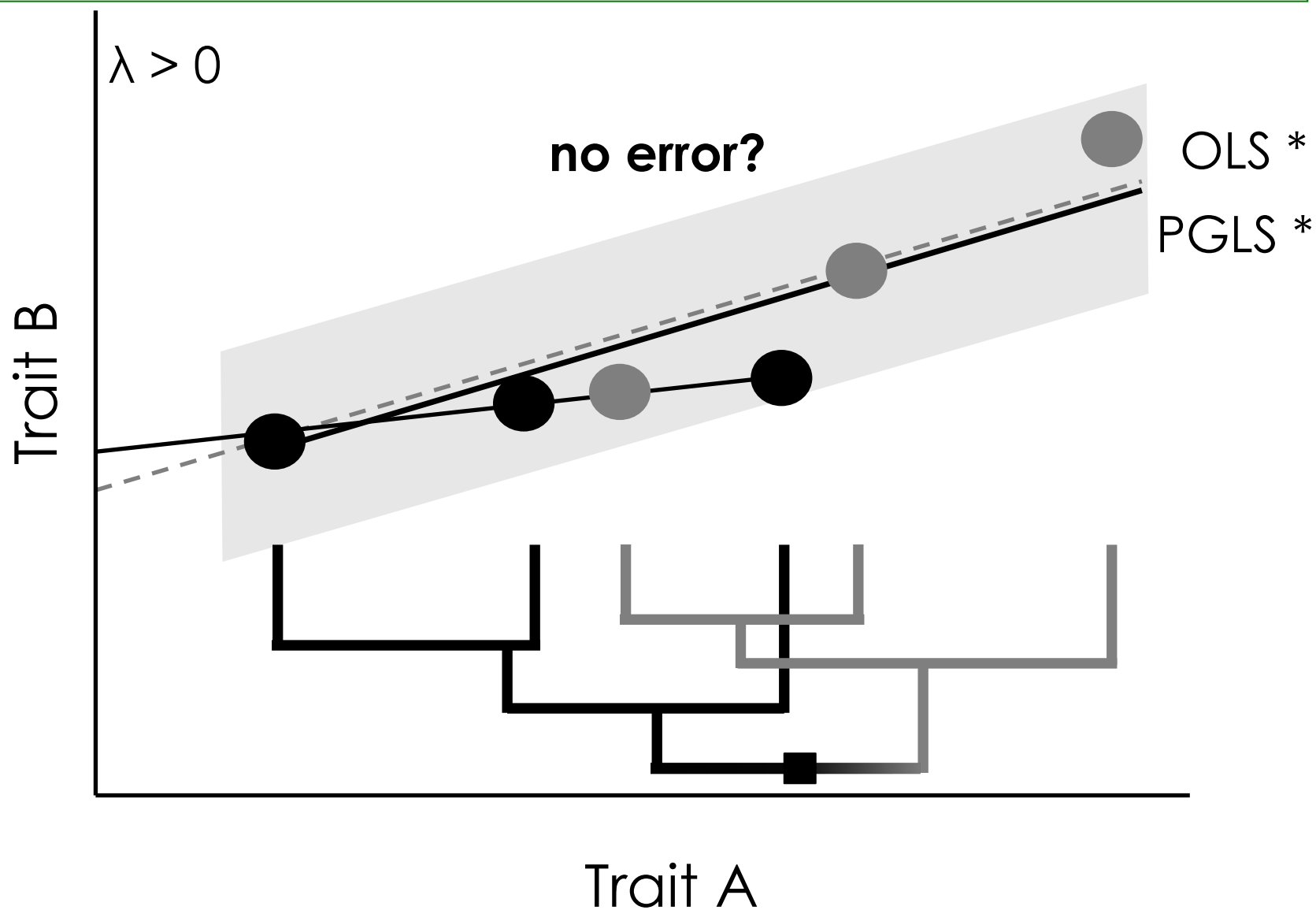


Accounting for phylogeny



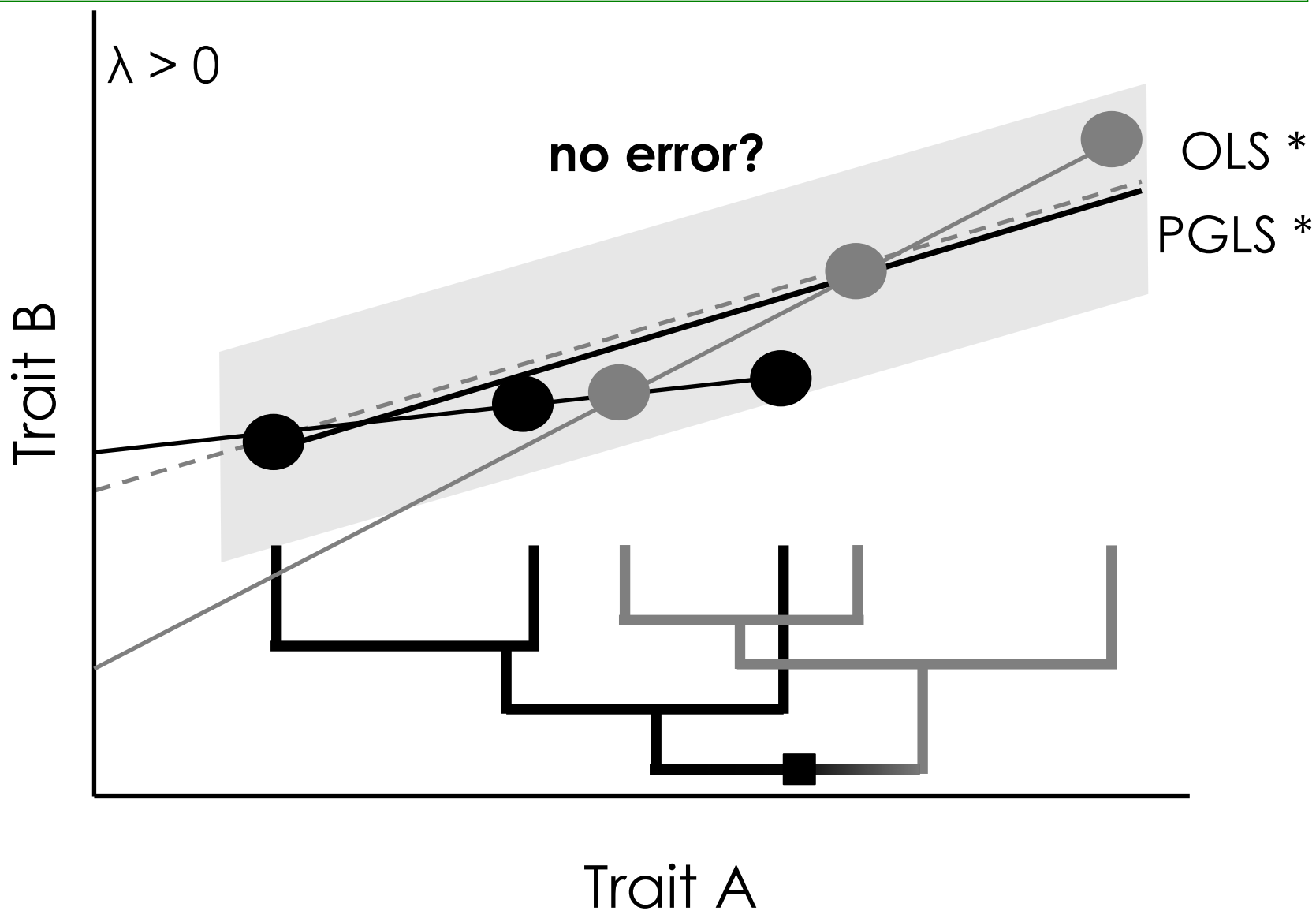


Accounting for phylogeny



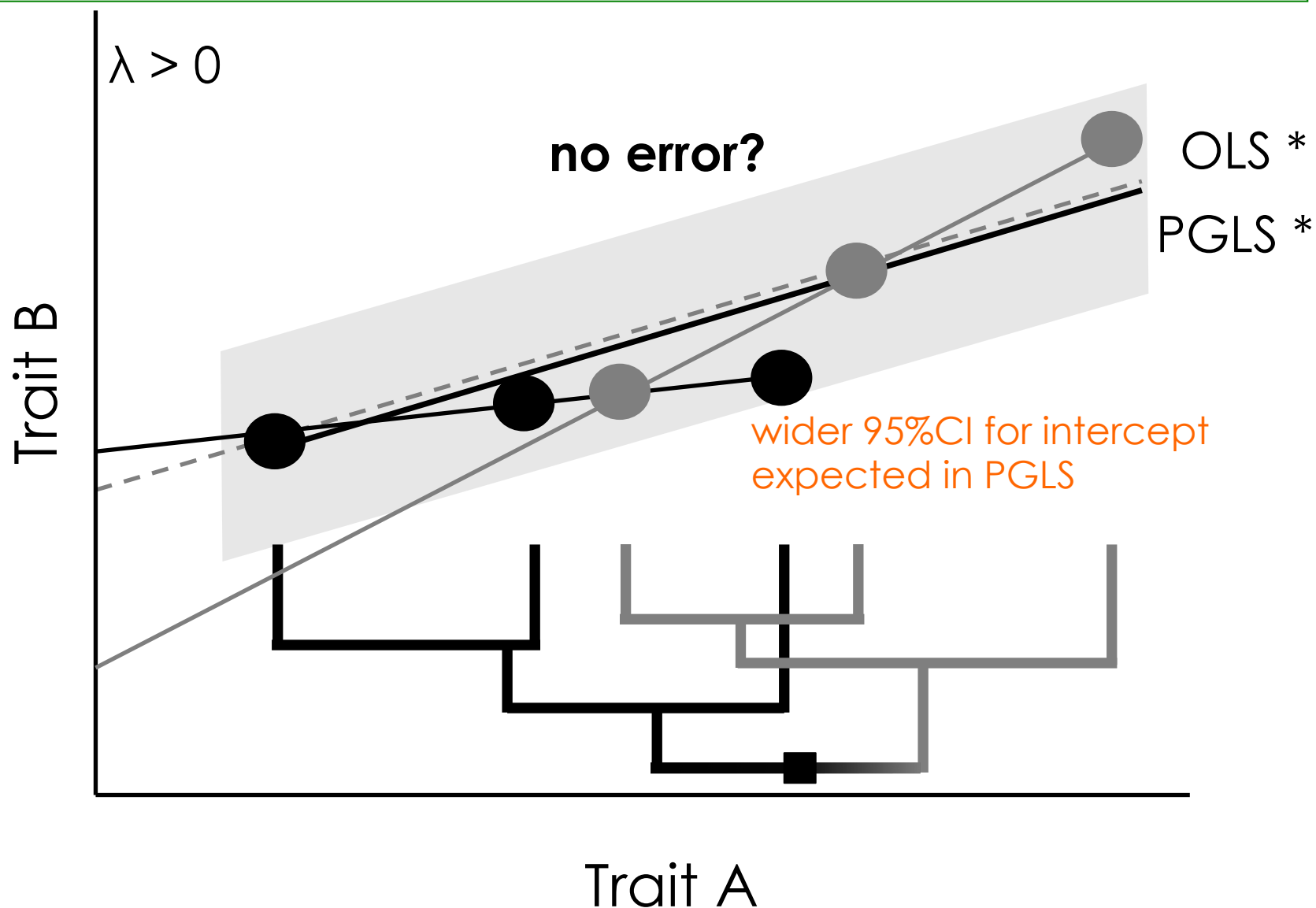


Accounting for phylogeny



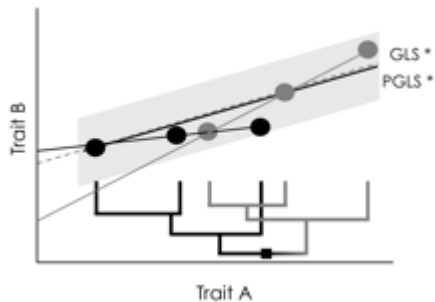


Accounting for phylogeny

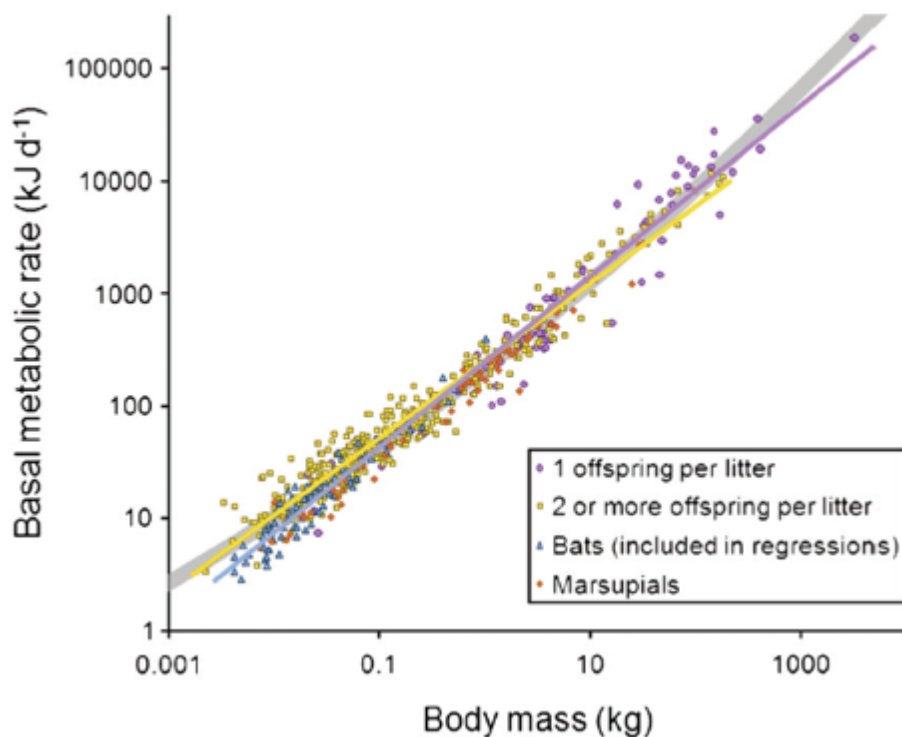




Example II: basal metabolic rate



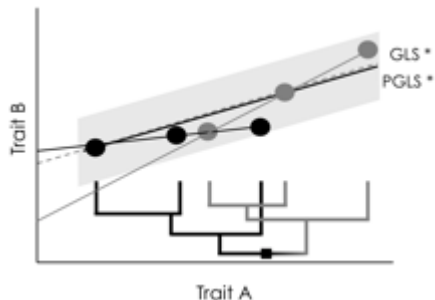
OLS: 2.38 (2.37-2.40) $BM^{0.72}$ (0.71-0.73)
PGLS: 2.25 (2.05-2.44) $BM^{0.73}$ (0.71-0.75)



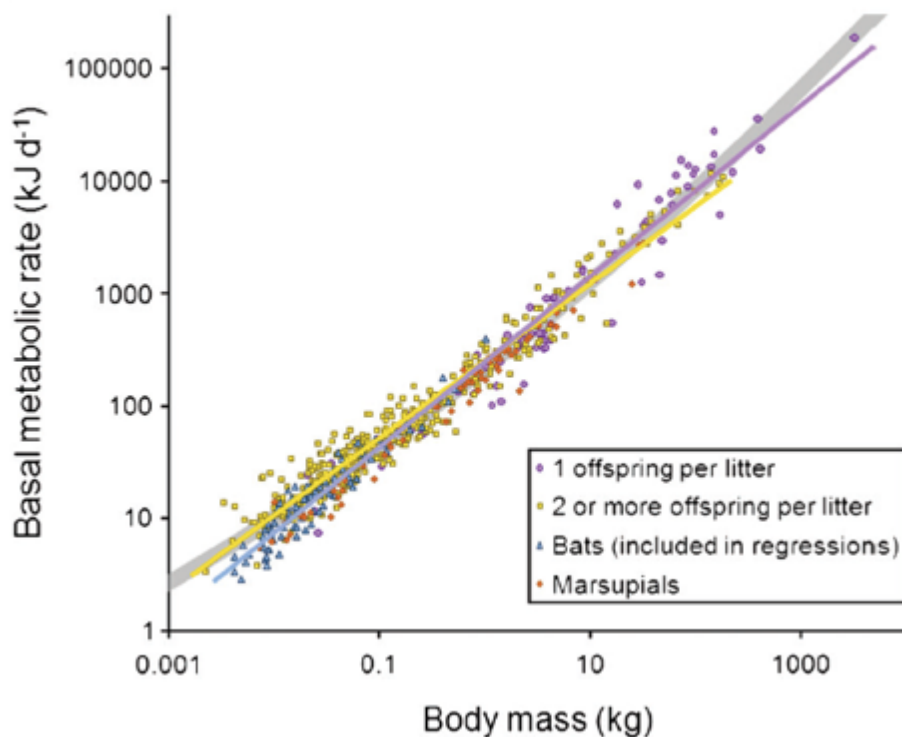
from Müller et al. (2012)



Example II: basal metabolic rate



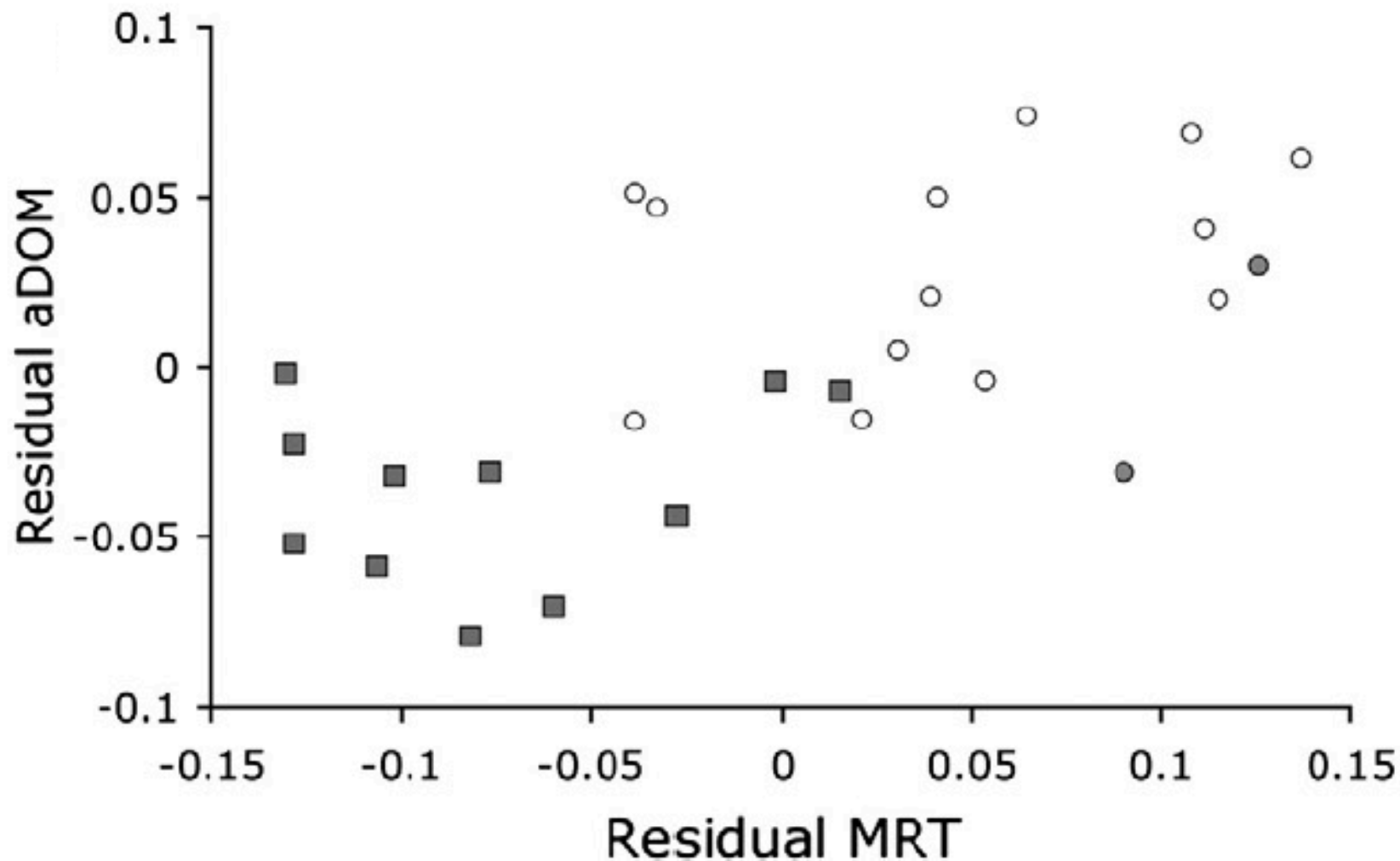
OLS: 2.38 (2.37-2.40) $BM^{0.72}$ (0.71-0.73)
PGLS: 2.25 (2.05-2.44) $BM^{0.73}$ (0.71-0.75)



from Müller et al. (2012)



Example III: retention/digestibility



from Müller et al. (2013)



Accounting for phylogeny

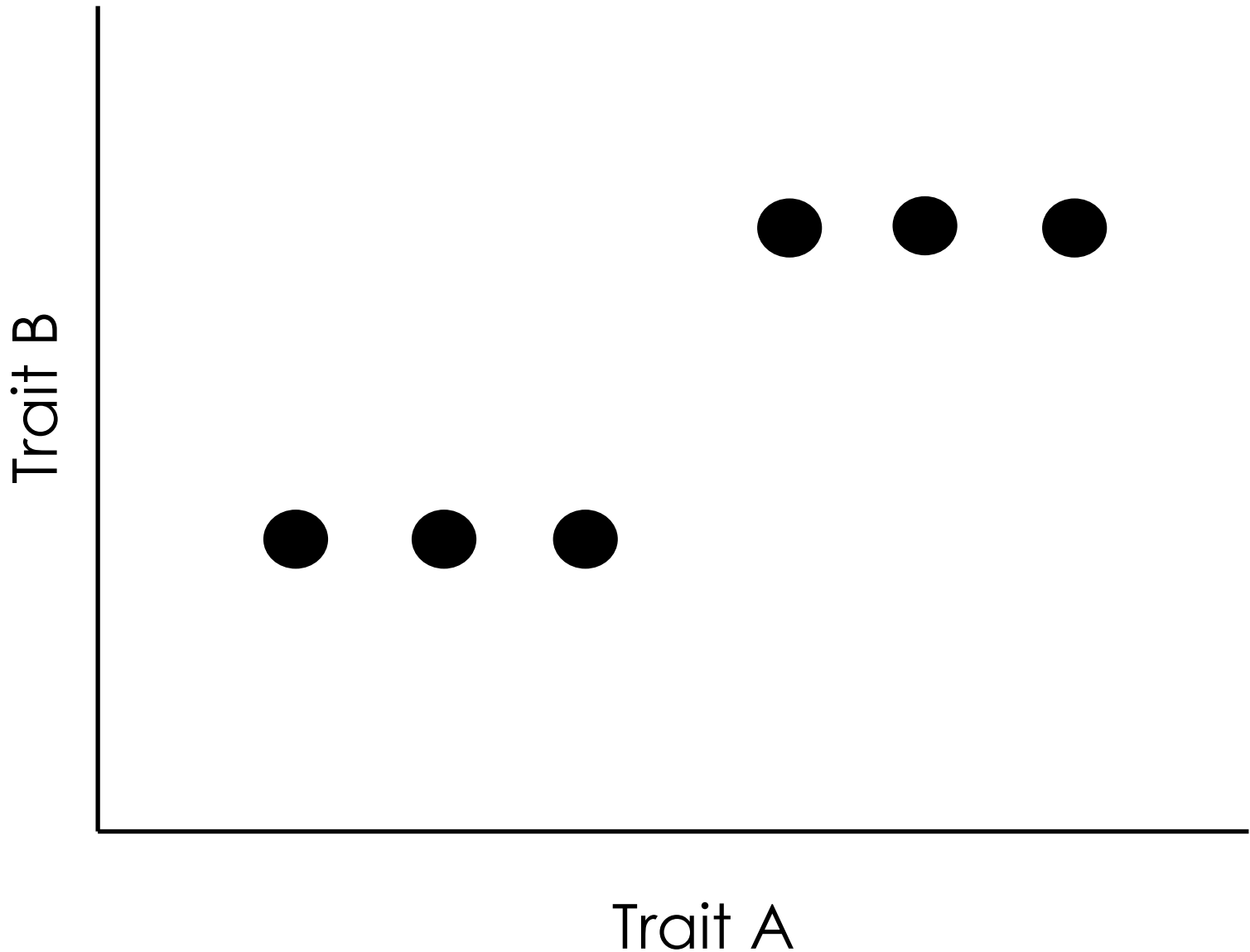
Trait B

Trait A

from Clauss et al. (2013)



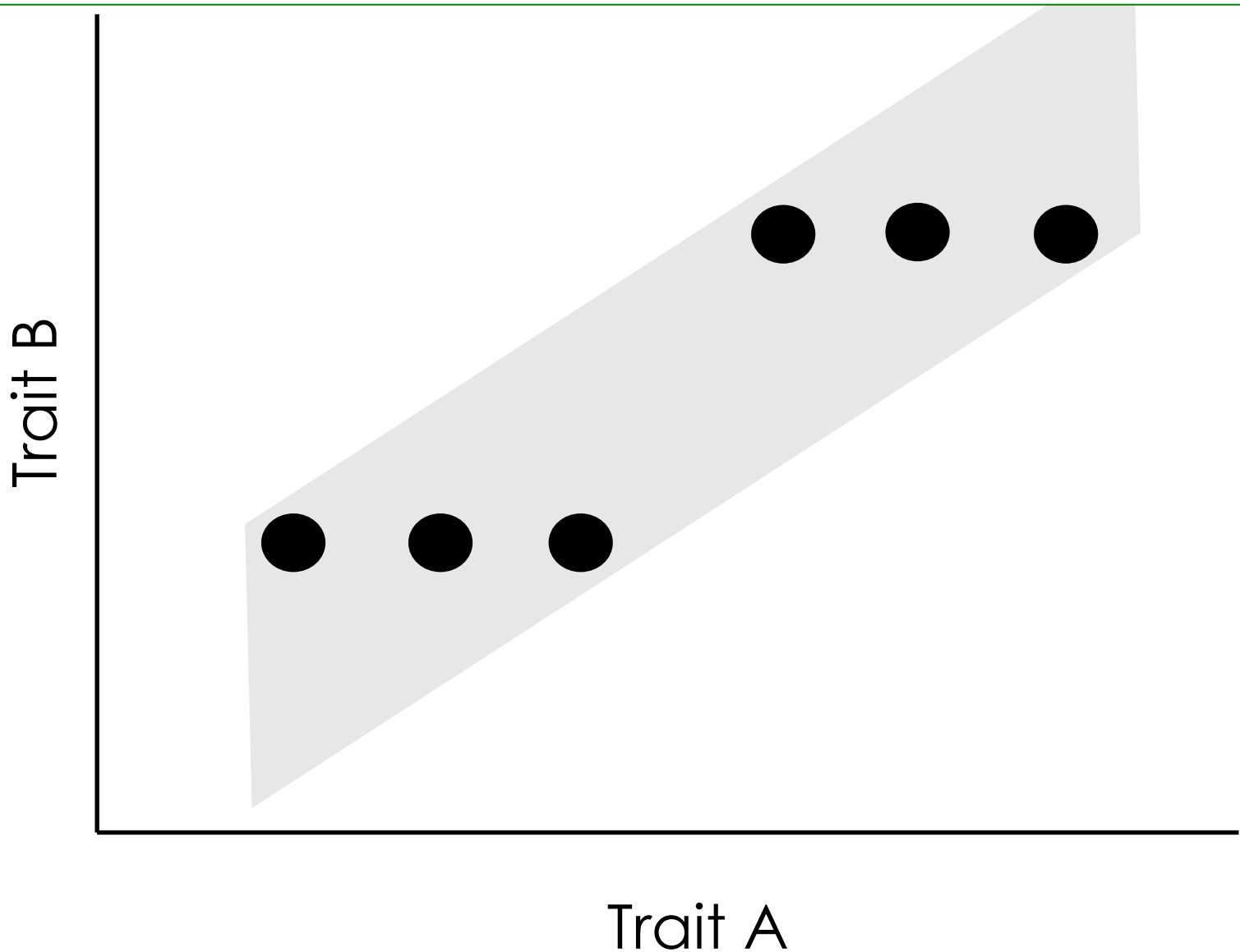
Accounting for phylogeny



from Clauss et al. (2013)



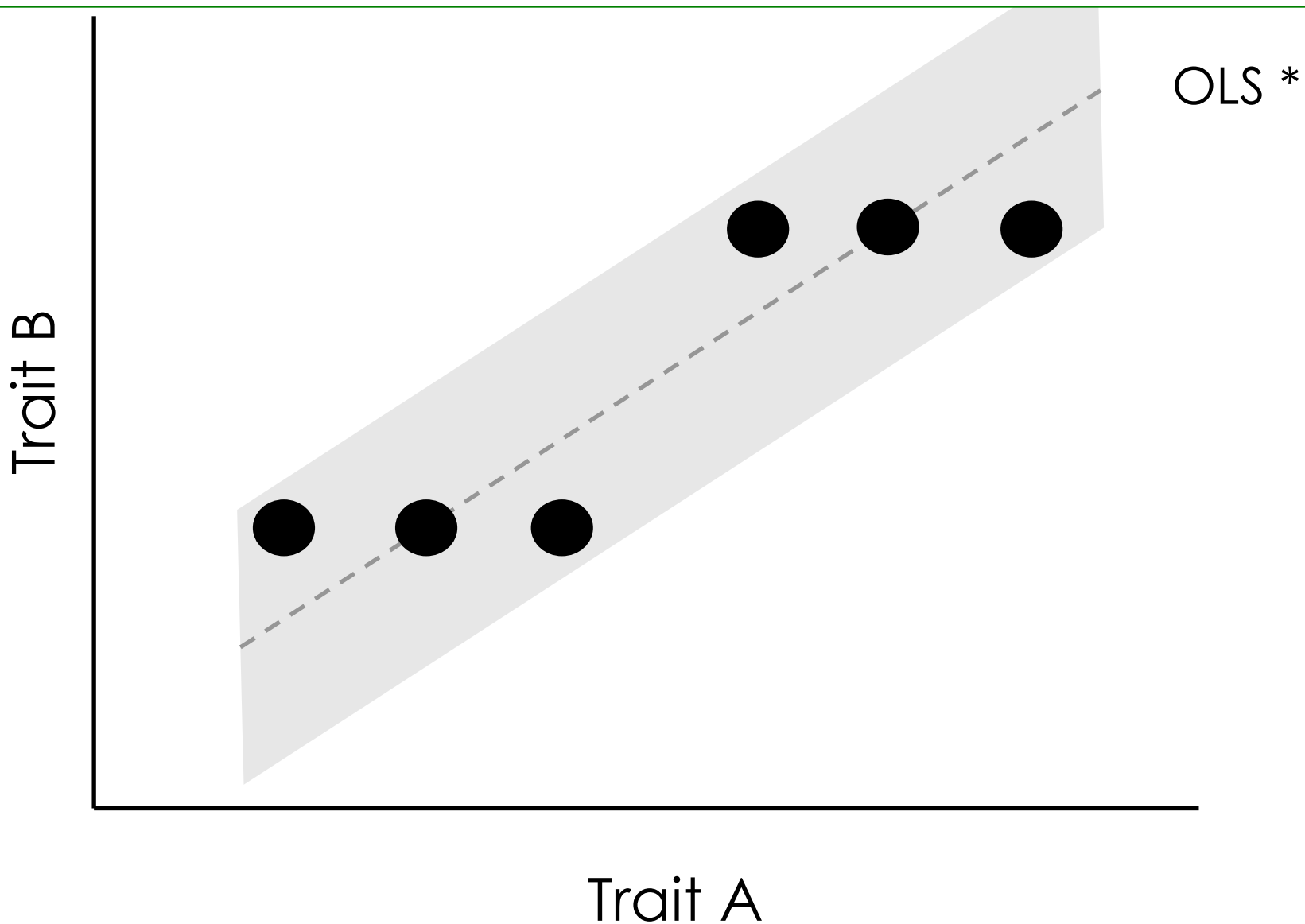
Accounting for phylogeny



from Clauss et al. (2013)



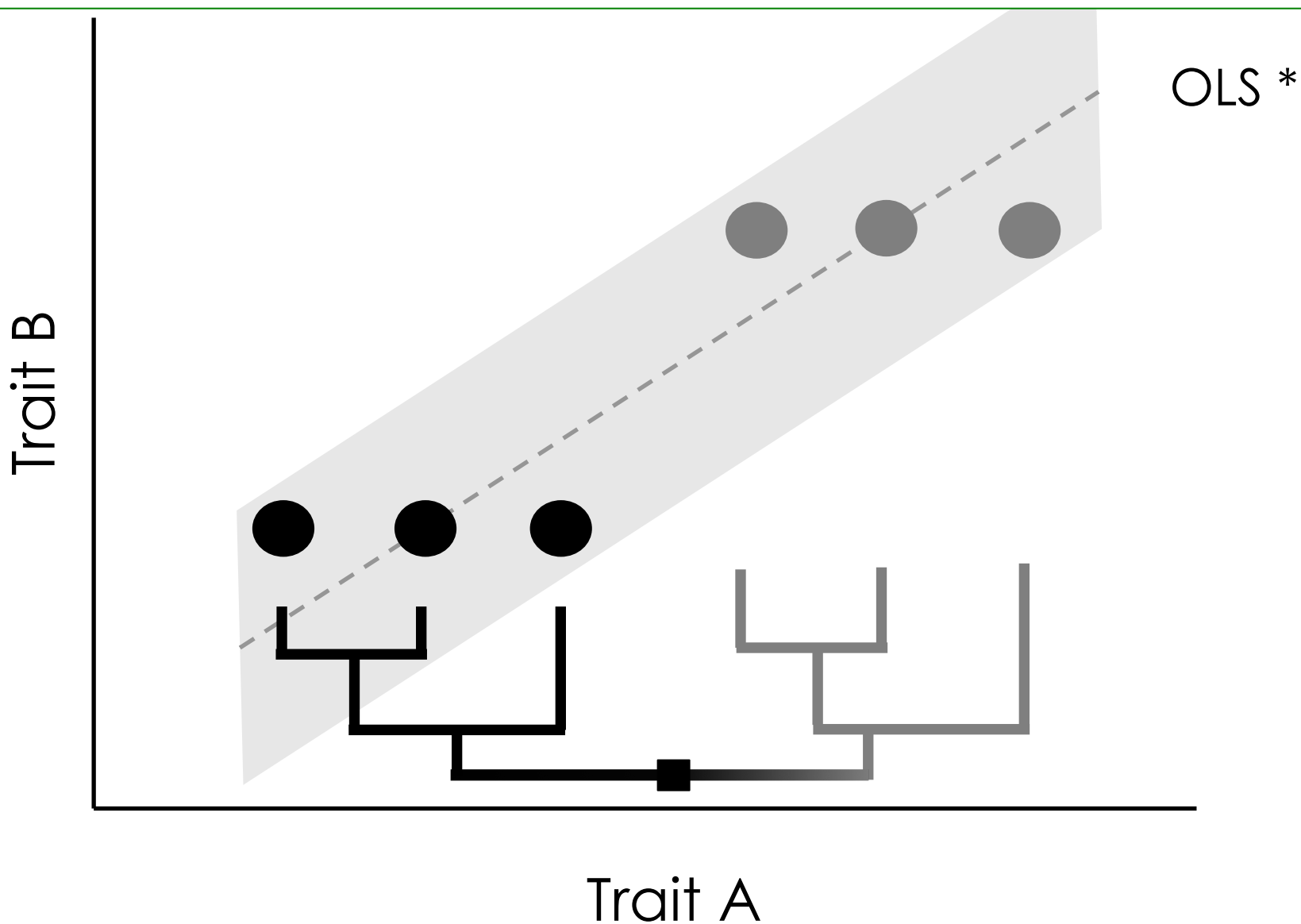
Accounting for phylogeny



from Clauss et al. (2013)



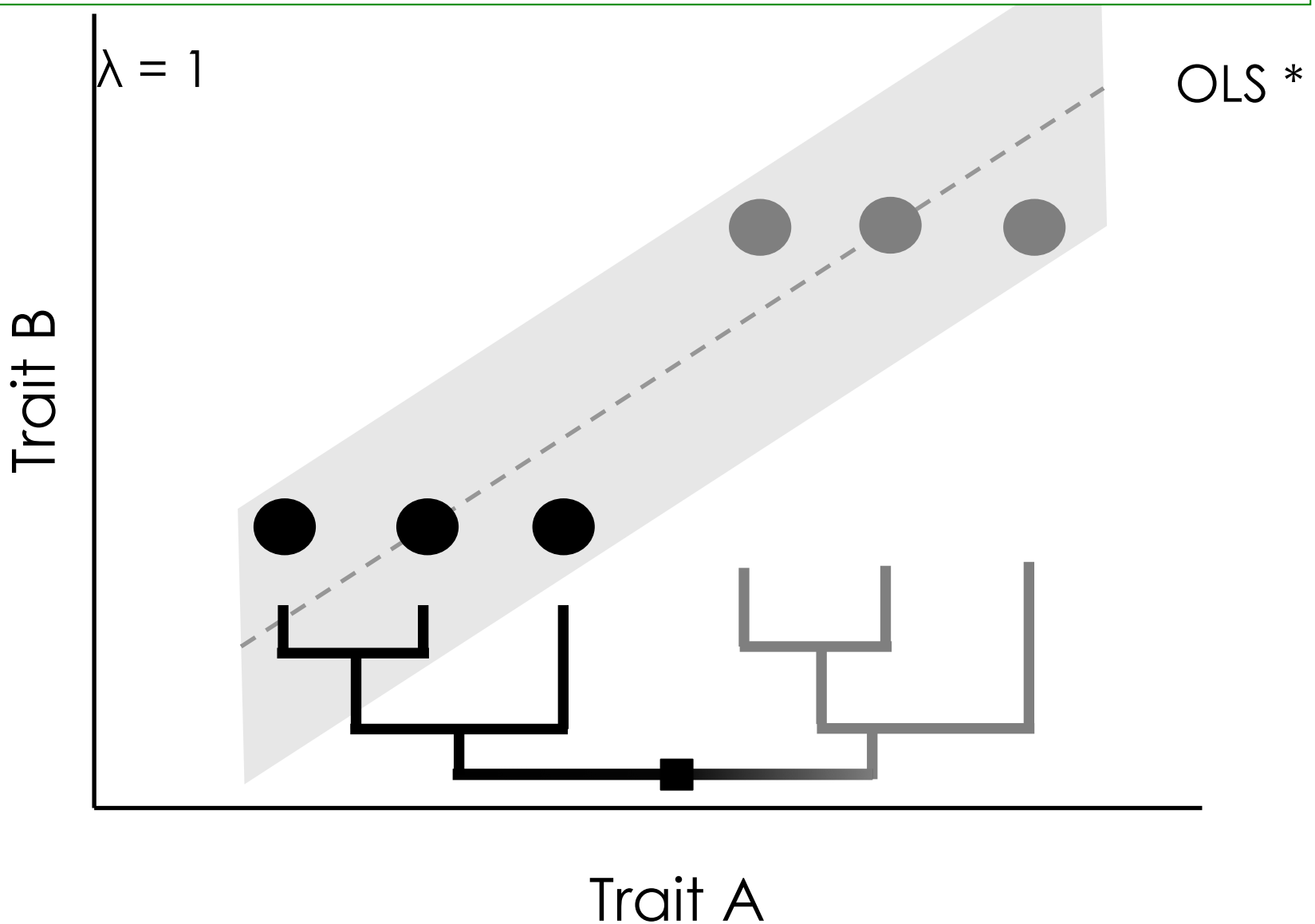
Accounting for phylogeny



from Clauss et al. (2013)

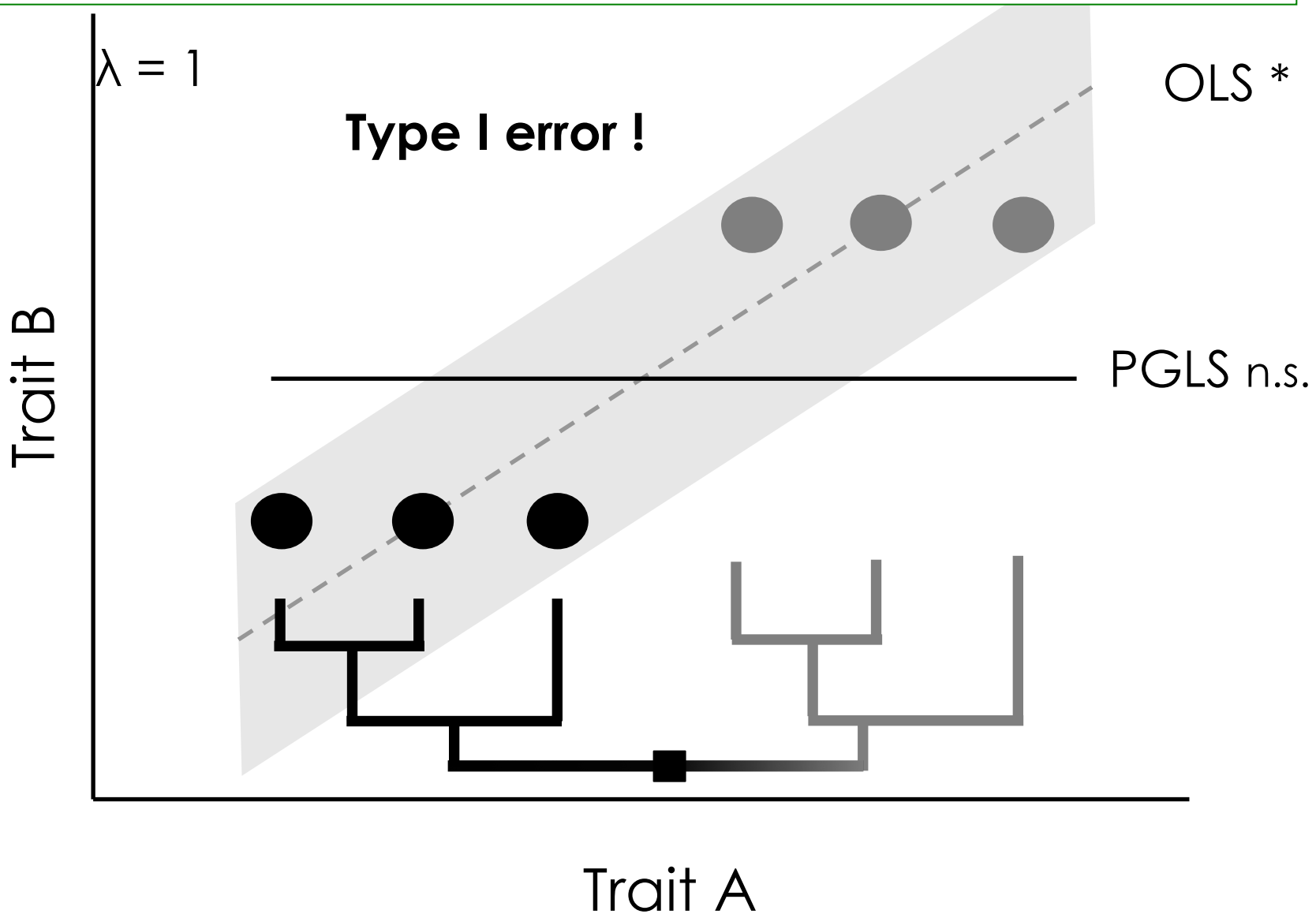


Accounting for phylogeny





Accounting for phylogeny

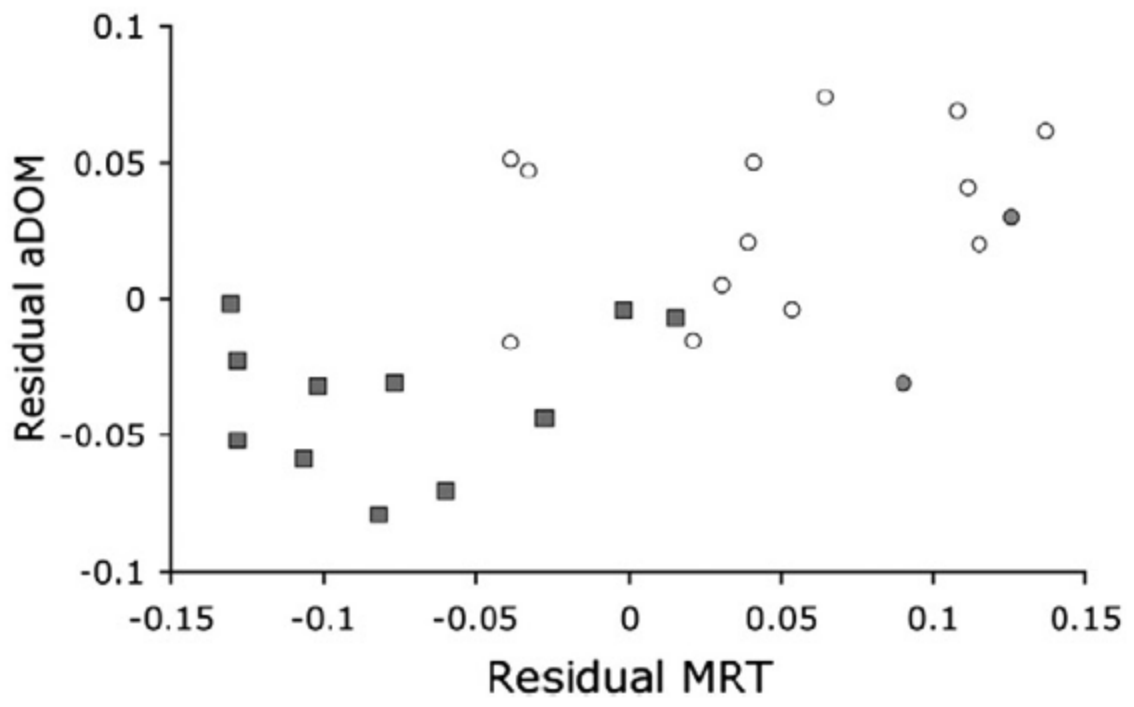




Example IIIa: retention/digestibility

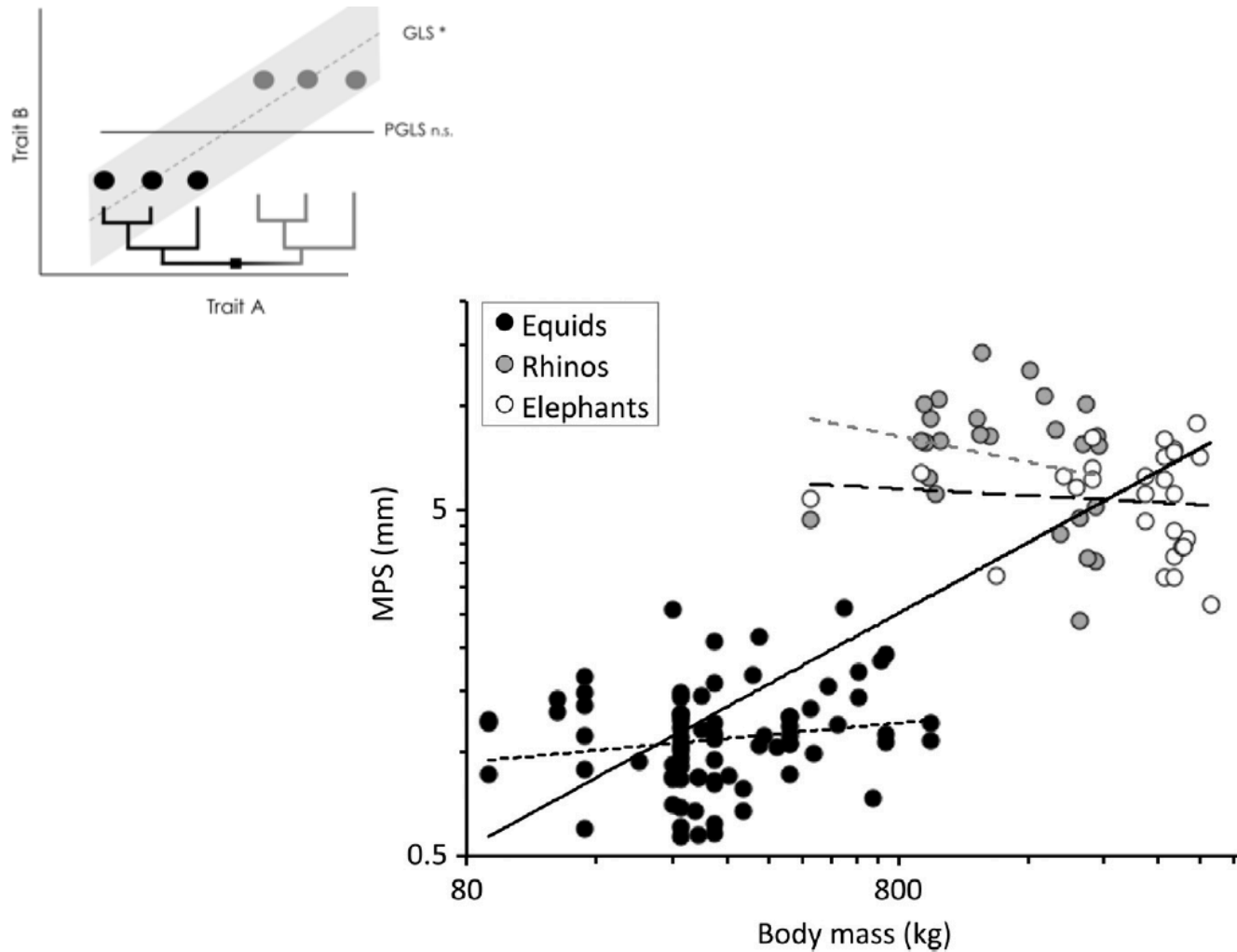


OLS: significant
PGLS: not significant



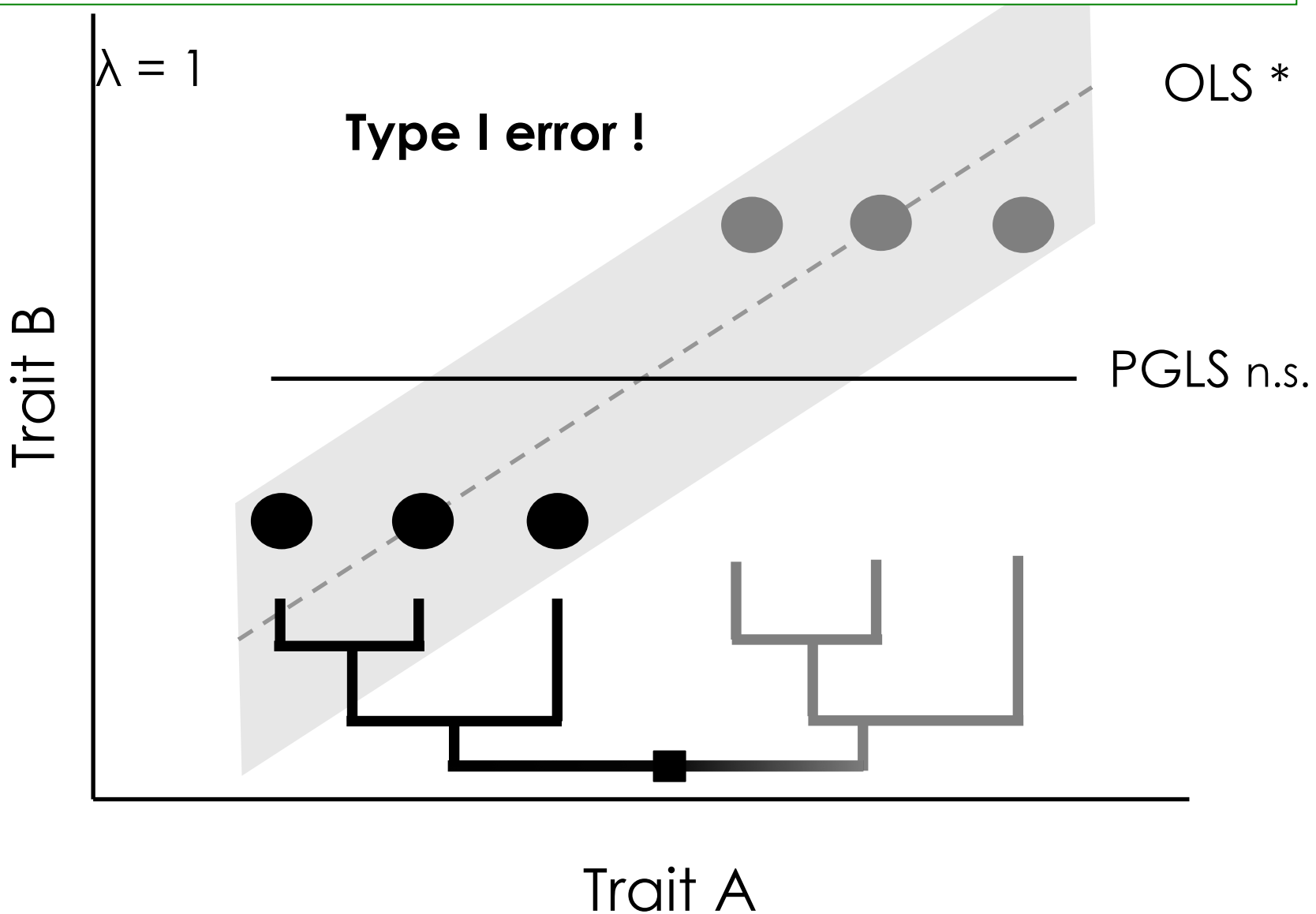


Example IIIb: fecal particle size





Accounting for phylogeny



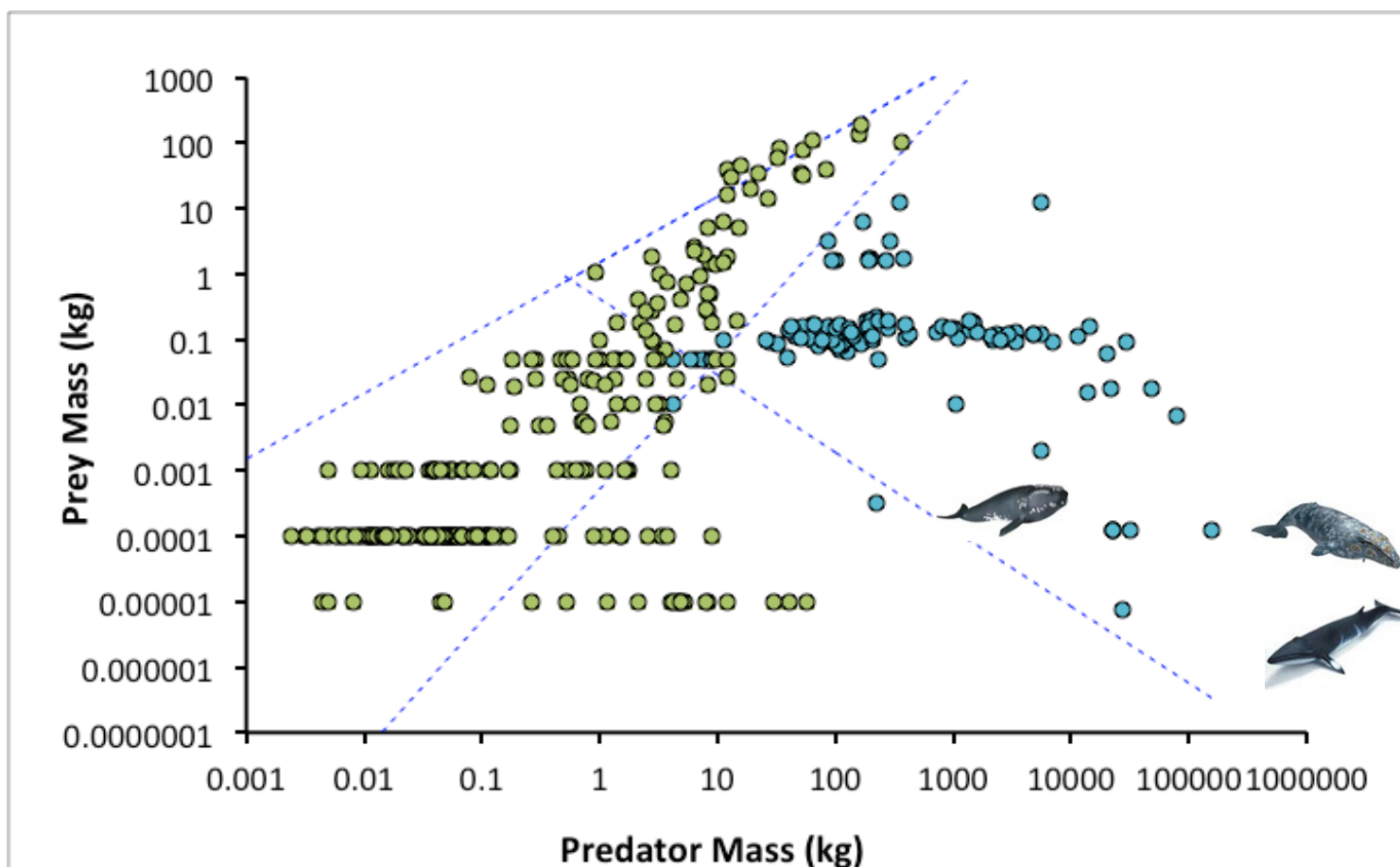


Geometric factors influencing the diet of vertebrate predators in marine and terrestrial environments

Chris Carbone,^{1*} Daryl Codron,^{2,3}

Conrad Scofield,¹ Marcus Clauss³

and Jon Bielby¹



from Carbone et al. (2014)

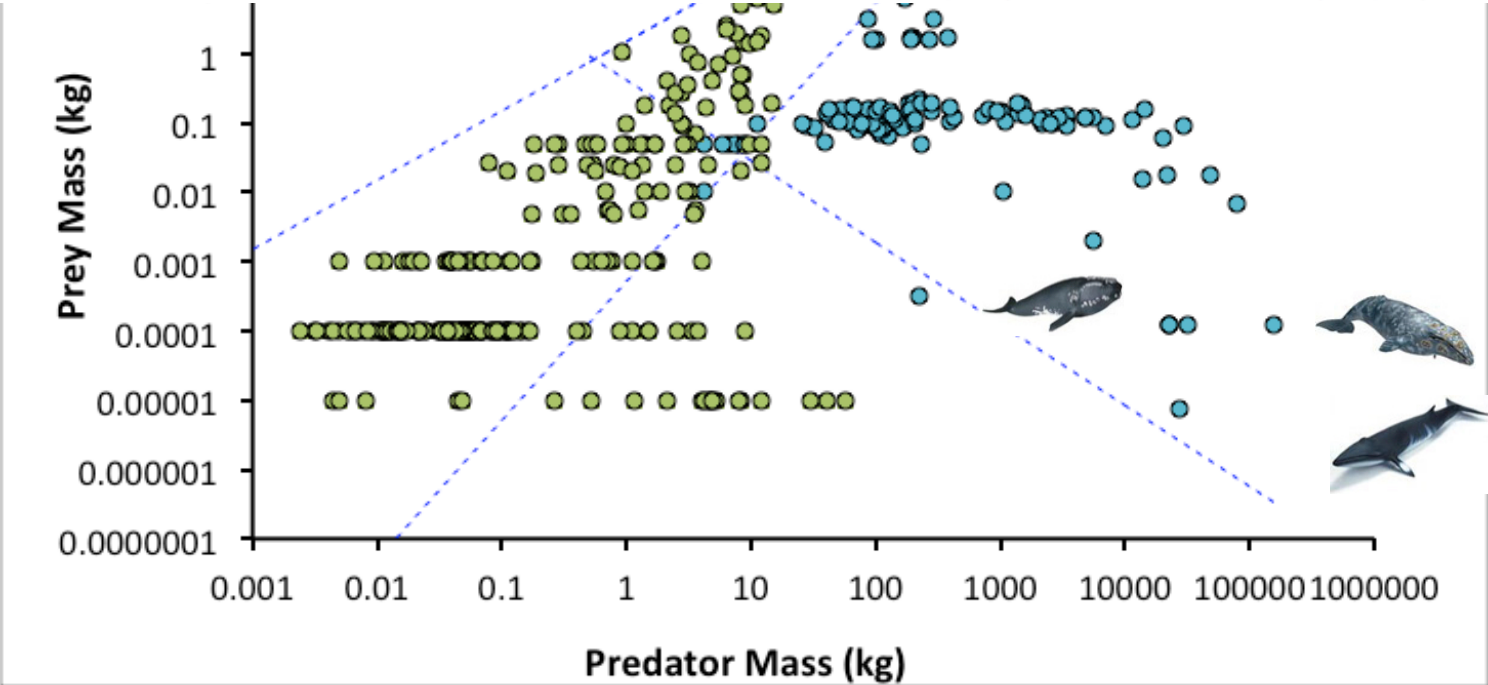


Geometric factors influencing the diet of vertebrate predators in marine and terrestrial environments

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Conrad Scofield,¹ Marcus Clauss³
and Jon Bielby¹

Table 2 Result of comparative analyses of how minimum prey size varies as a function of predator size

Taxonomic group/biome	n	Body mass (kg, mode, range)		Stat	λ^*	a (95% CI)	t	P	b (95% CI)	t	P
		Predator	Prey								
Terrestrial mammals	270	0.112 (0.002–371)	0.0001 (0.000001–189)	OLS	(0)	0.007 (0.004; 0.010)	−22.456	0.000	1.05 (0.90; 1.20)	13.709	0.000
				PGLS [§]	0.929 [‡]	0.0003 (0.00001; 0.013)	−4.276	0.000	0.82 (0.60; 1.03)	7.381	0.000
				PGLS [†]	1.0 [†]	0.0001 (0.00001; 0.001)	−7.923	0.000	0.36 (0.15; 0.57)	3.293	0.001
Marine Mammals	126	23000 (4–154160)	0.100 (0.00003–12)	OLS	(0)	0.546 (0.215; 1.386)	−1.274	0.205	−0.30 (−0.45; −0.15)	−3.975	0.000
				PGLS [†]	0.978 [†]	0.013 (0.001; 0.232)	−2.940	0.004	0.16 (−0.13; 0.44)	1.054	0.294

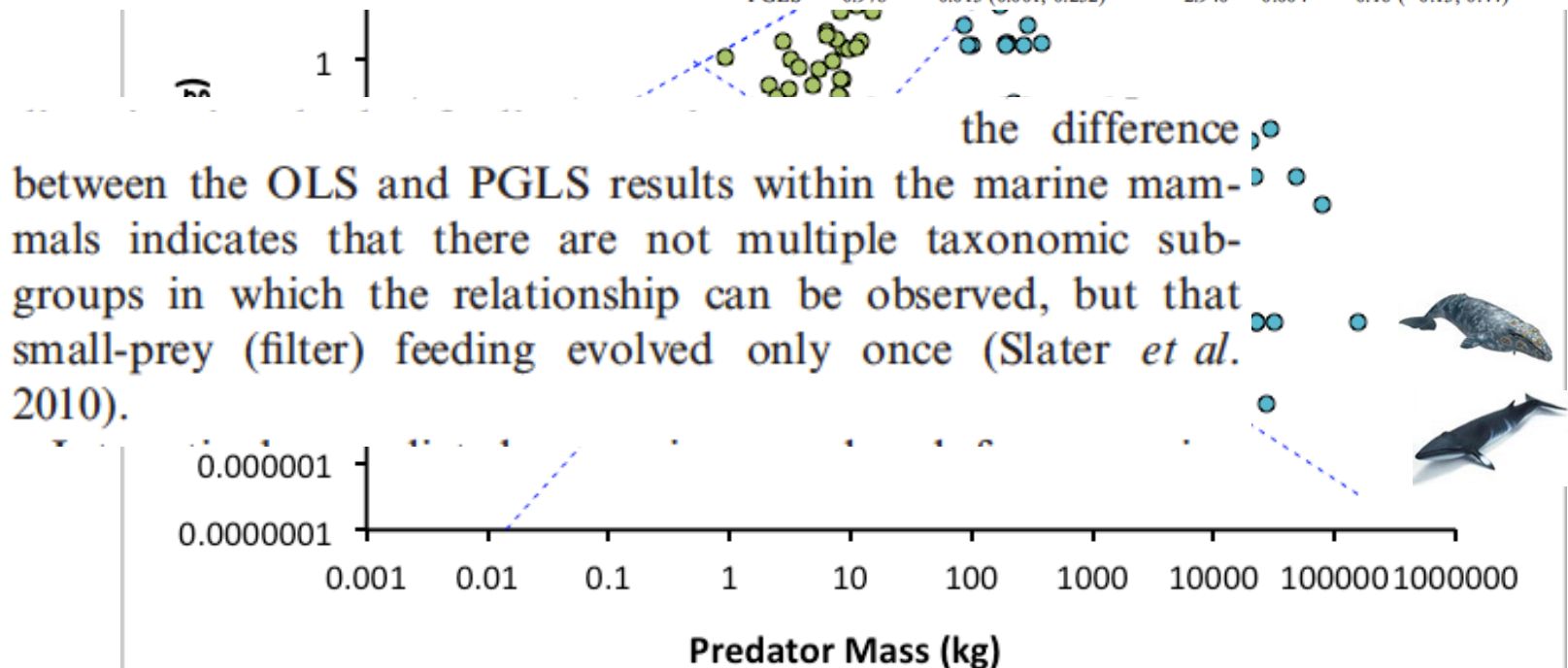


from Carbone et al. (2014)

Geometric factors influencing the diet of vertebrate predators in marine and terrestrial environments

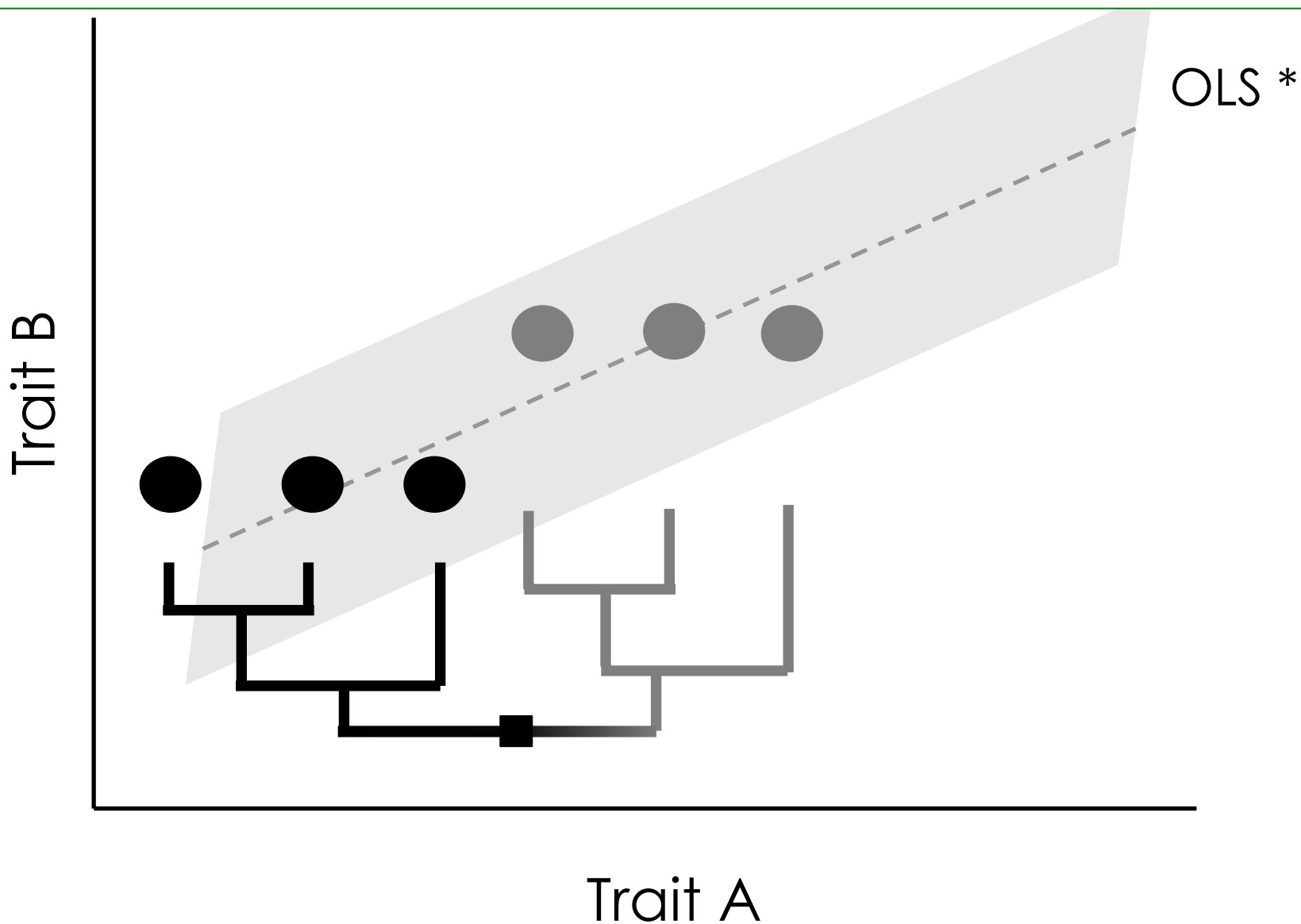
Chris Carbone,^{1*} Daryl Codron,^{2,3}Conrad Scofield,¹ Marcus Clauss³and Jon Bielby¹**Table 2** Result of comparative analyses of how minimum prey size varies as a function of predator size

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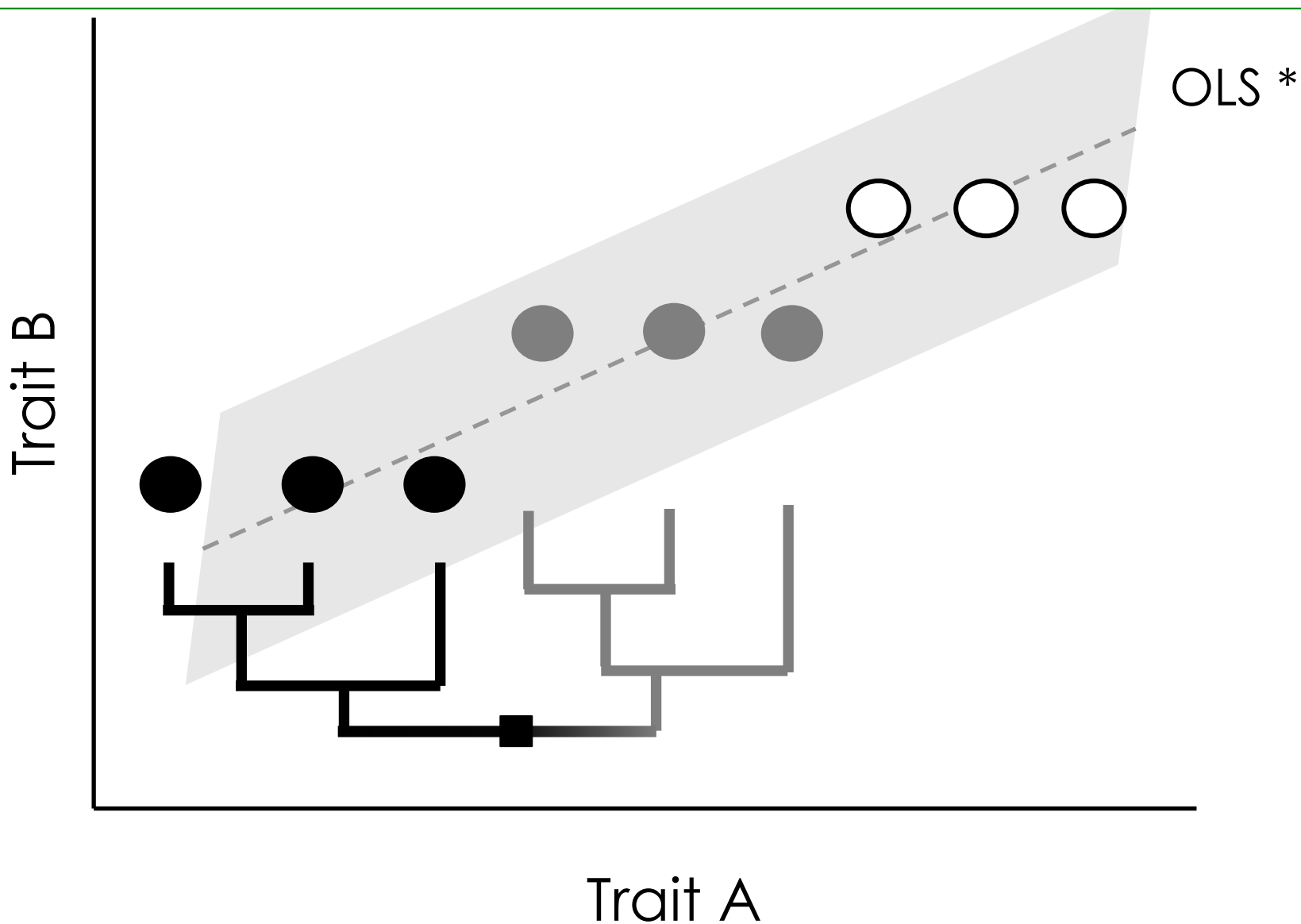
Accounting for phylogeny



from Clauss et al. (2013)



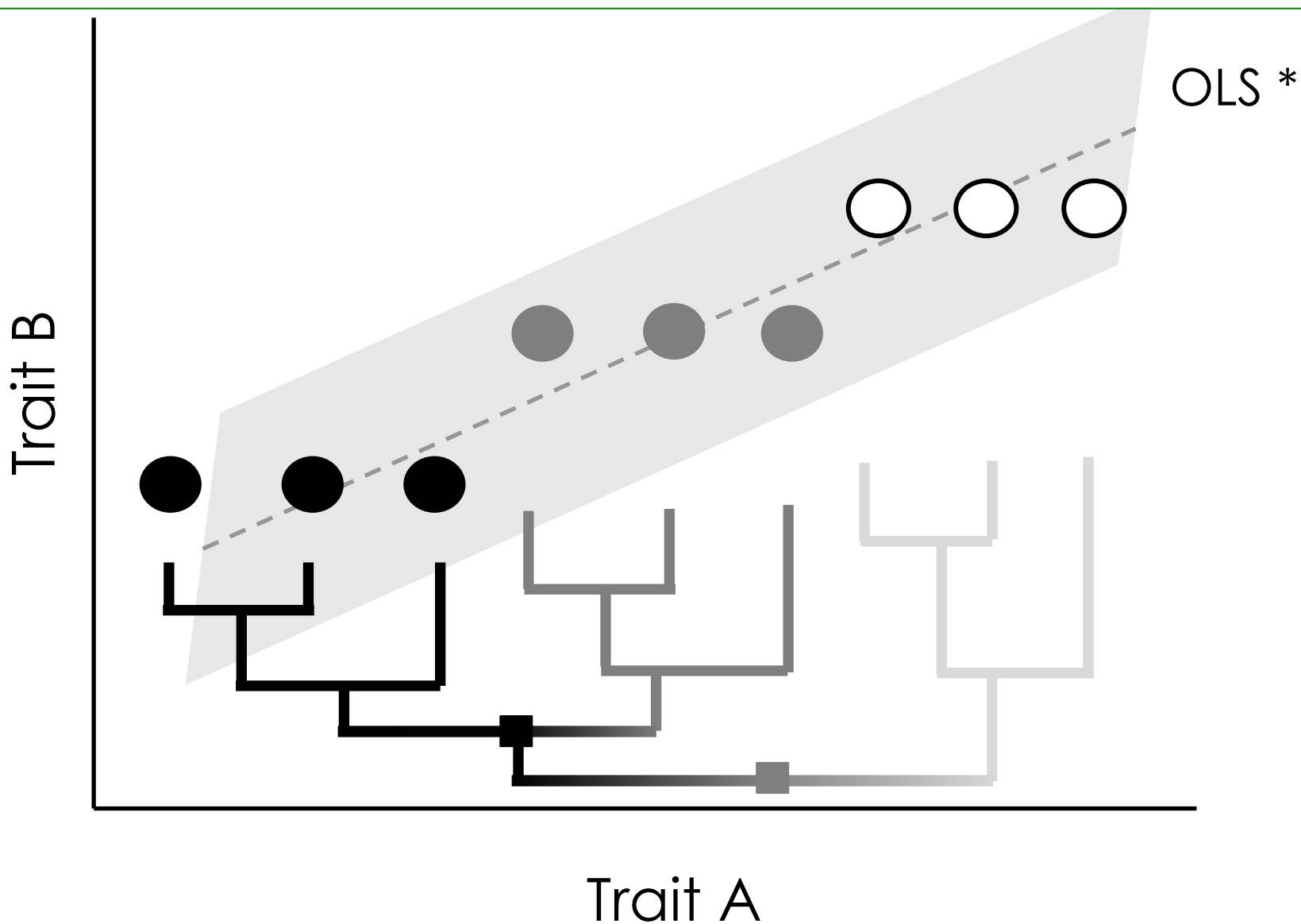
Accounting for phylogeny



from Clauss et al. (2013)



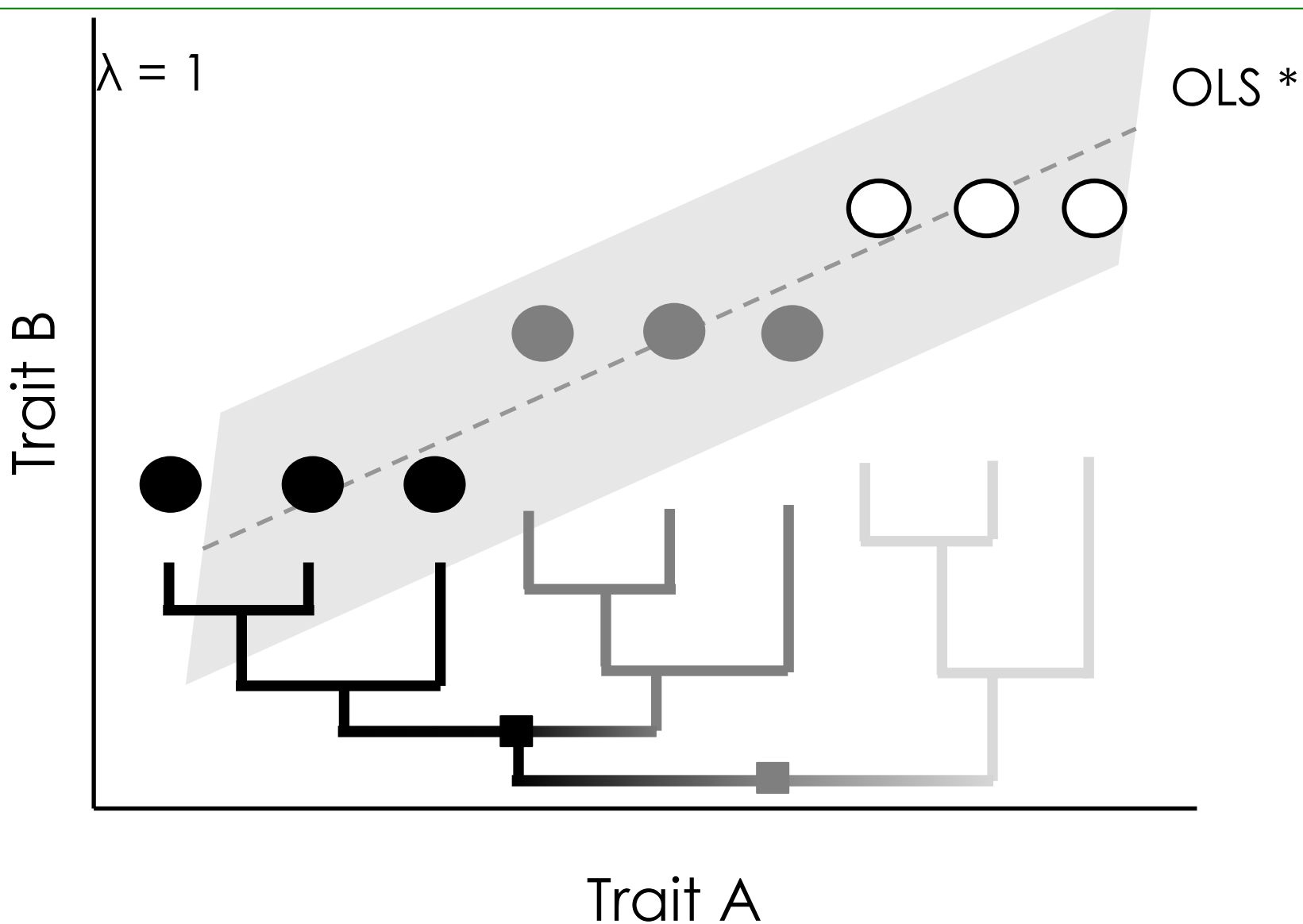
Accounting for phylogeny



from Clauss et al. (2013)

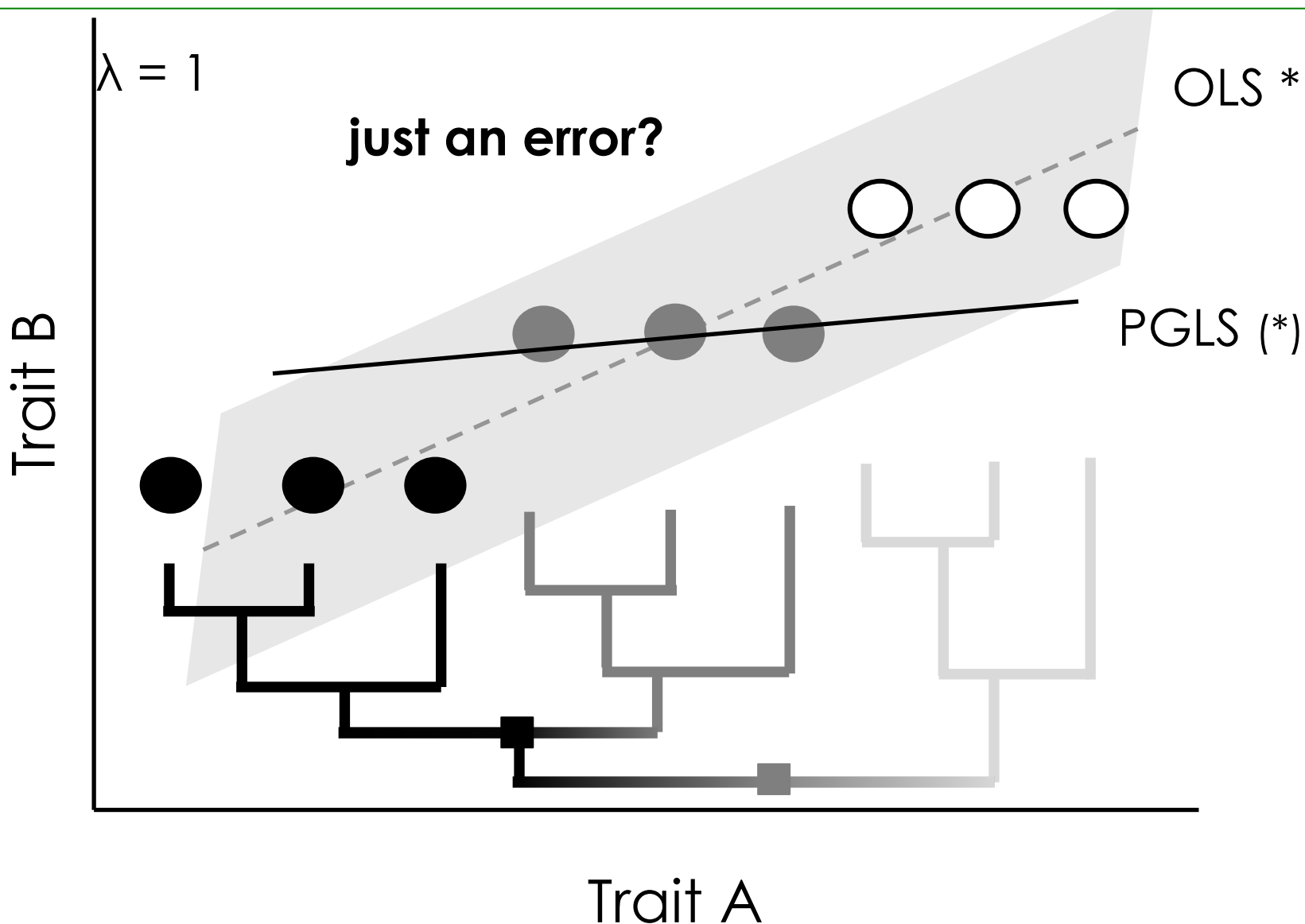


Accounting for phylogeny



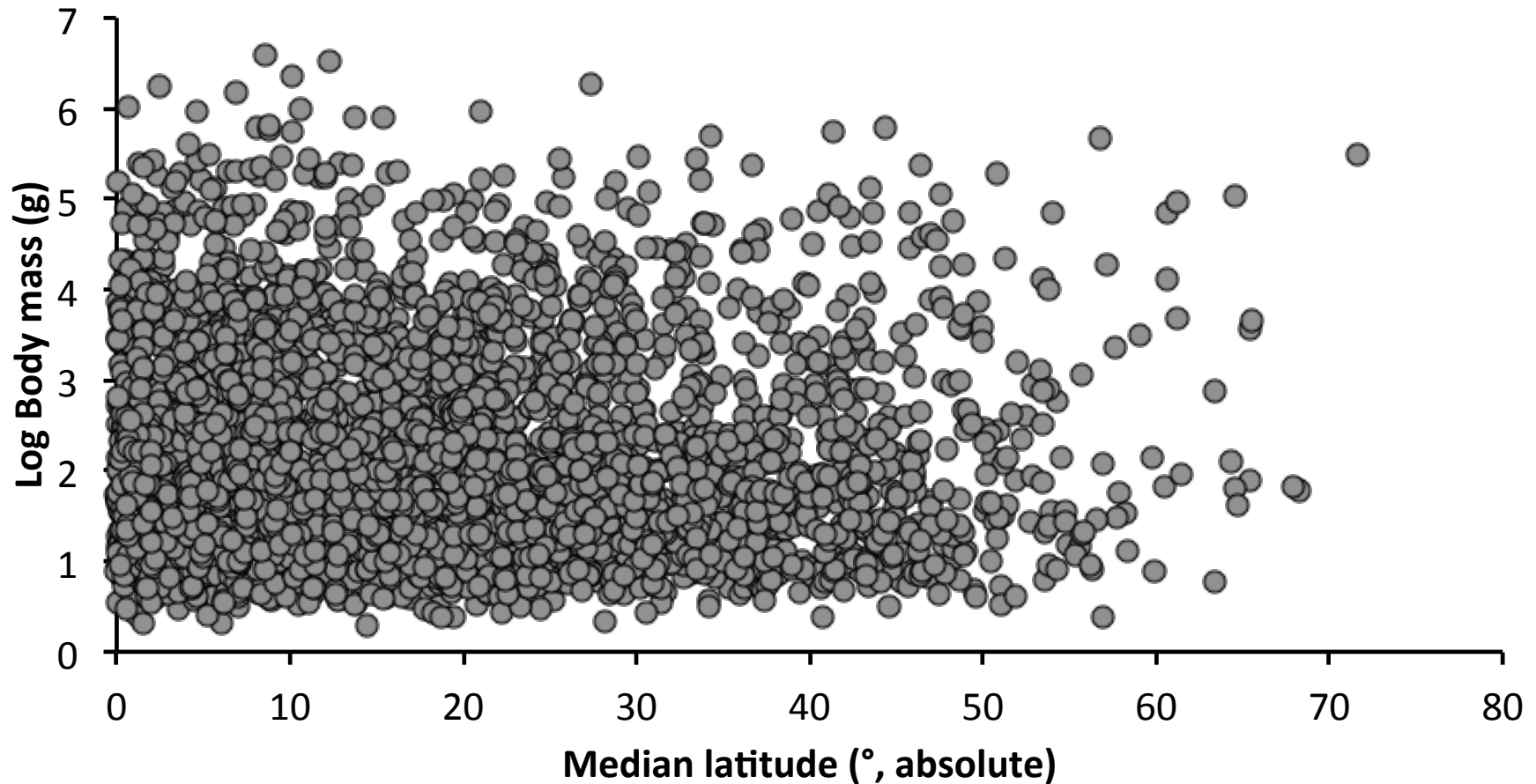


Accounting for phylogeny





Example IV: Bergmann's rule



from Clauss et al. (2013)



Accounting for phylogeny

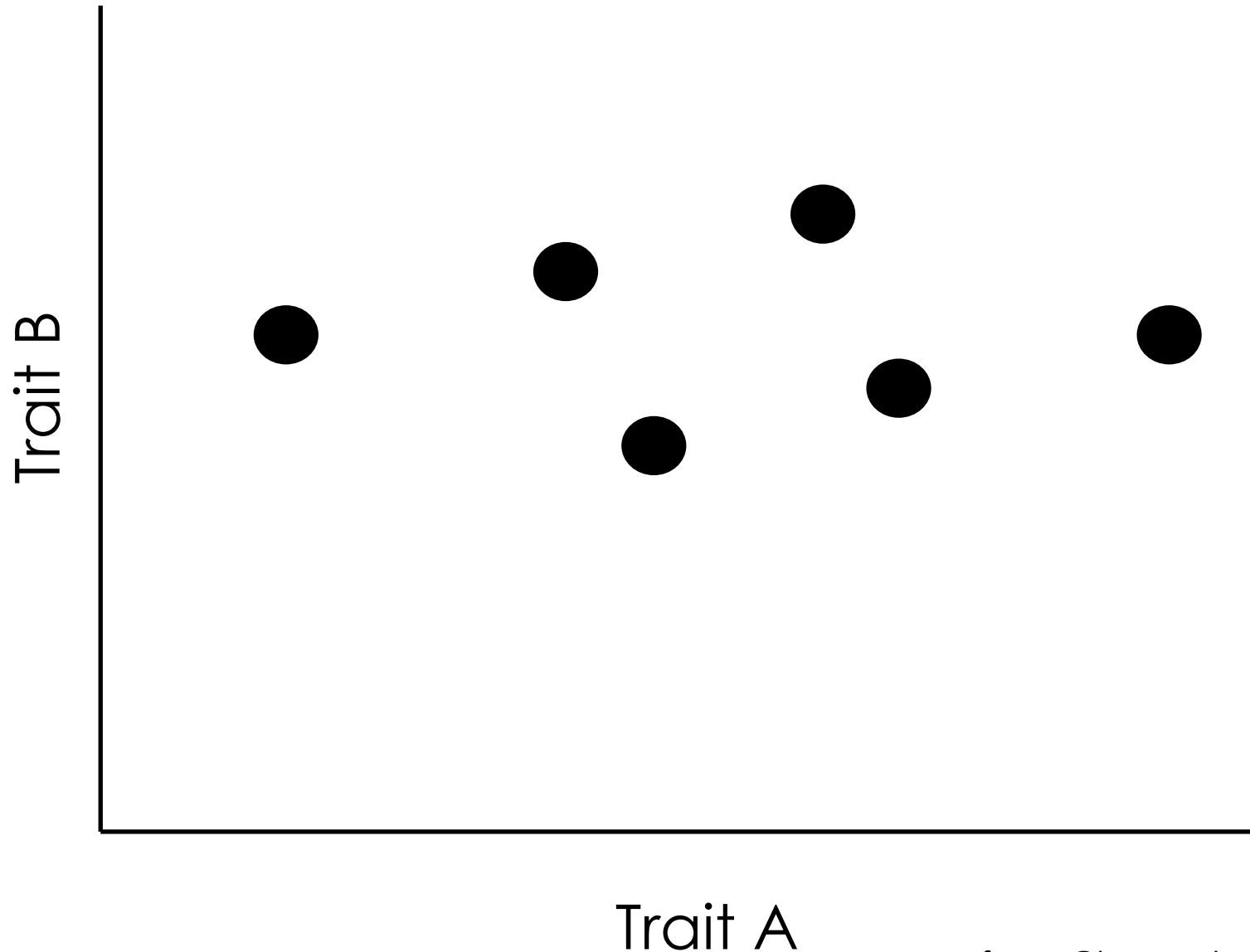
Trait B

Trait A

from Clauss et al. (2013)



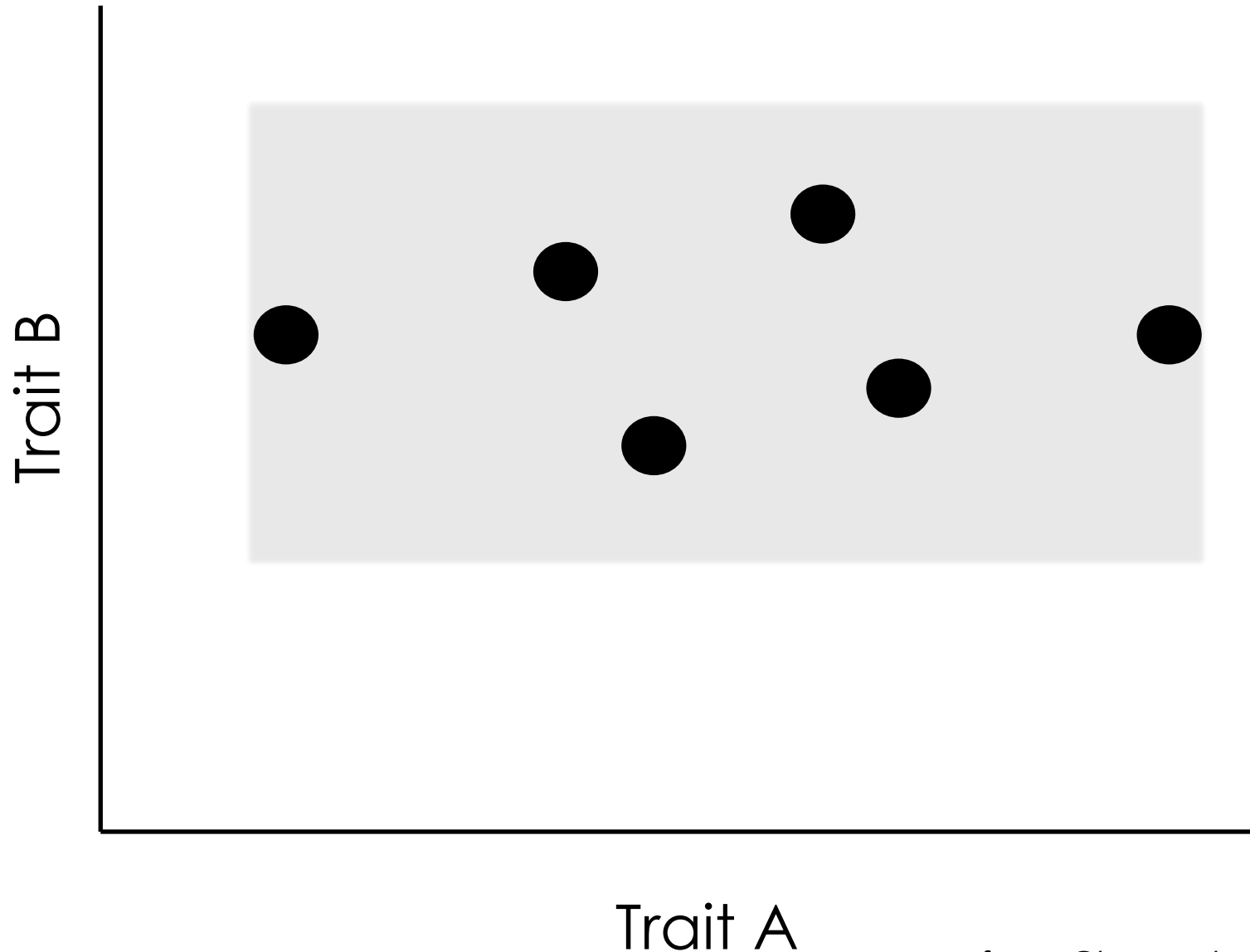
Accounting for phylogeny



from Clauss et al. (2013)



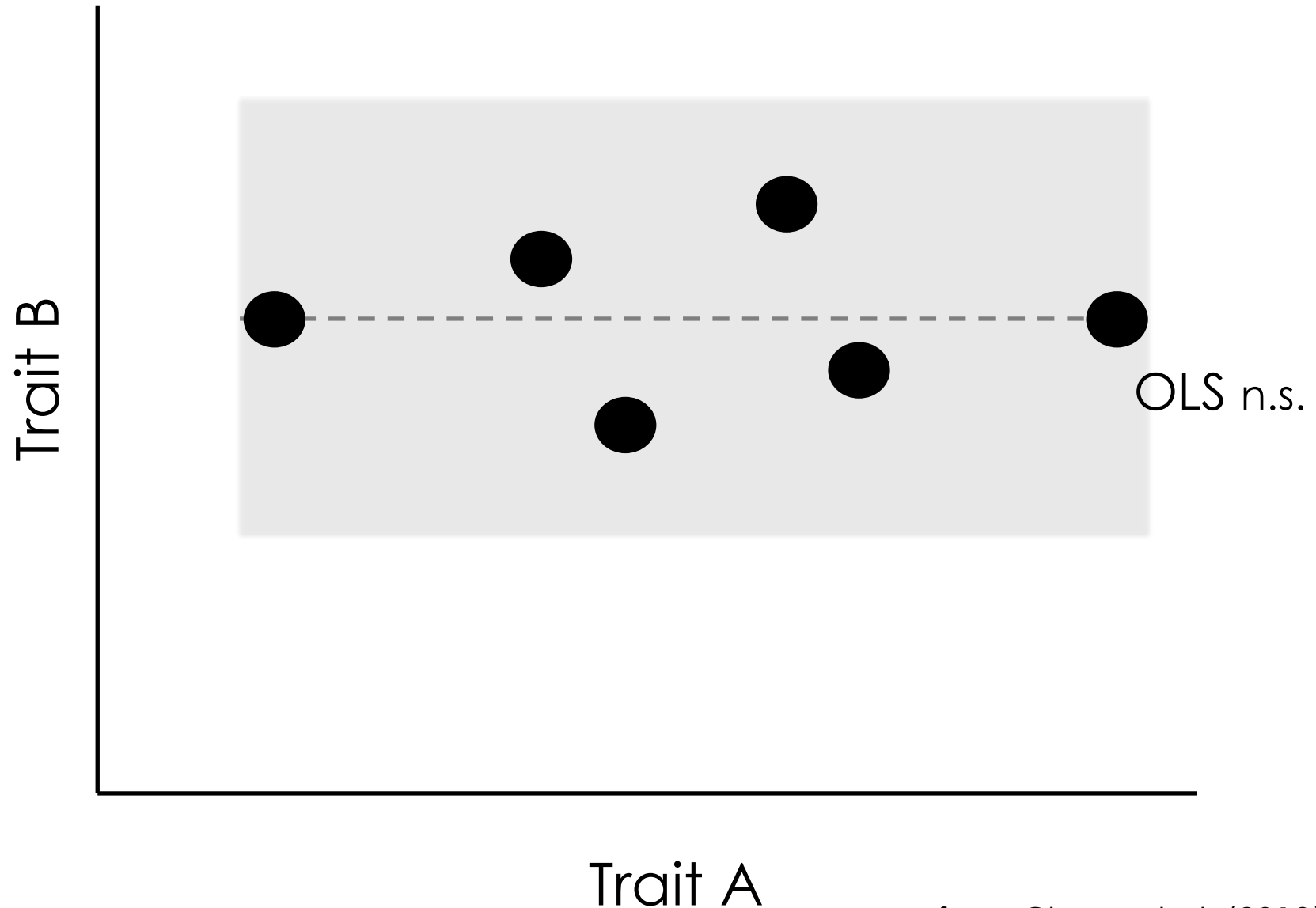
Accounting for phylogeny



from Clauss et al. (2013)



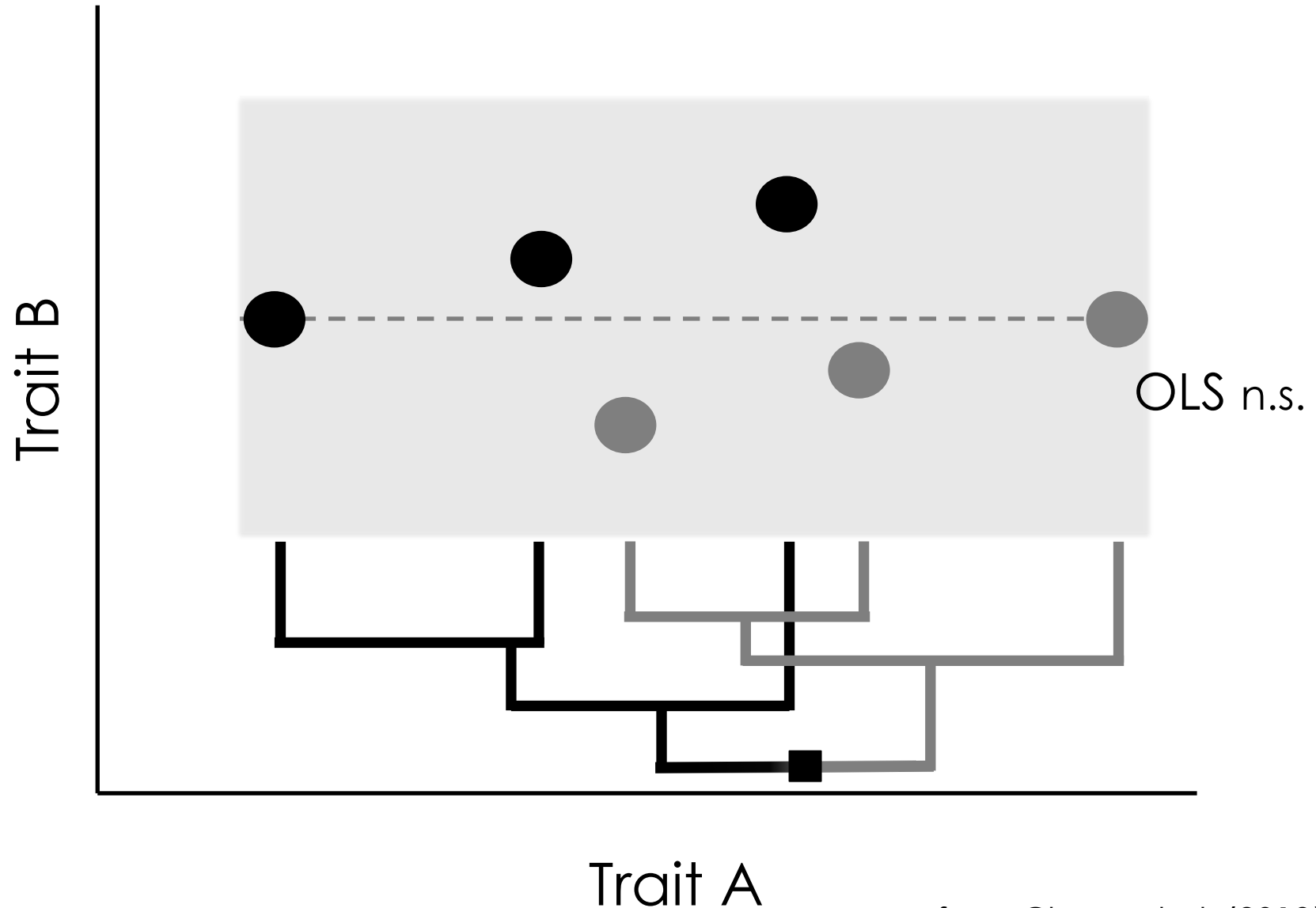
Accounting for phylogeny



from Clauss et al. (2013)



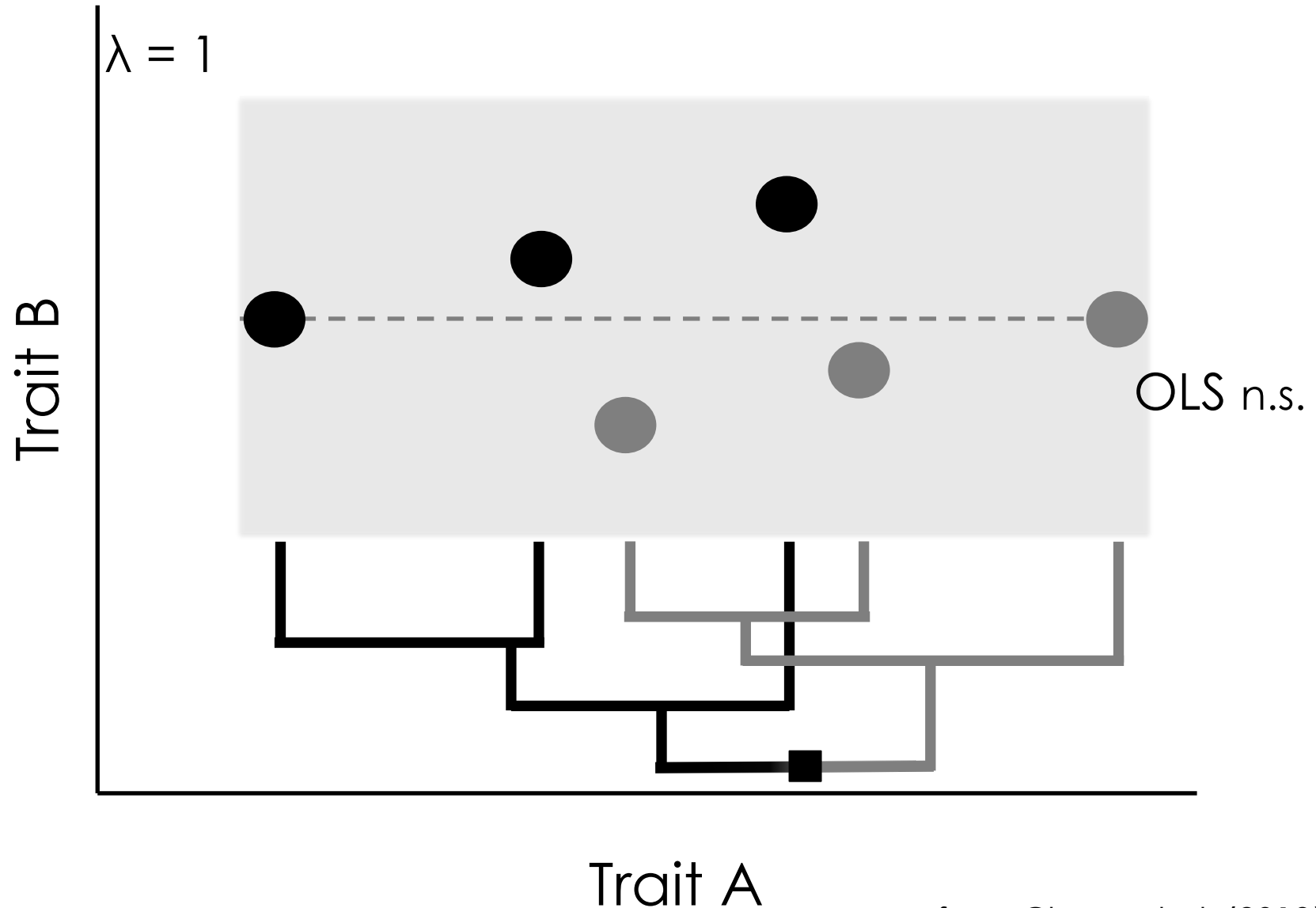
Accounting for phylogeny



from Clauss et al. (2013)

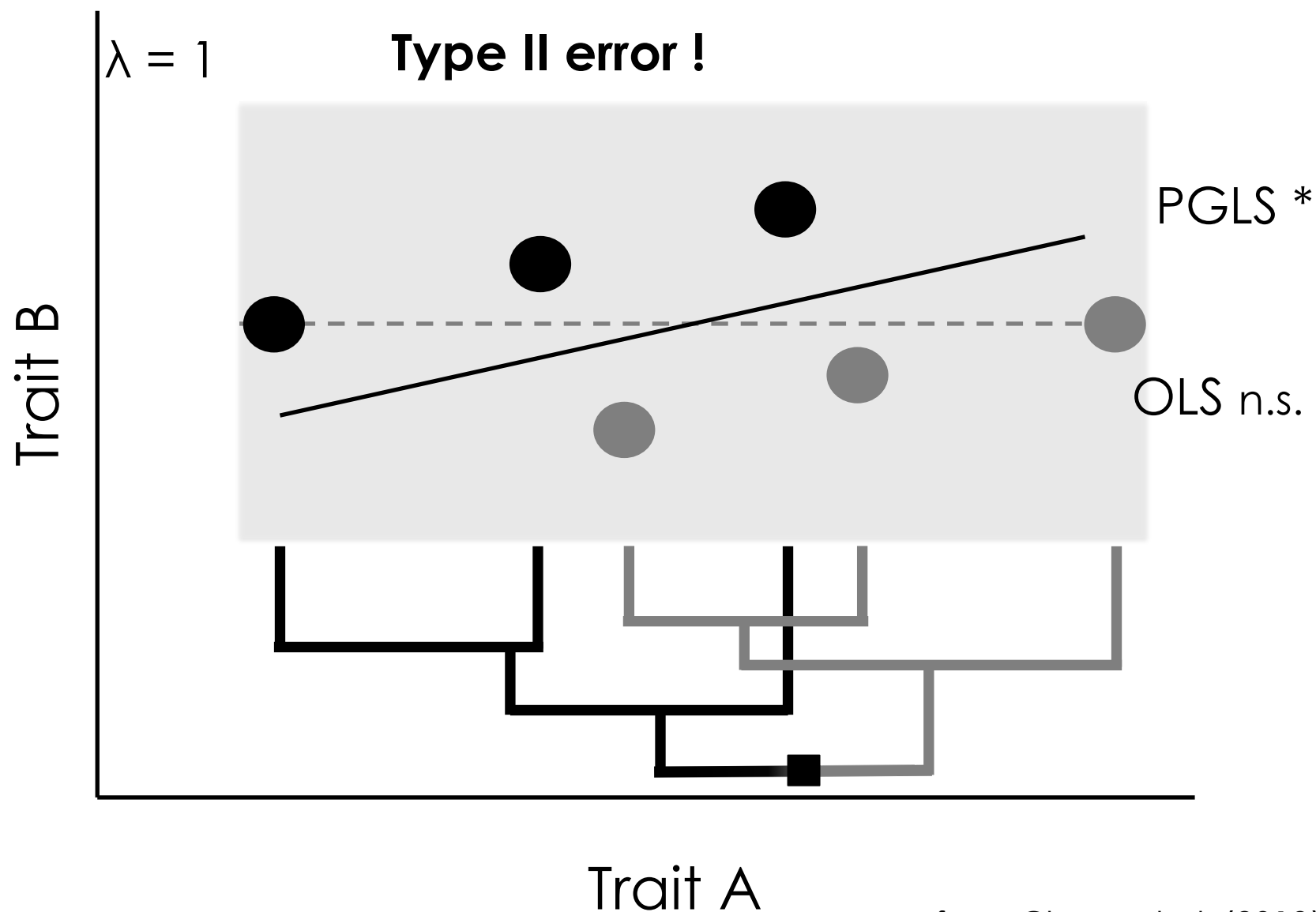


Accounting for phylogeny



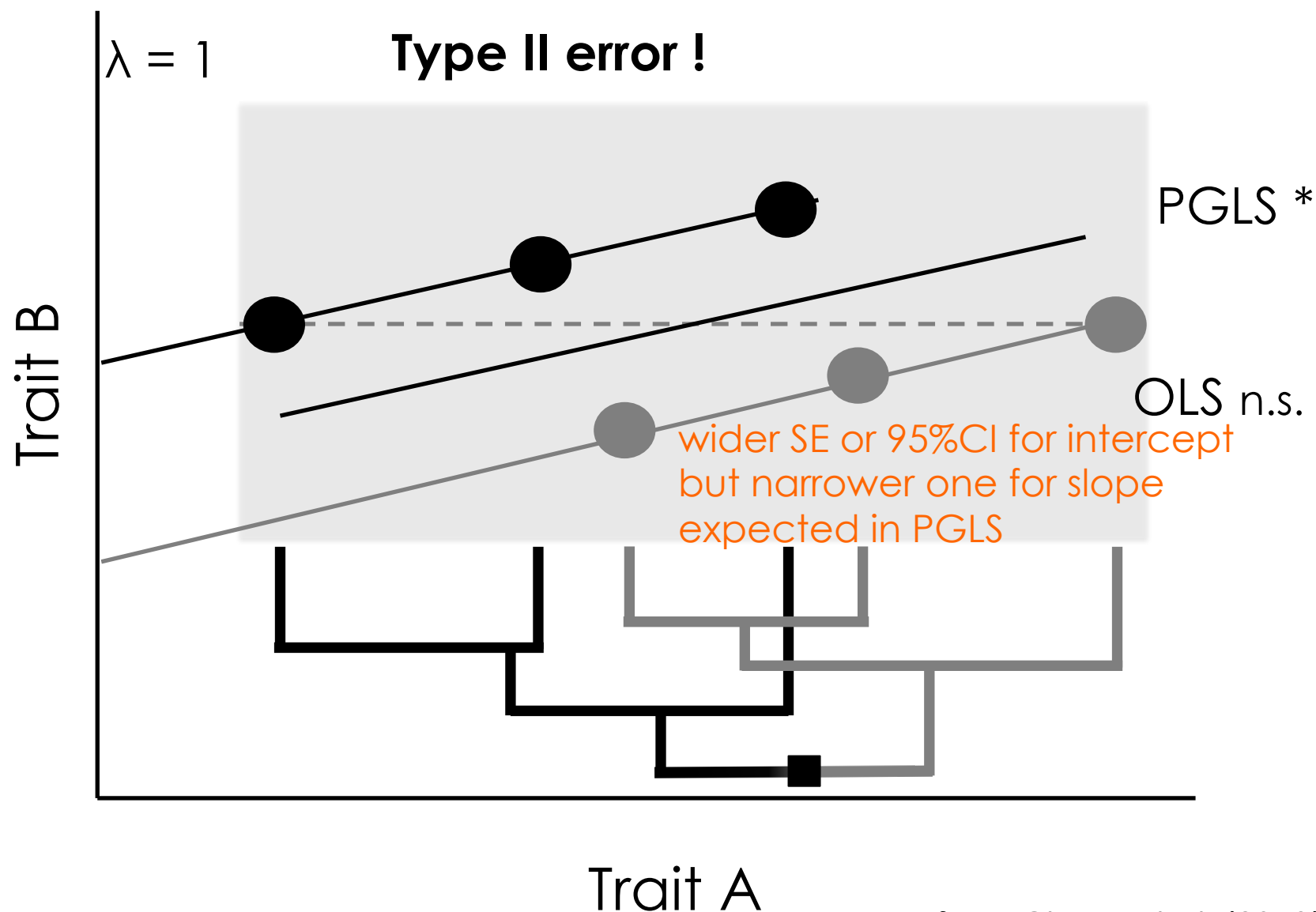


Accounting for phylogeny



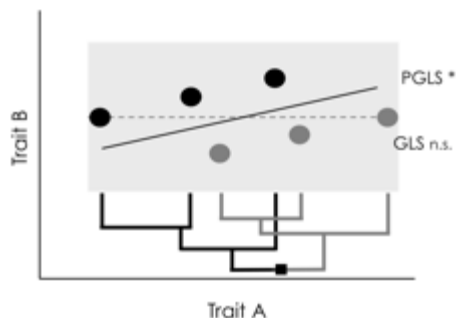


Accounting for phylogeny

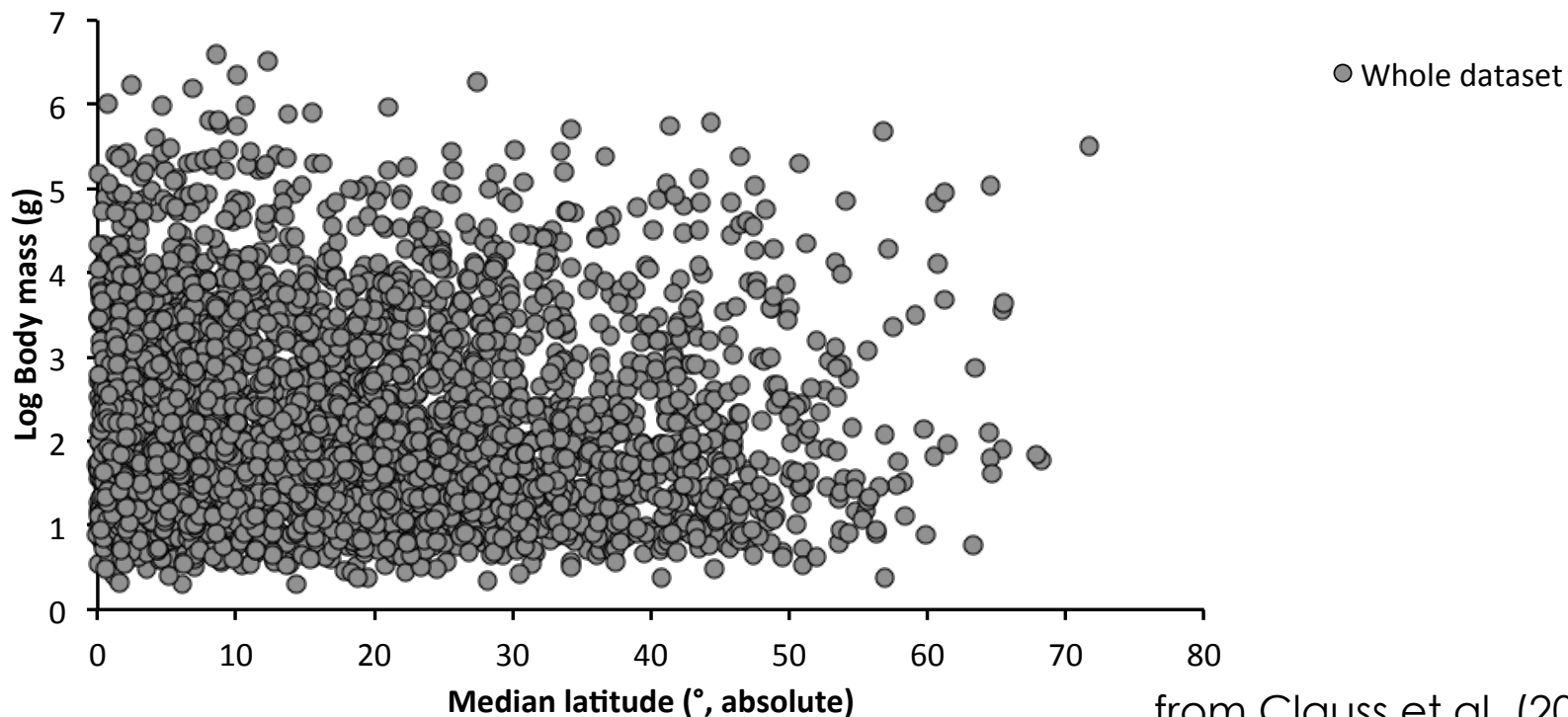




Example IV: Bergmann's rule



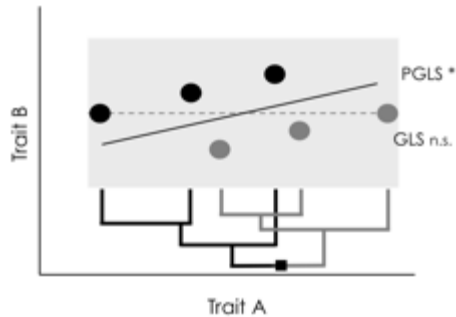
OLS: $2.19 (\pm 0.03) - 0.0012 (\pm 0.0013) \text{ Lat.}$
PGLS: $2.79 (\pm 0.47) + 0.0016 (\pm 0.0005) \text{ Lat.}$



from Clauss et al. (2013)

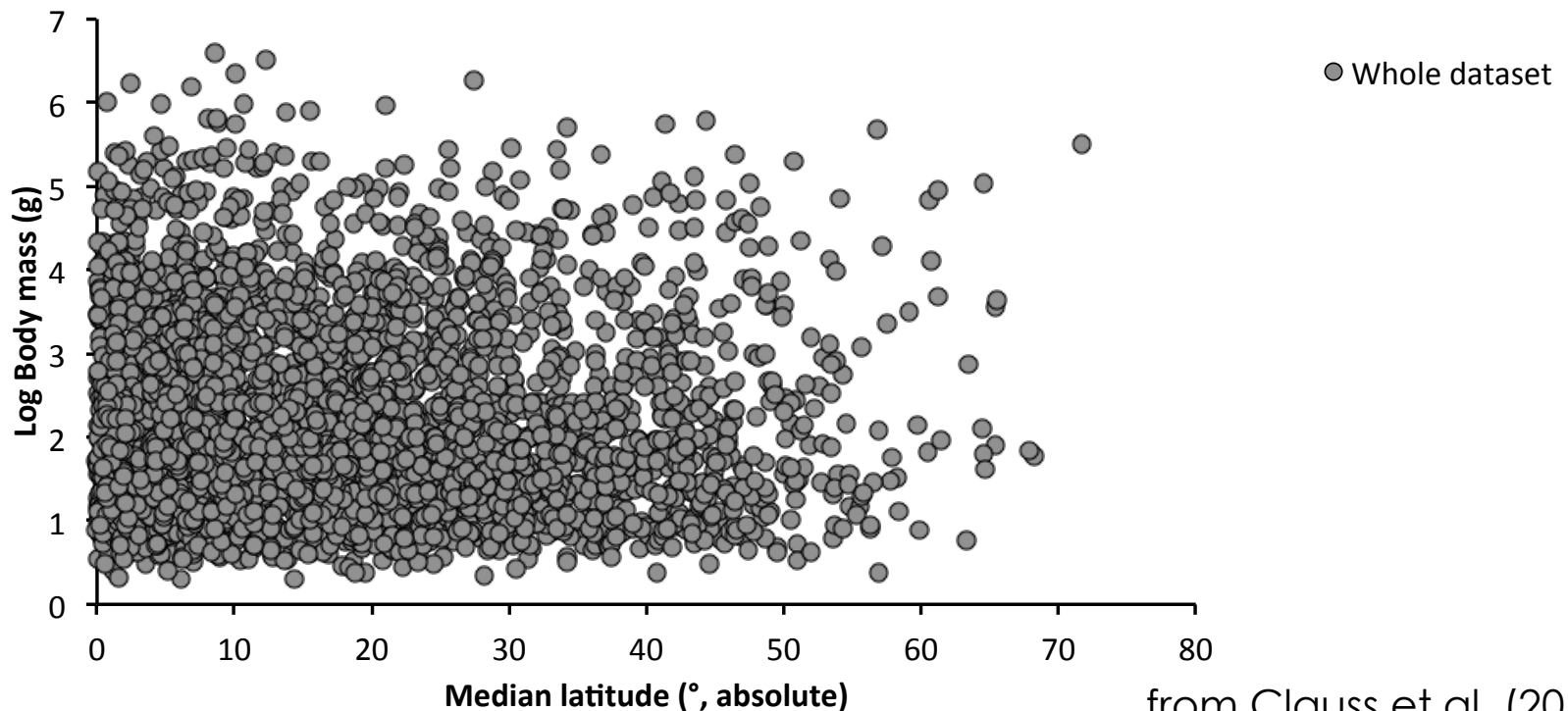


Example IV: Bergmann's rule



OLS: $2.19 (\pm 0.03) - 0.0012 (\pm 0.0013) \text{ Lat.}$

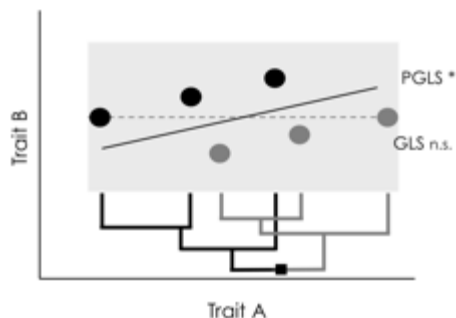
PGLS: $2.79 (\pm 0.47) + 0.0016 (\pm 0.0005) \text{ Lat.}$



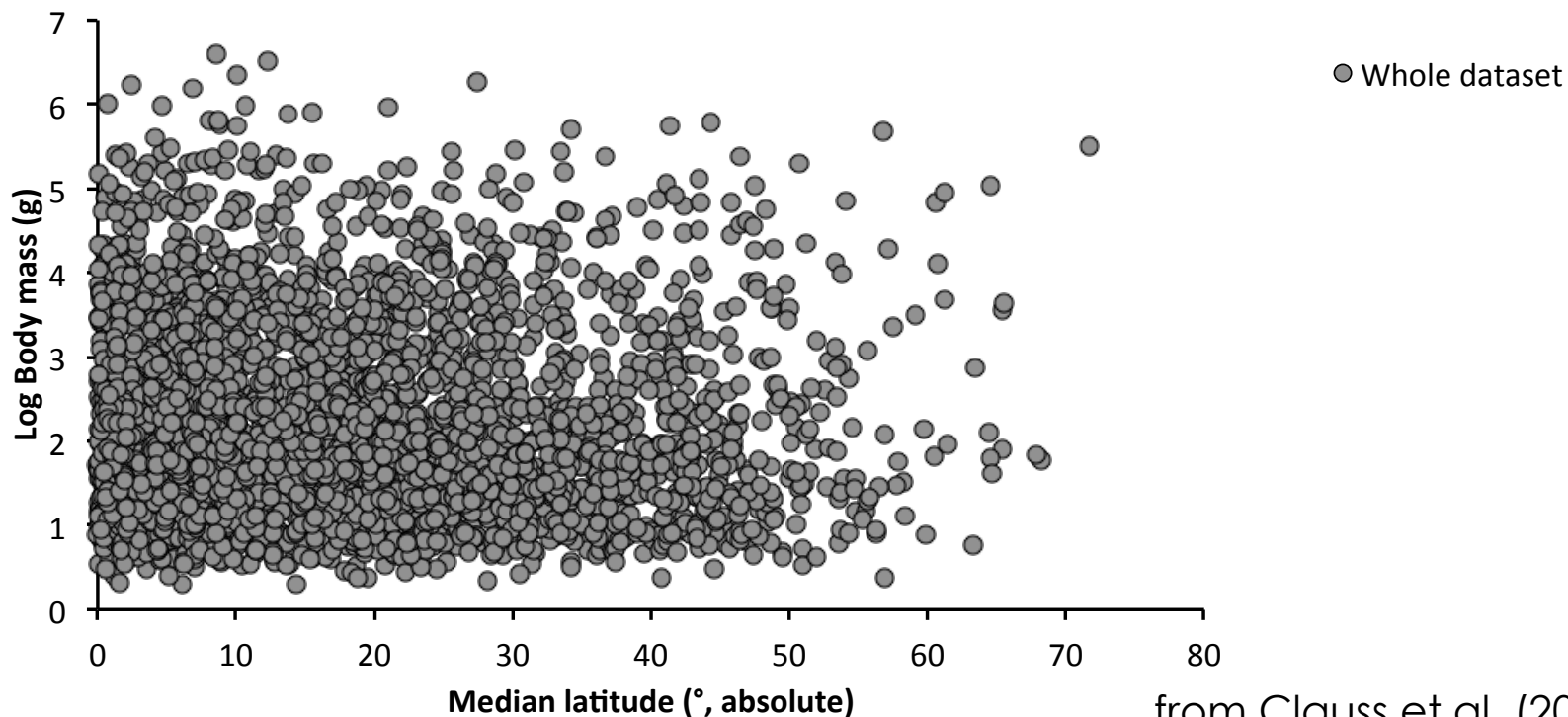
from Clauss et al. (2013)



Example IV: Bergmann's rule



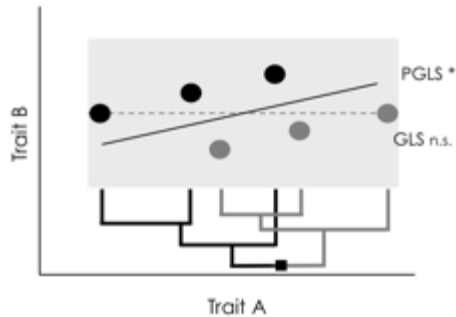
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from Clauss et al. (2013)

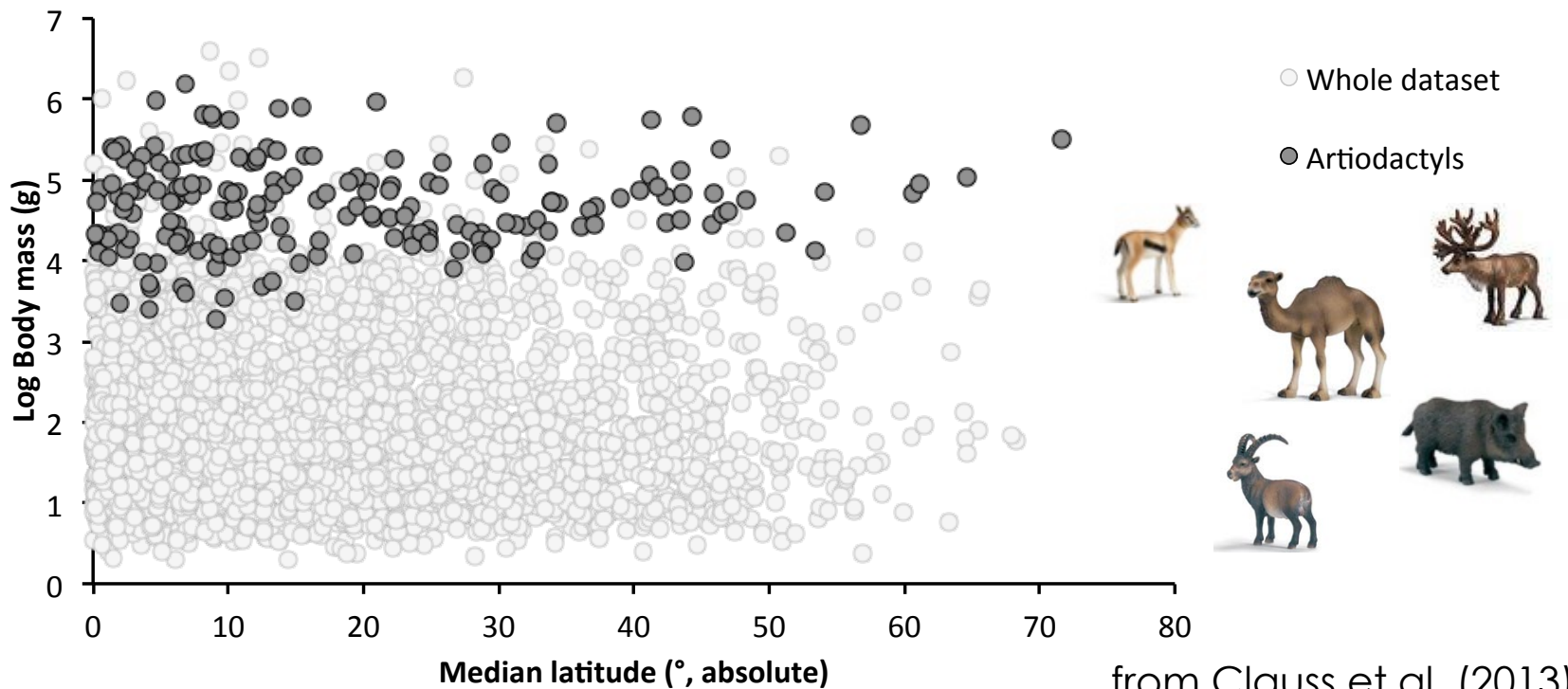


Example IV: Bergmann's rule



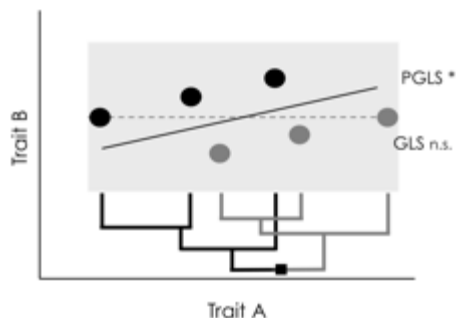
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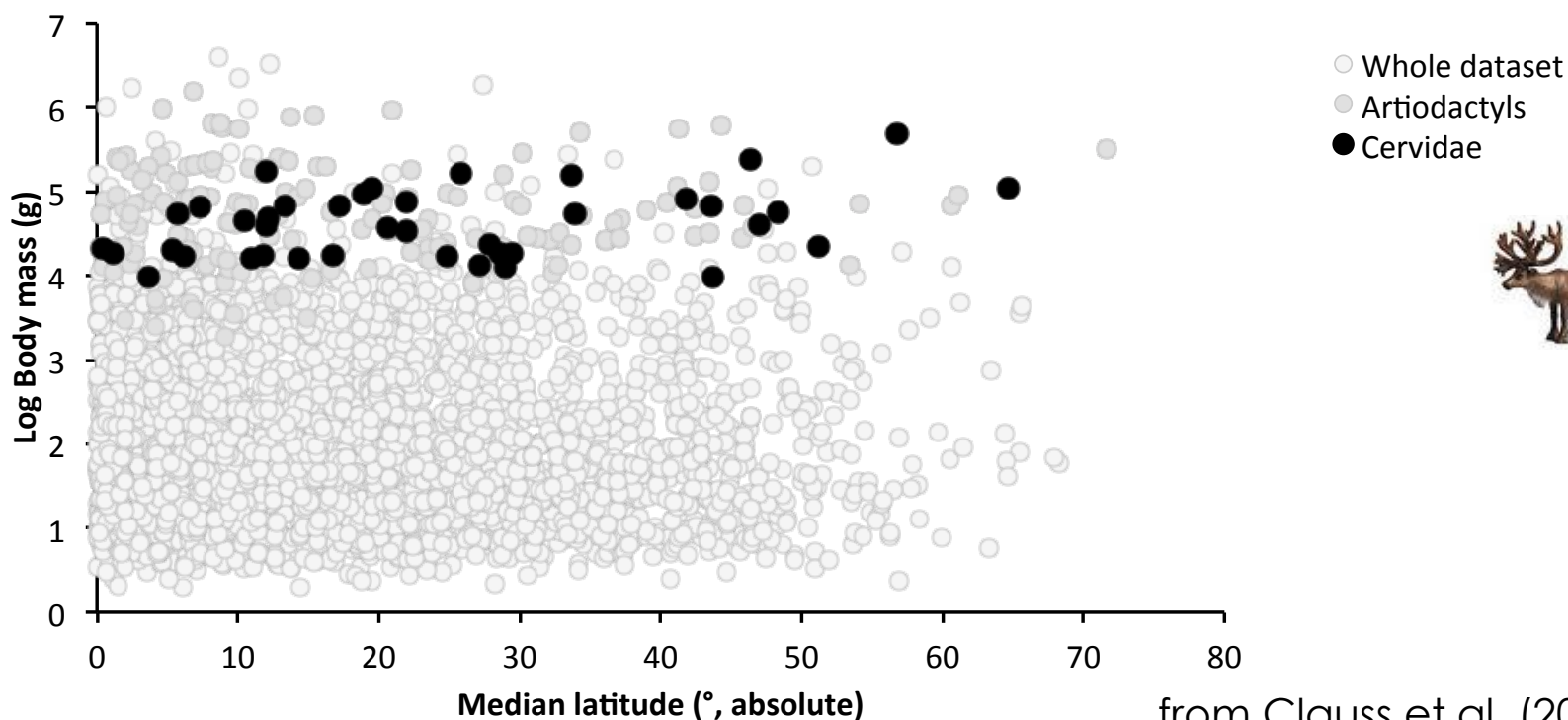




Example IV: Bergmann's rule

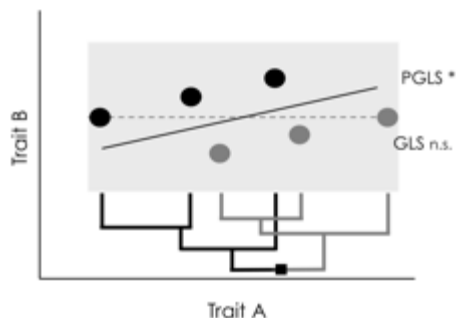


OLS: $2.19 (\pm 0.03) - 0.0012 (\pm 0.0013) \text{ Lat.}$
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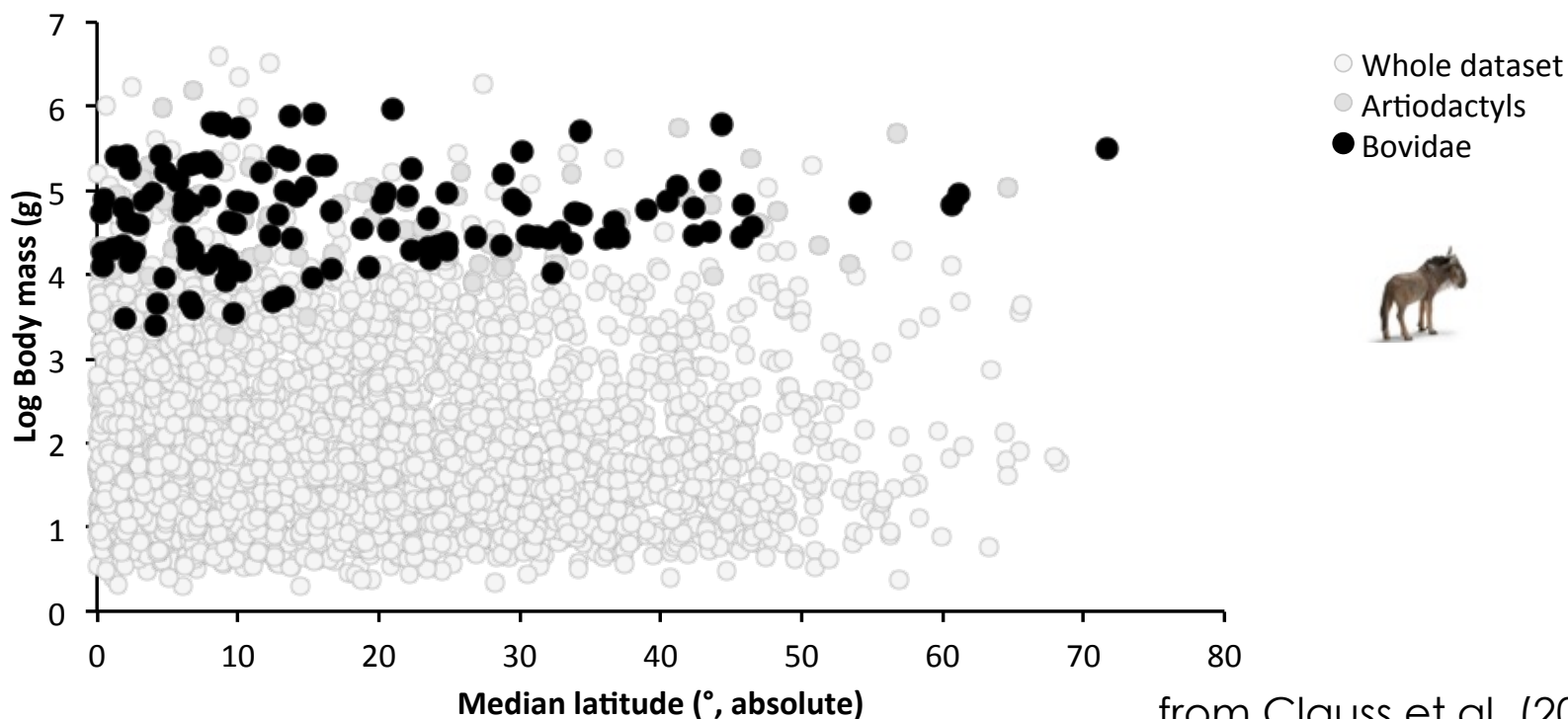




Example IV: Bergmann's rule

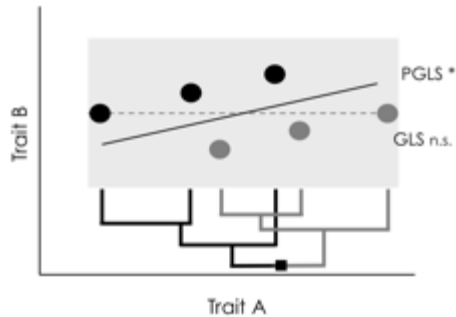


OLS: $2.19 (\pm 0.03) - 0.0012 (\pm 0.0013) \text{ Lat.}$
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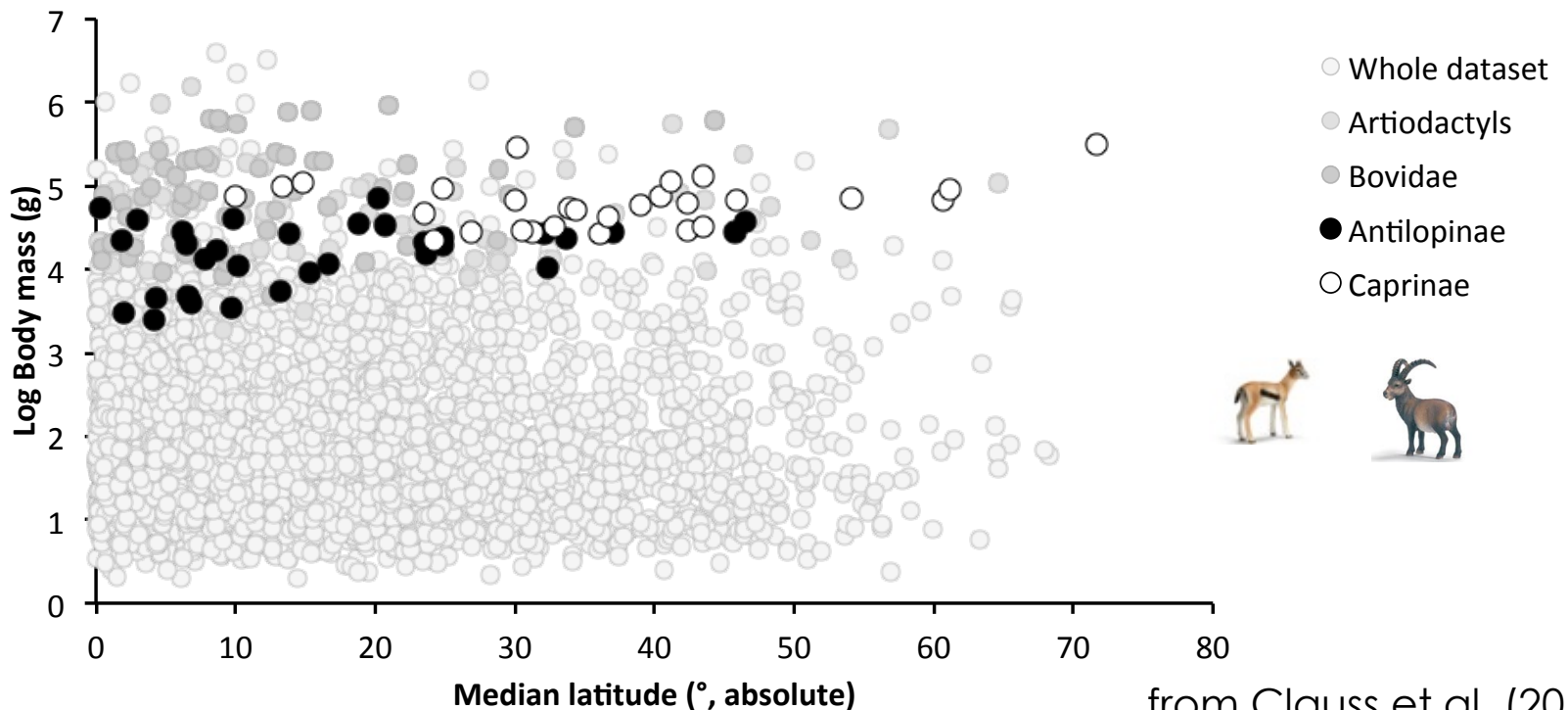




Example IV: Bergmann's rule

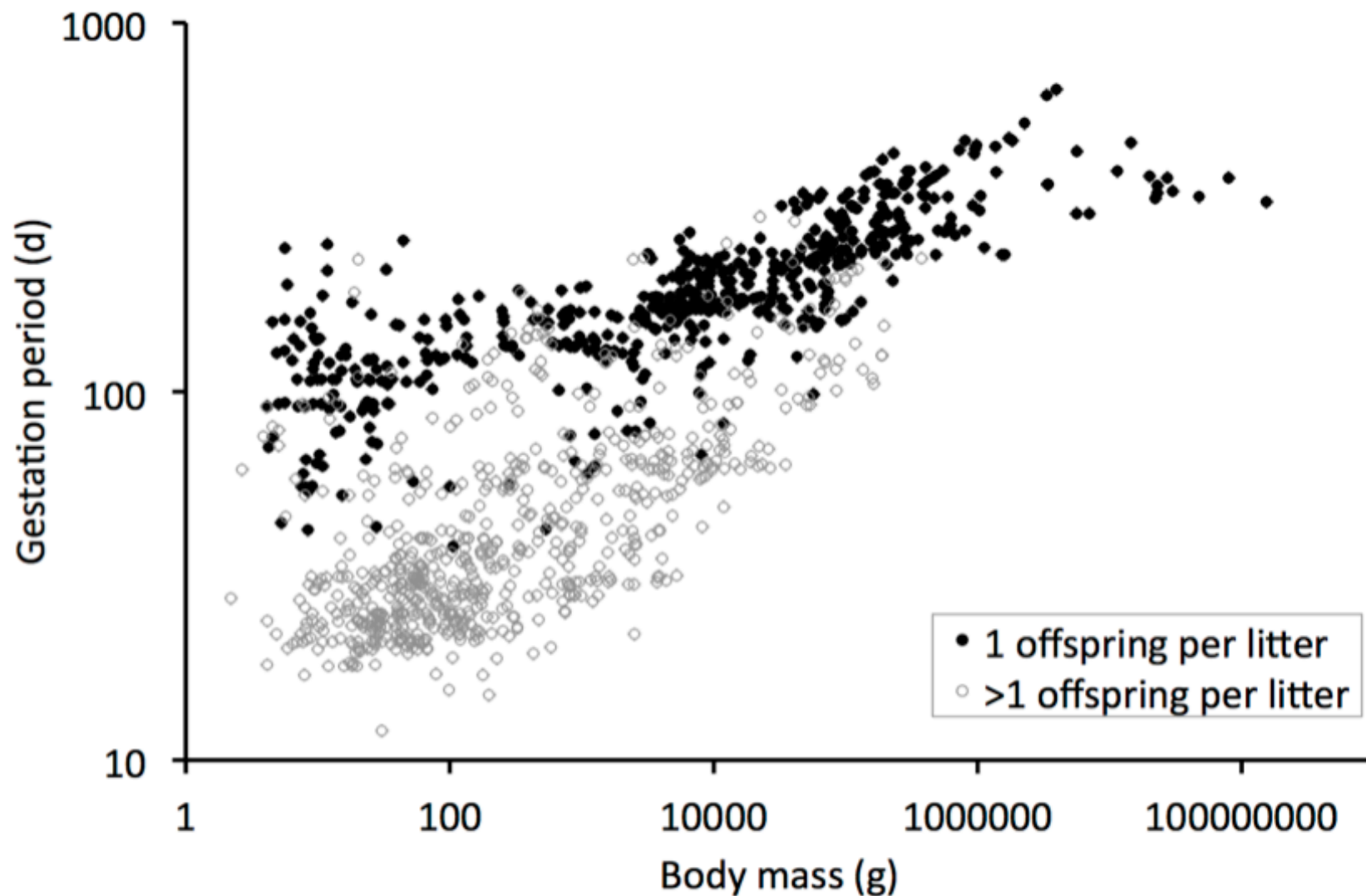


OLS: $2.19 (\pm 0.03) - 0.0012 (\pm 0.0013) \text{ Lat.}$
PGLS: $2.79 (\pm 0.47) + 0.0016 (\pm 0.0005) \text{ Lat.}$





Example V: Gestation time



from Clauss et al. (2014)



Accounting for phylogeny

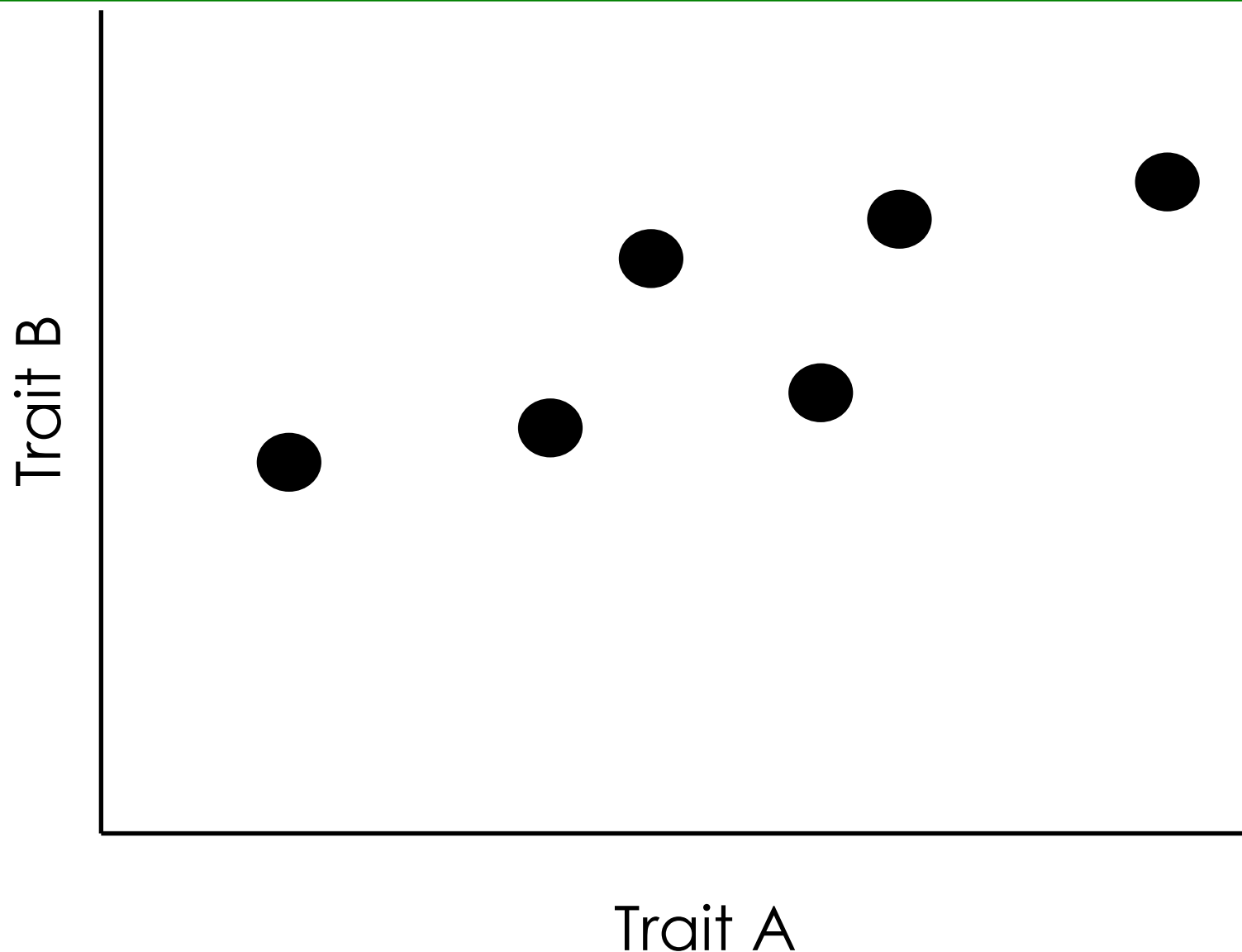
Trait B

Trait A

from Clauss et al. (2013)



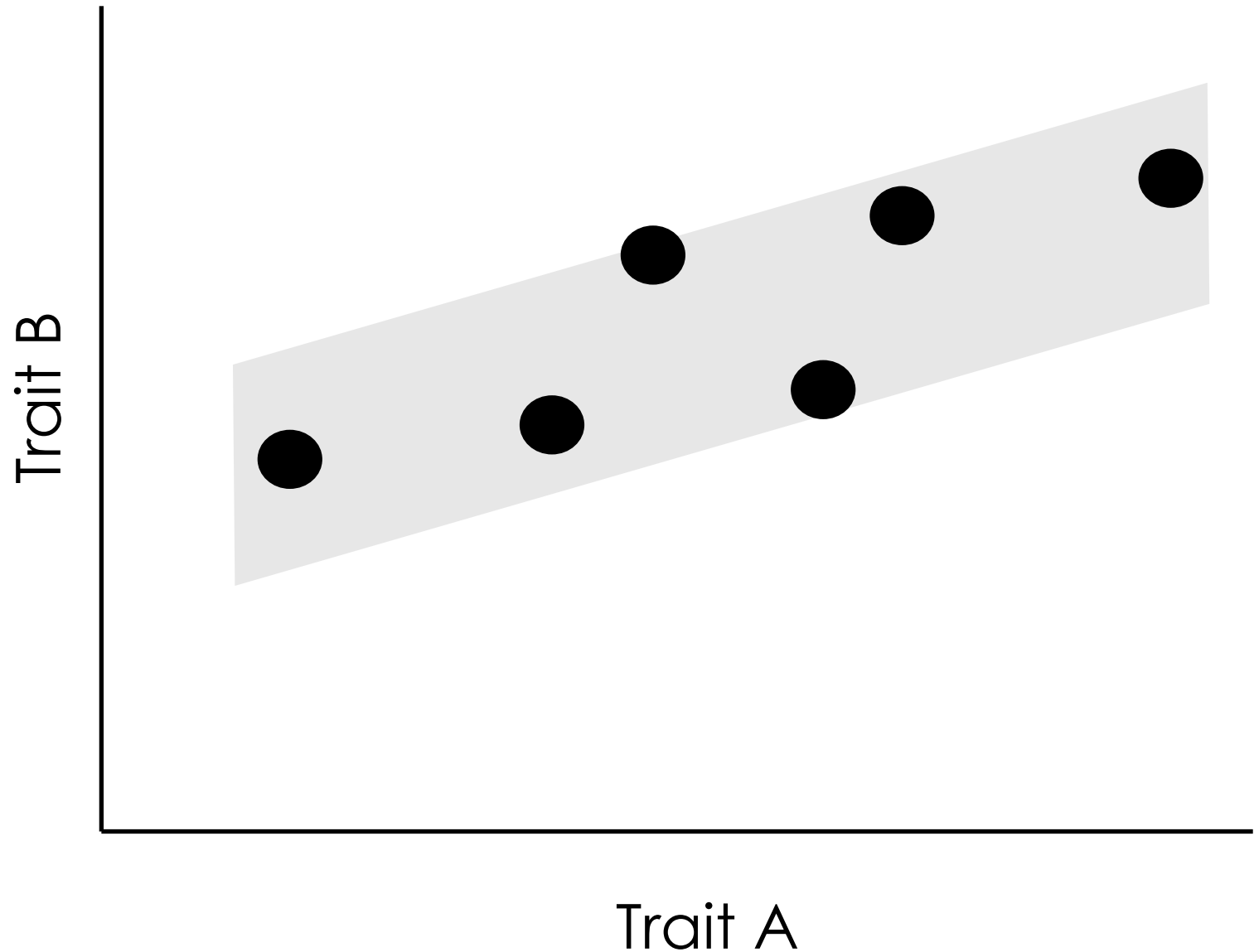
Accounting for phylogeny



from Clauss et al. (2013)



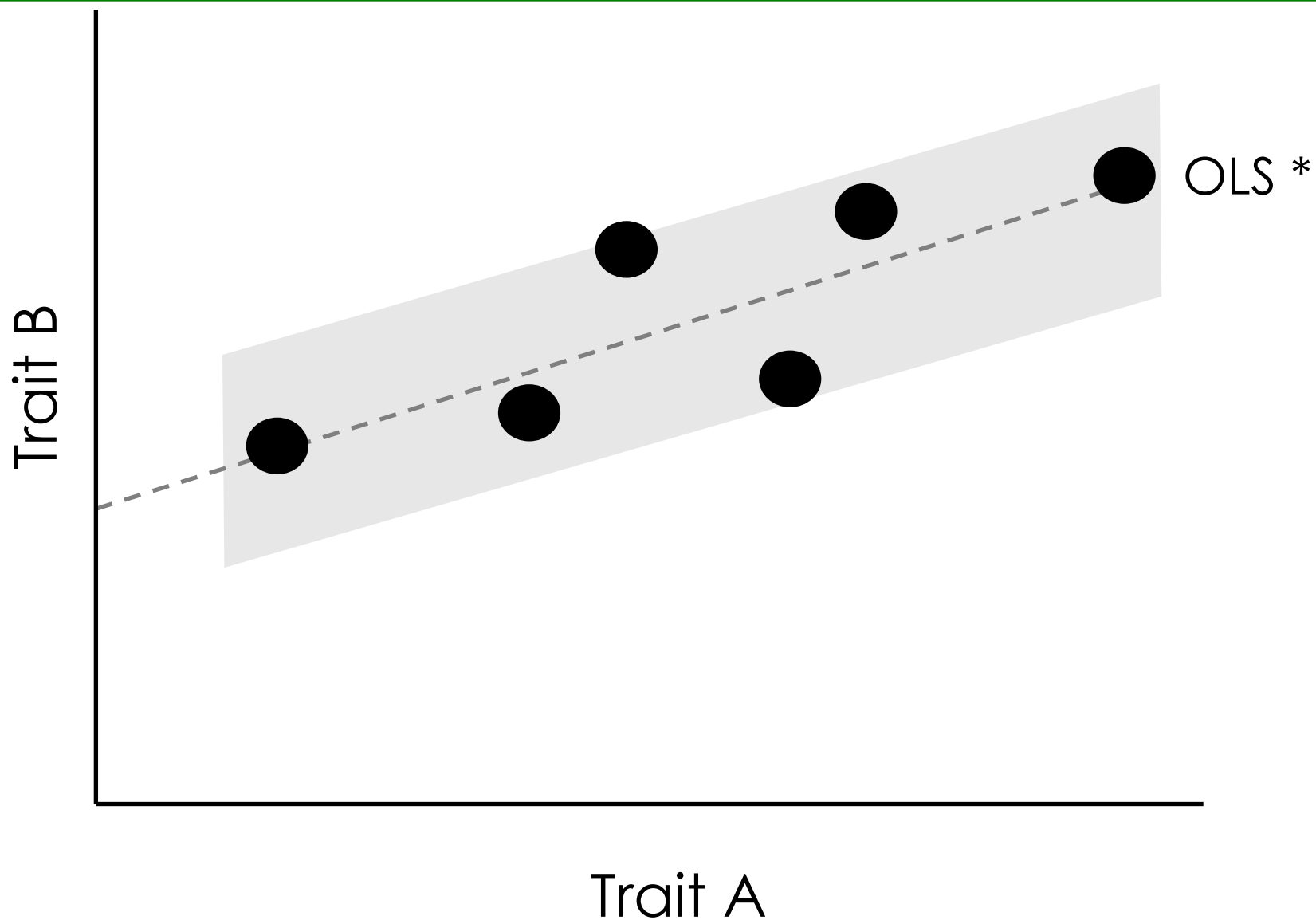
Accounting for phylogeny



from Clauss et al. (2013)



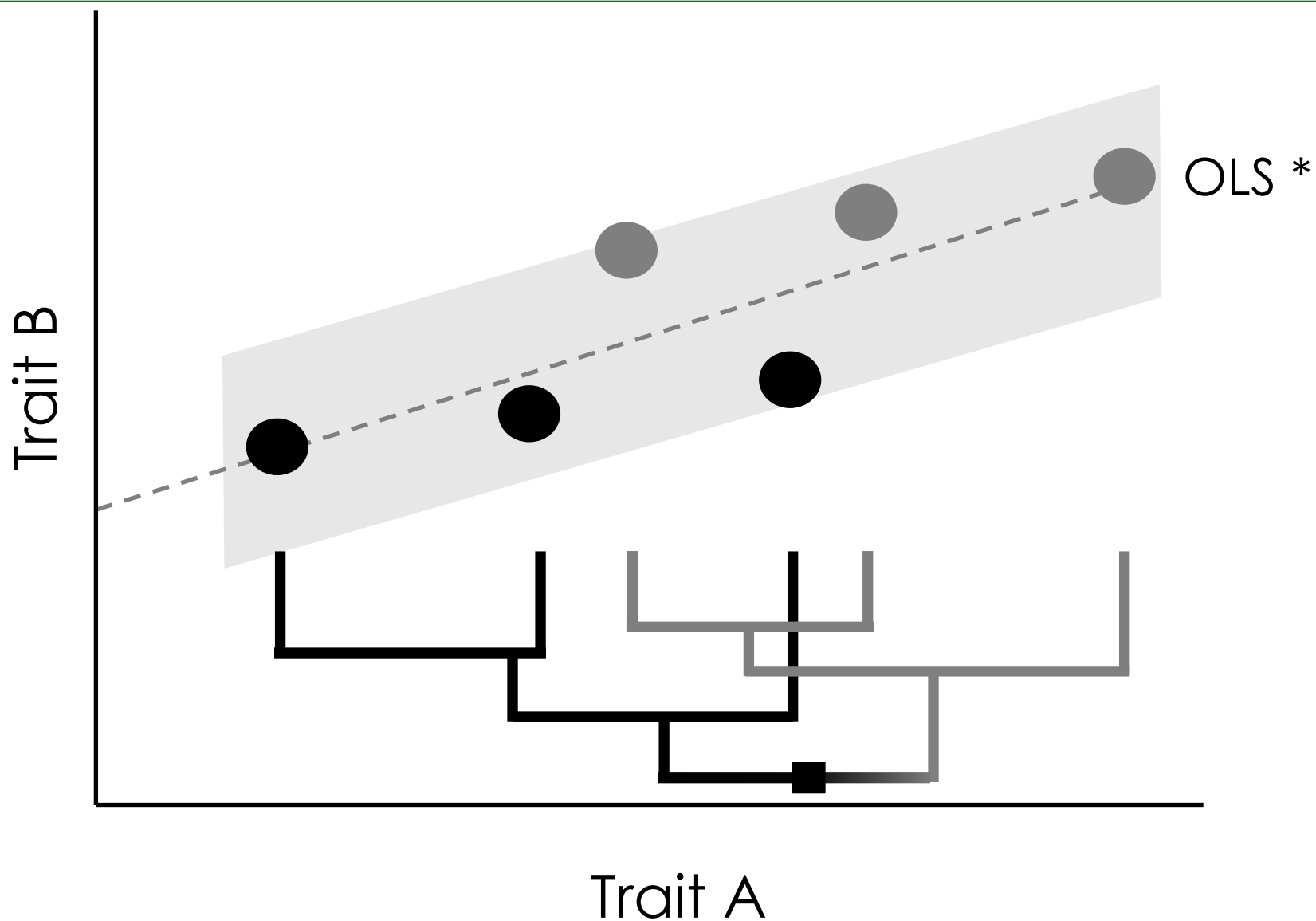
Accounting for phylogeny



from Clauss et al. (2013)



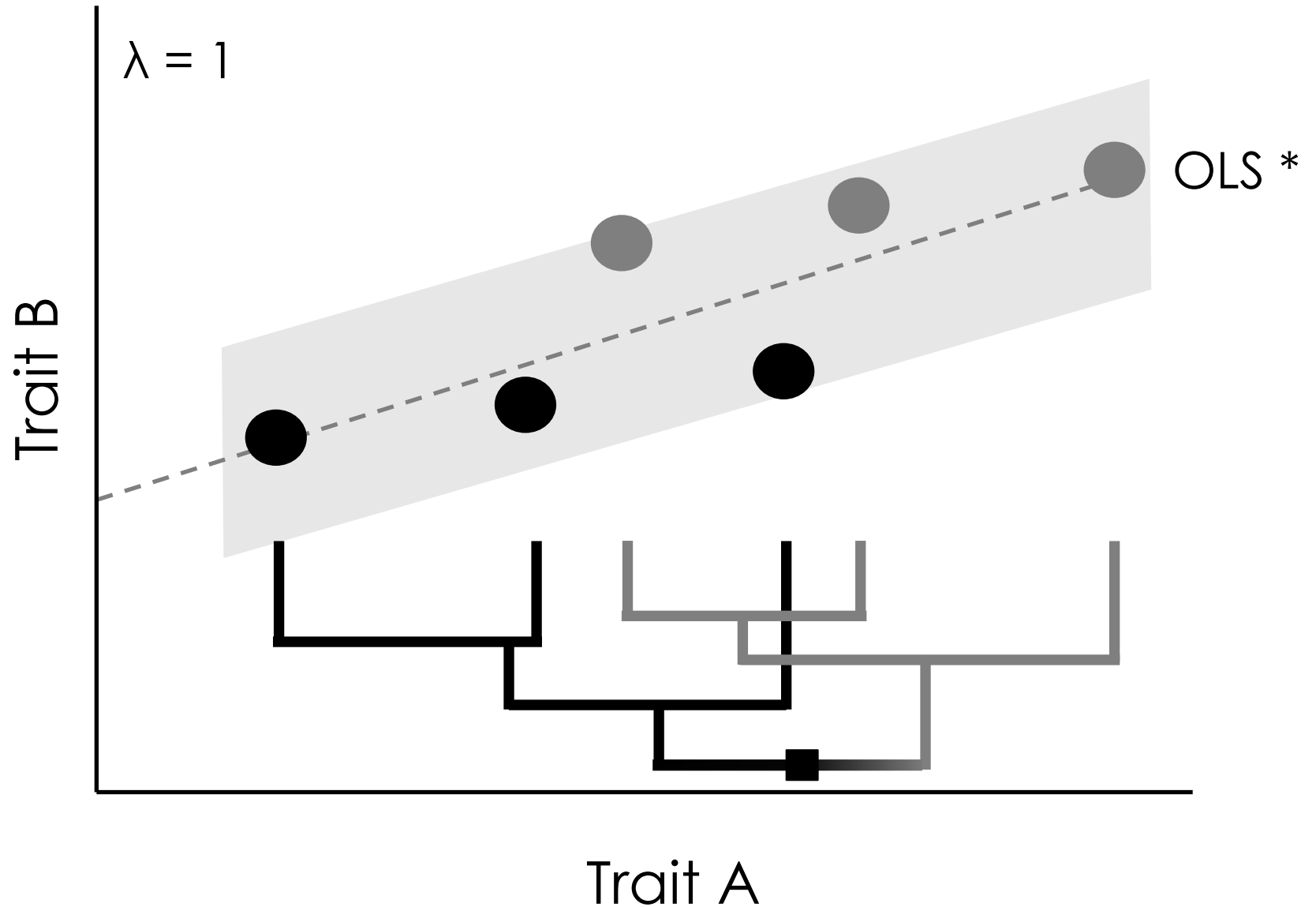
Accounting for phylogeny

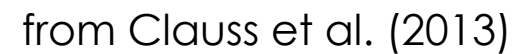


from Clauss et al. (2013)



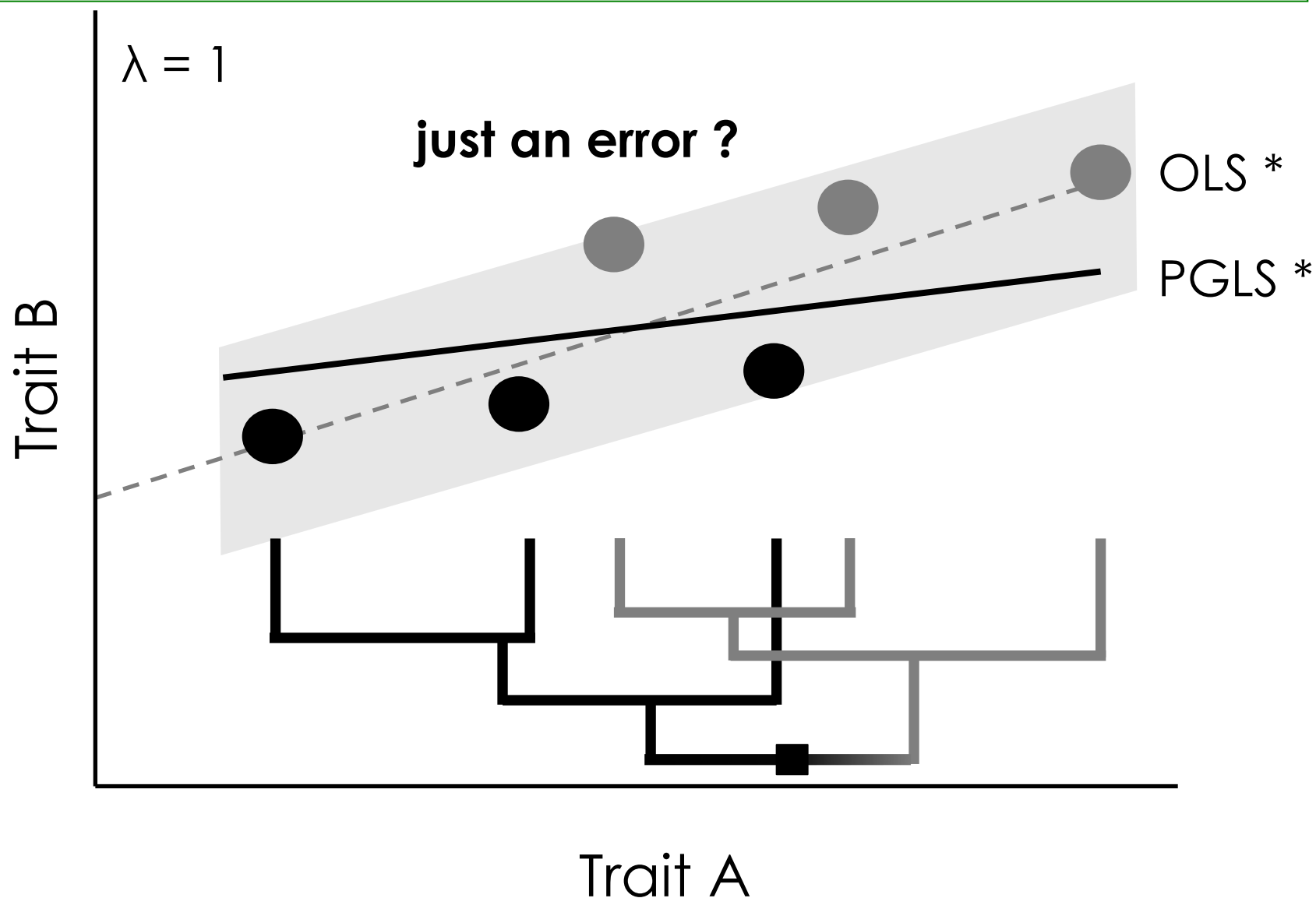
Accounting for phylogeny





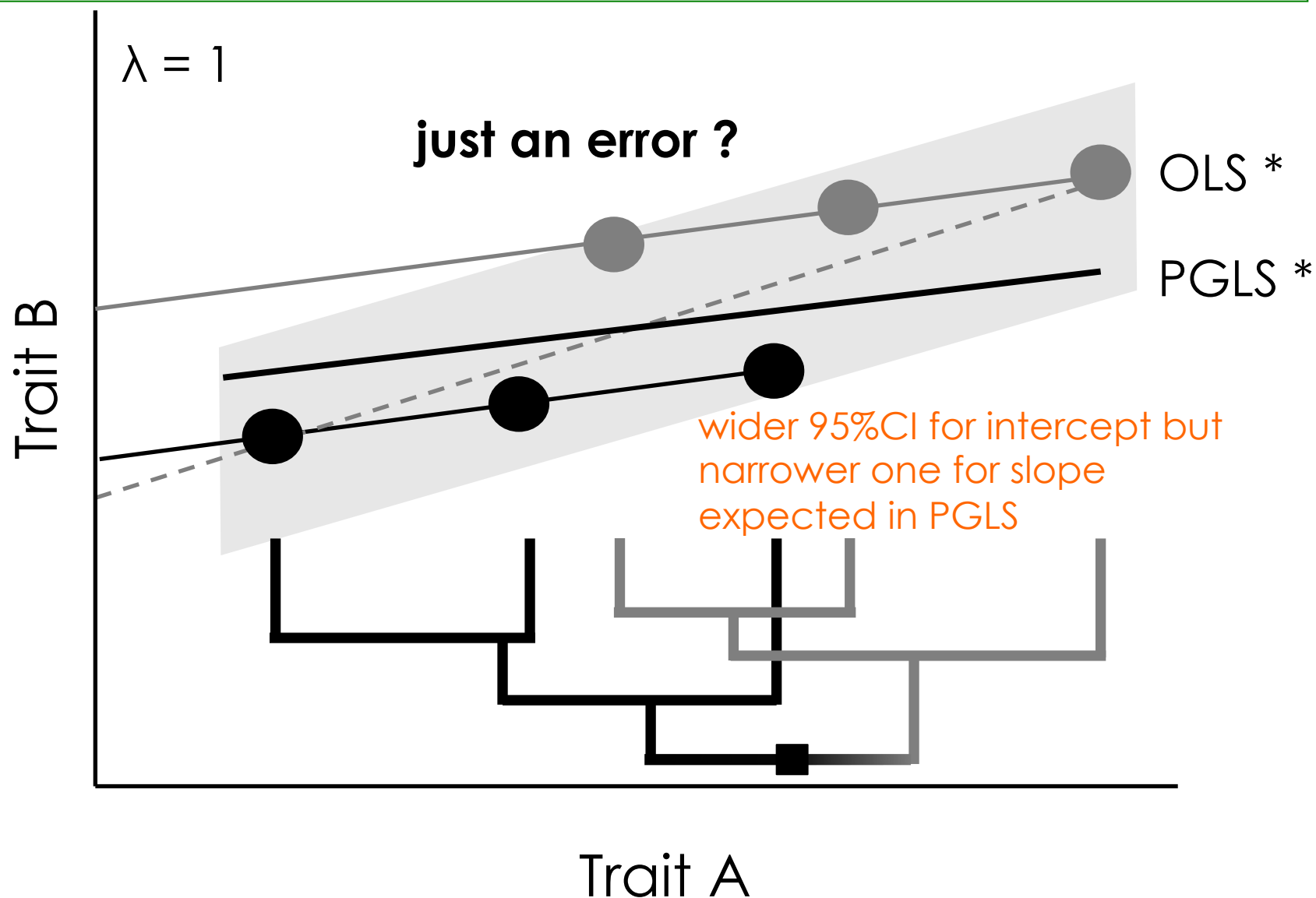


Accounting for phylogeny



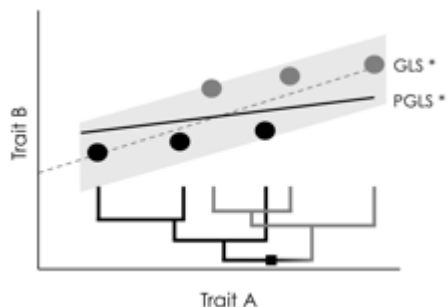


Accounting for phylogeny



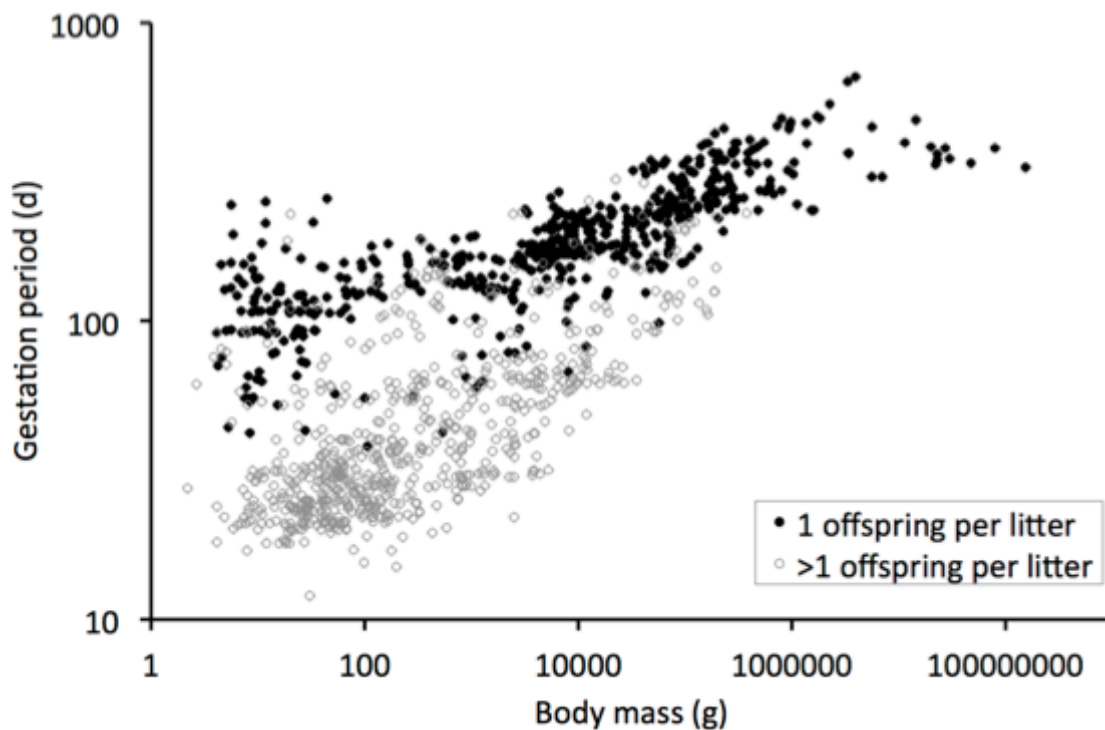


Example V: Gestation time



OLS: 21.5 (19.9-23.3) $BM^{0.19}$ (0.18-0.20)

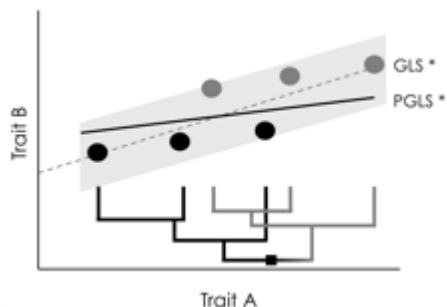
PGLS: 52.4 (41.3-66.3) $BM^{0.09}$ (0.08-0.10)



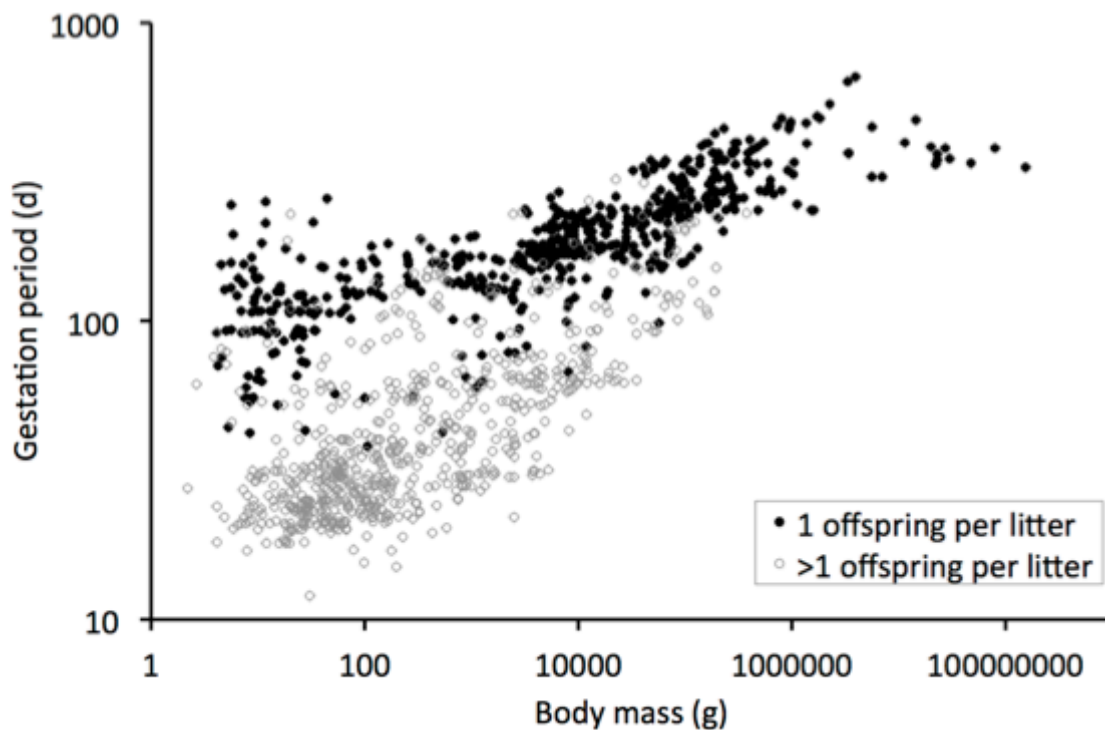
from Clauss et al. (2014)



Example V: Gestation time



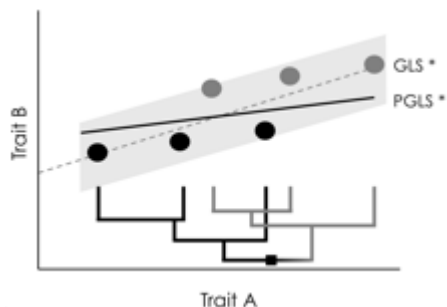
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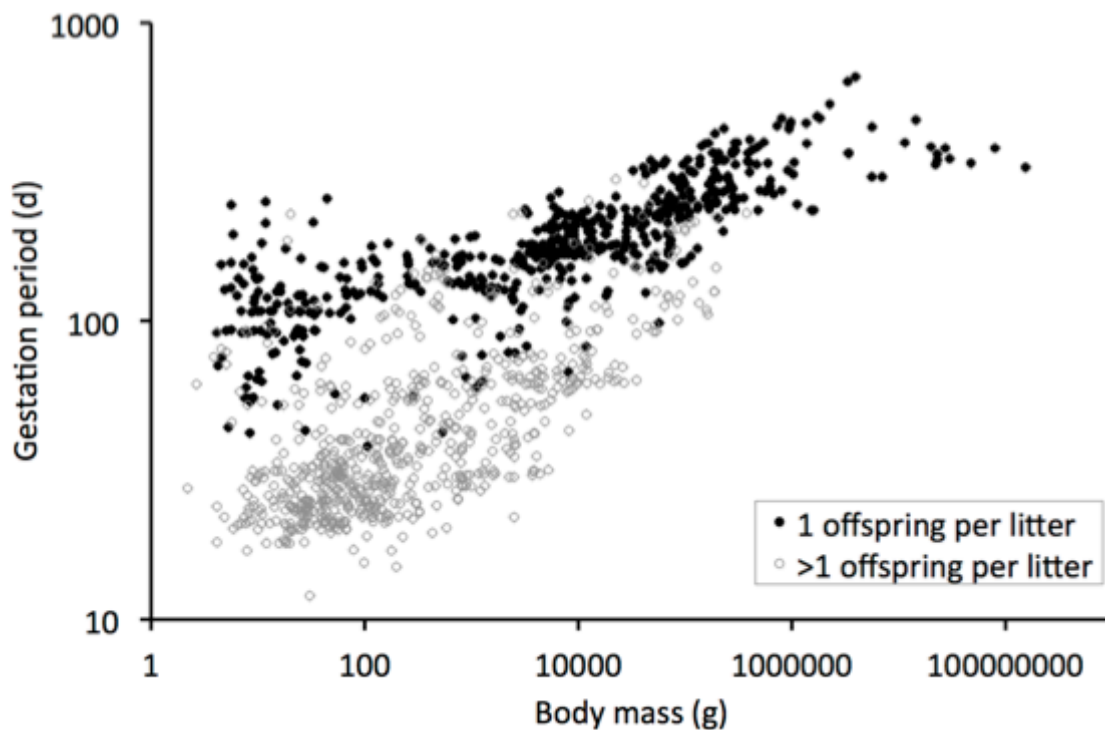
from Clauss et al. (2014)



Example V: Gestation time



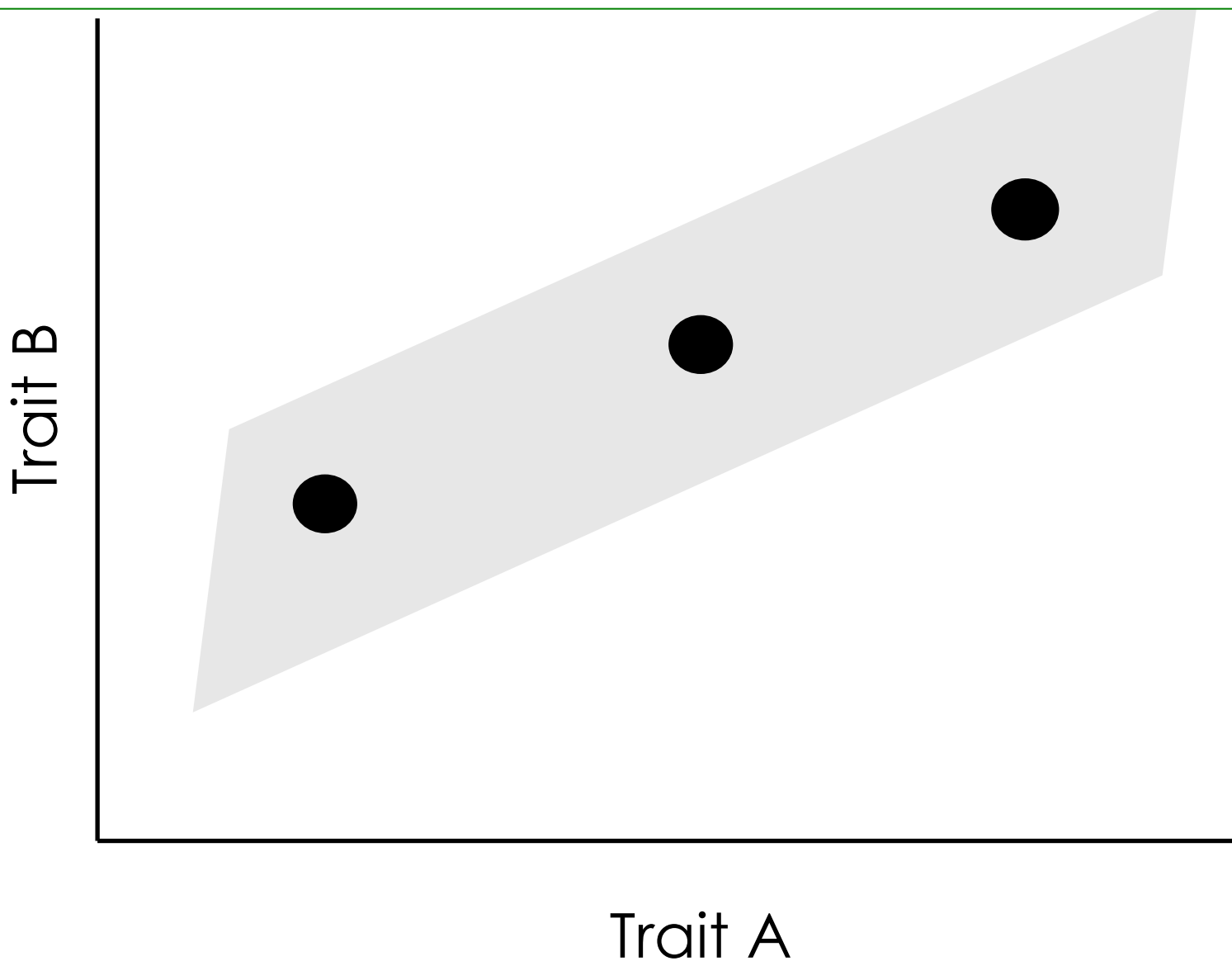
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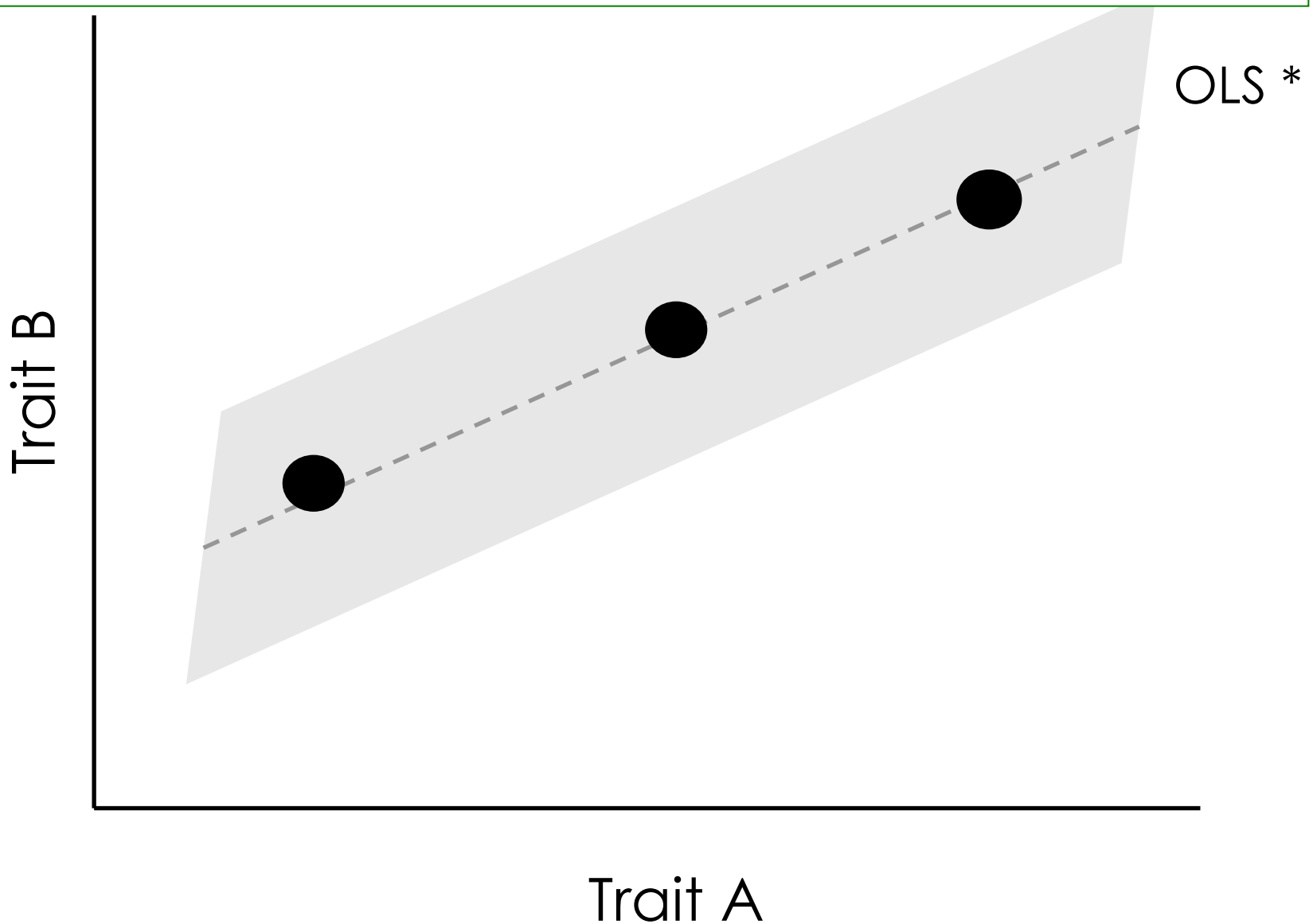
Accounting for phylogeny



from Clauss et al. (2013)



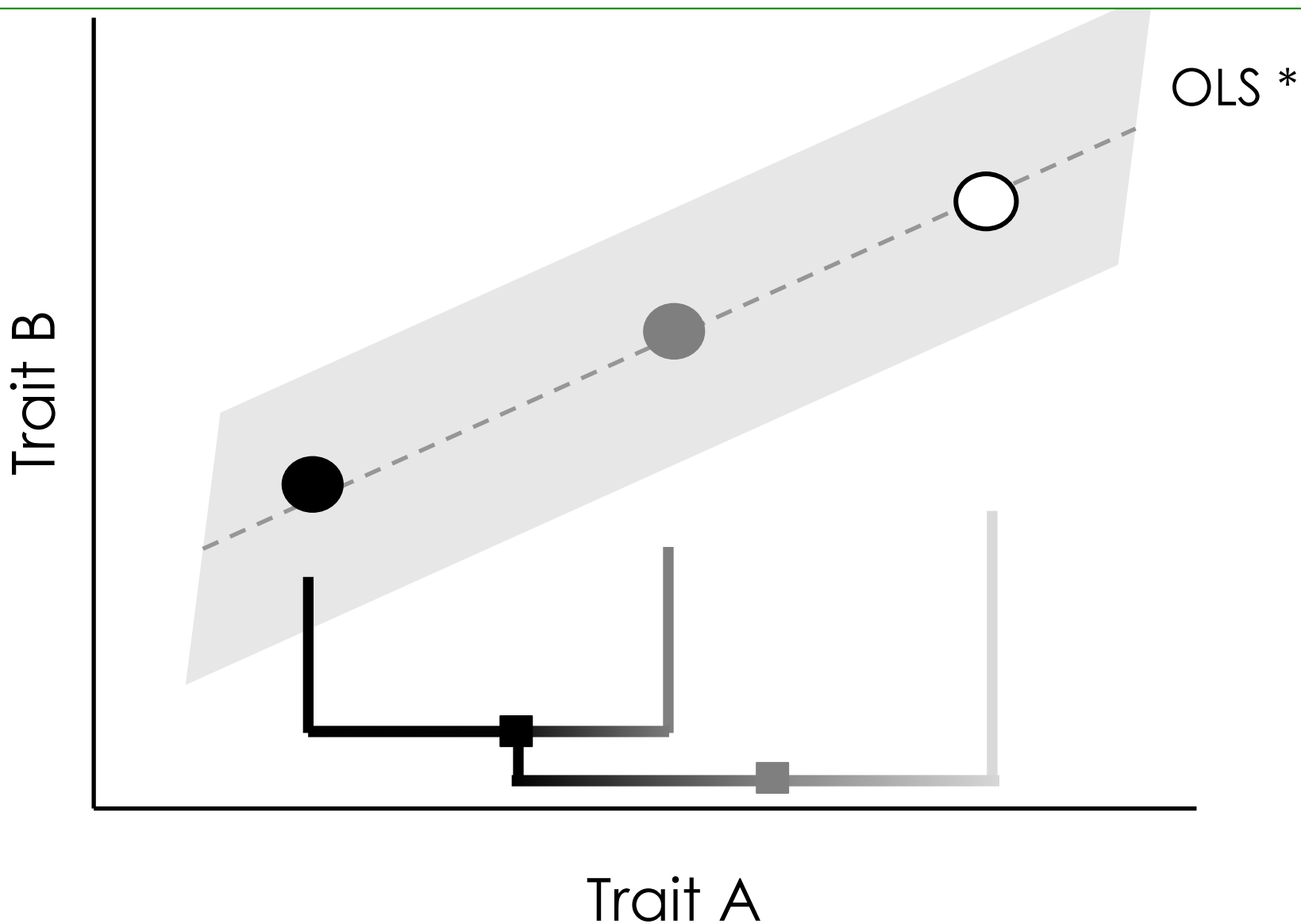
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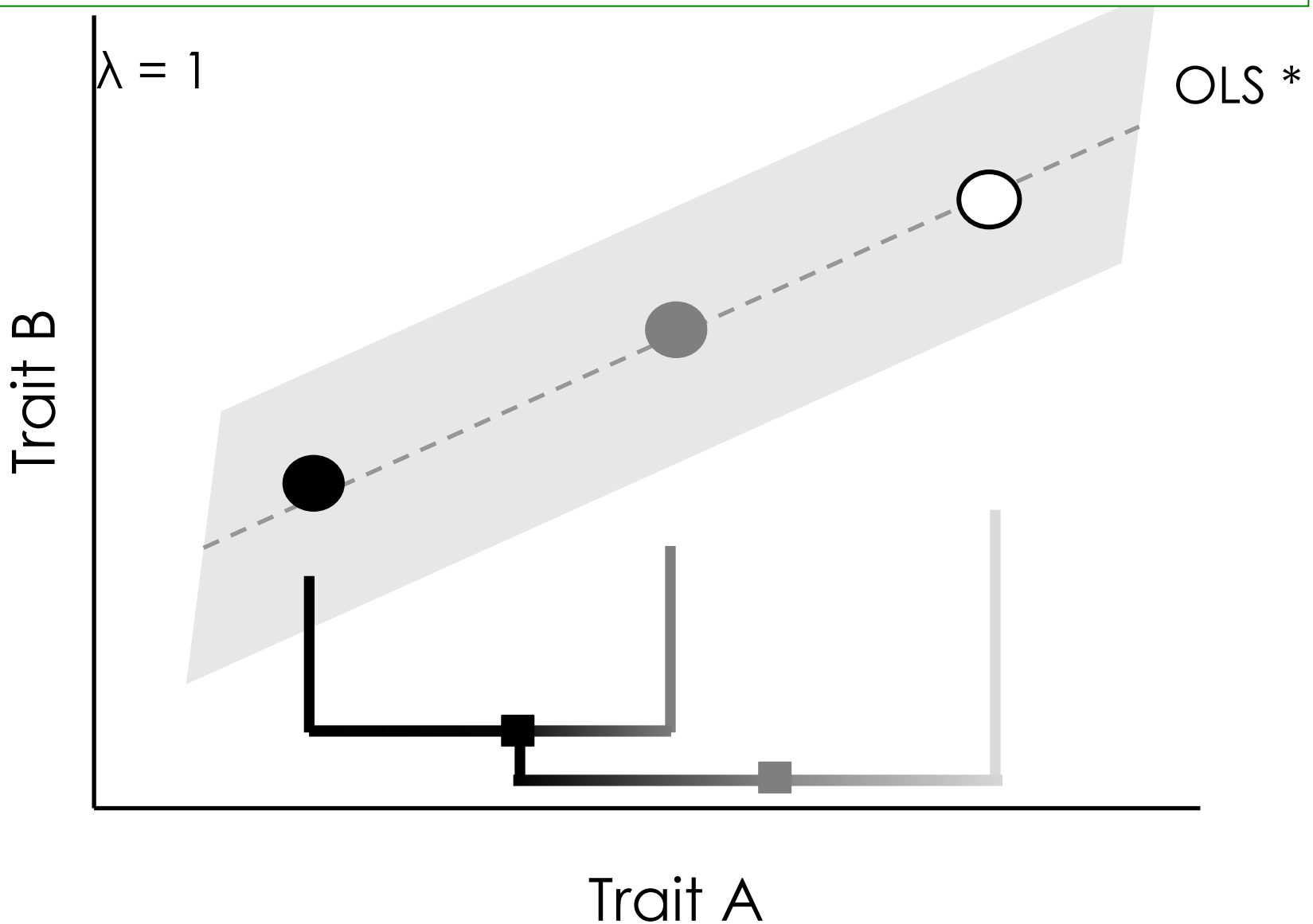
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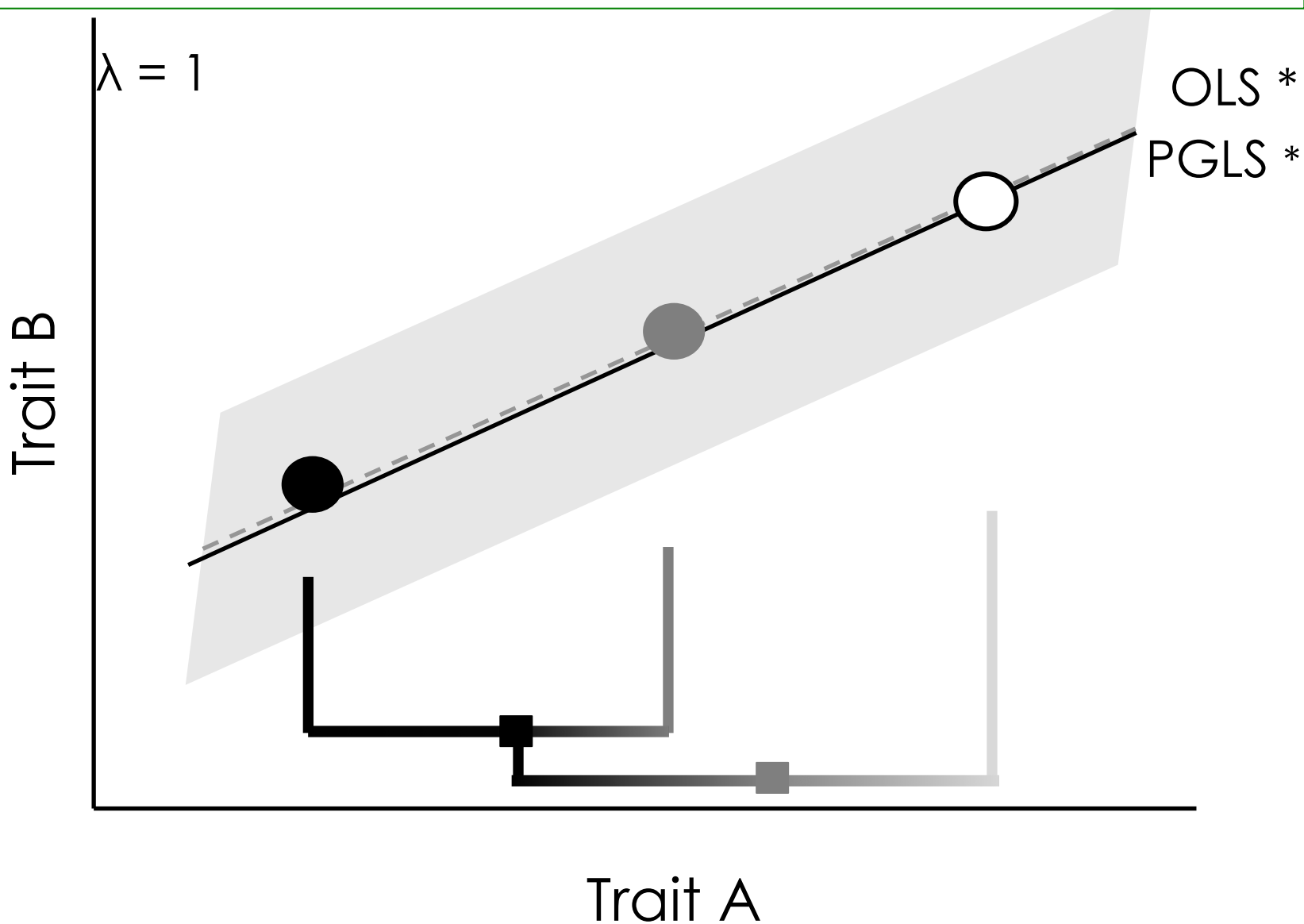
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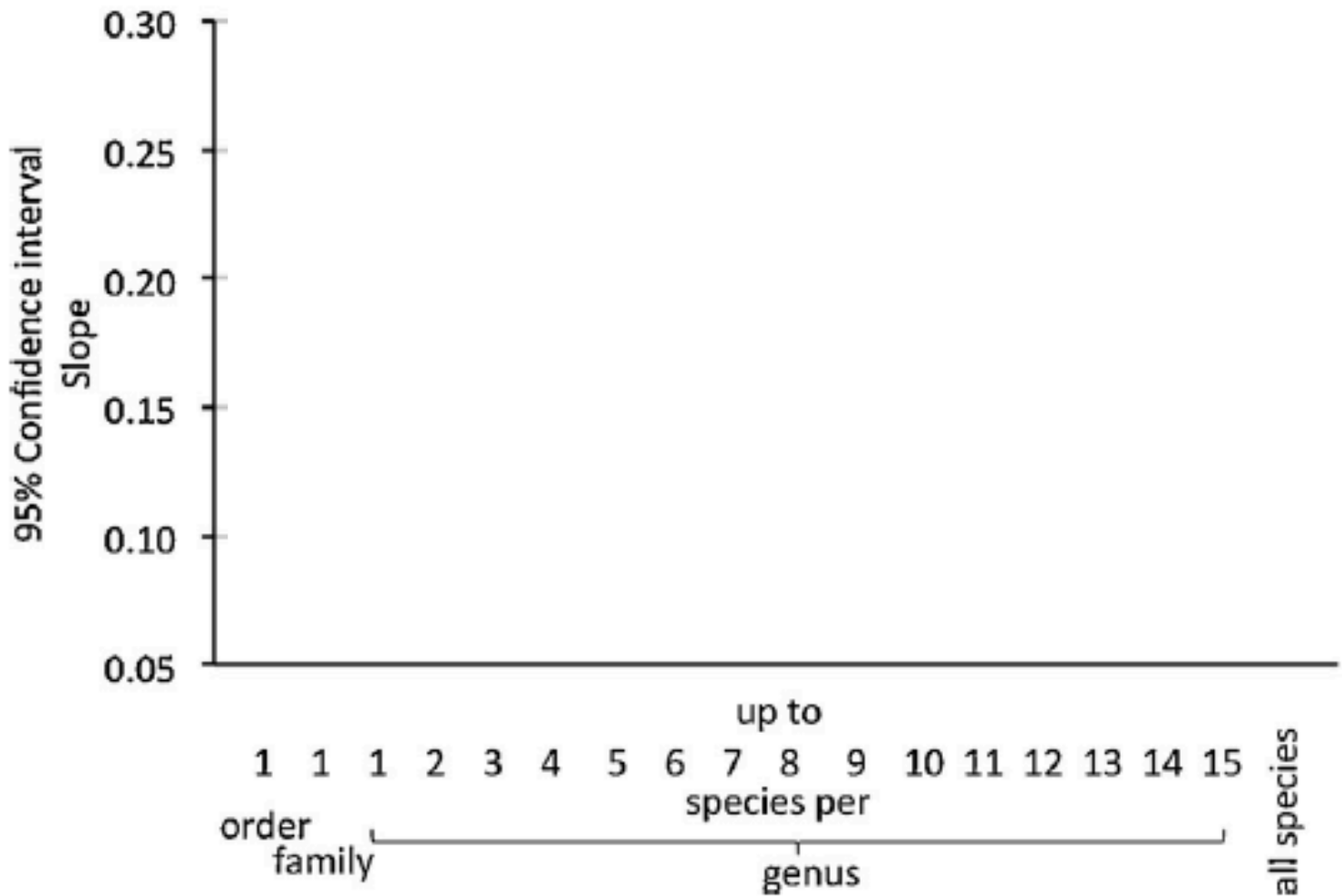
Accounting for phylogeny





Low scaling of a life history variable: Analysing eutherian gestation periods with and without phylogeny-informed statistics

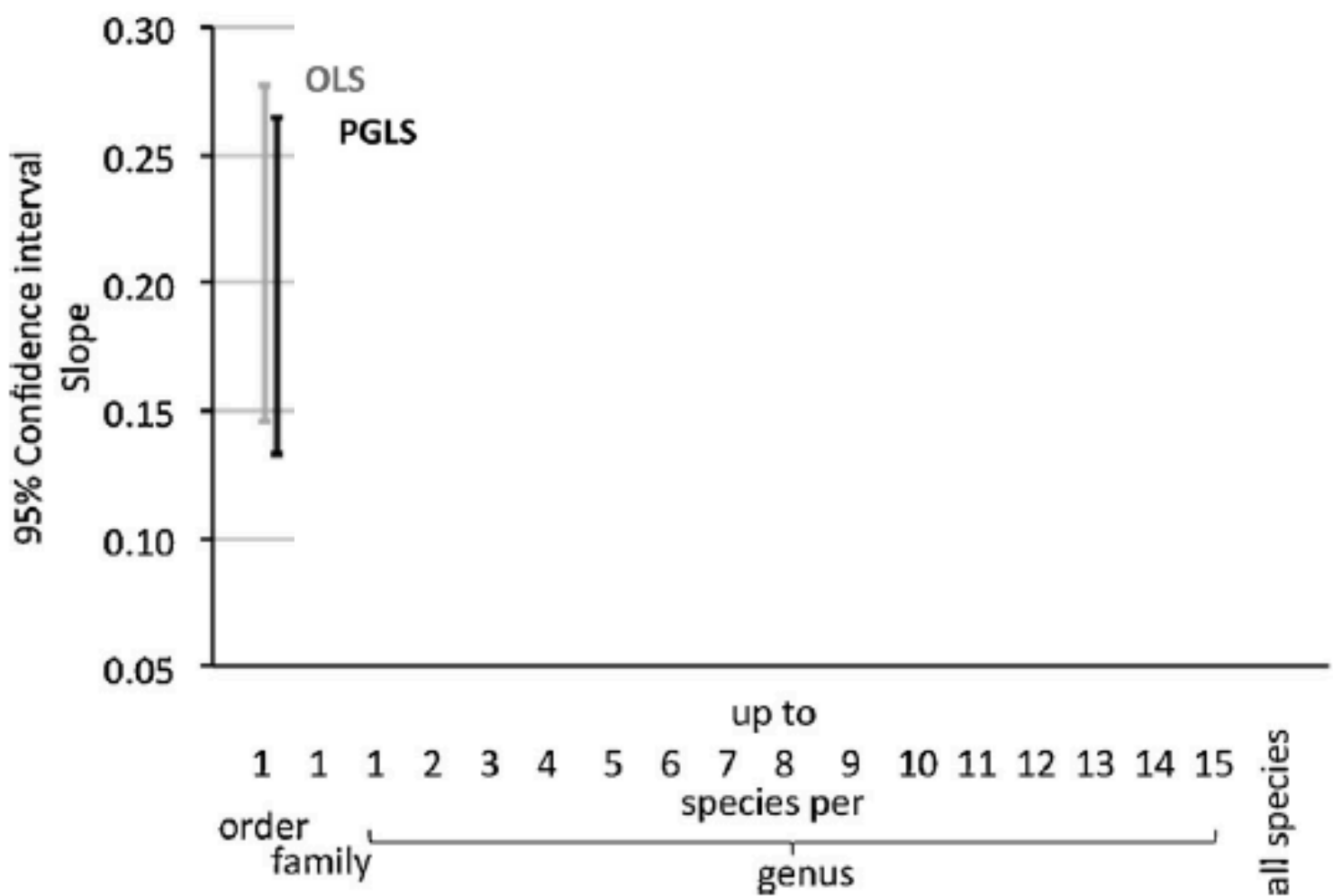
Marcus Clauss^{a,*}, Marie T. Dittmann^b, Dennis W.H. Müller^{a,c}, Philipp Zerbe^{a,d}, Daryl Codron^{a,e}
Mammalian Biology 79 (2014) 9–16





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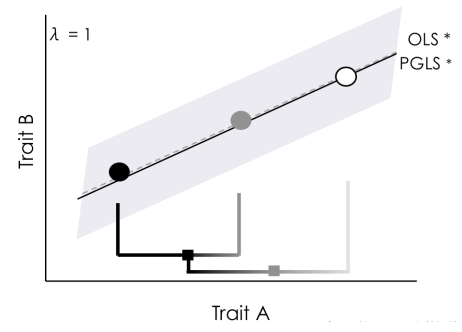
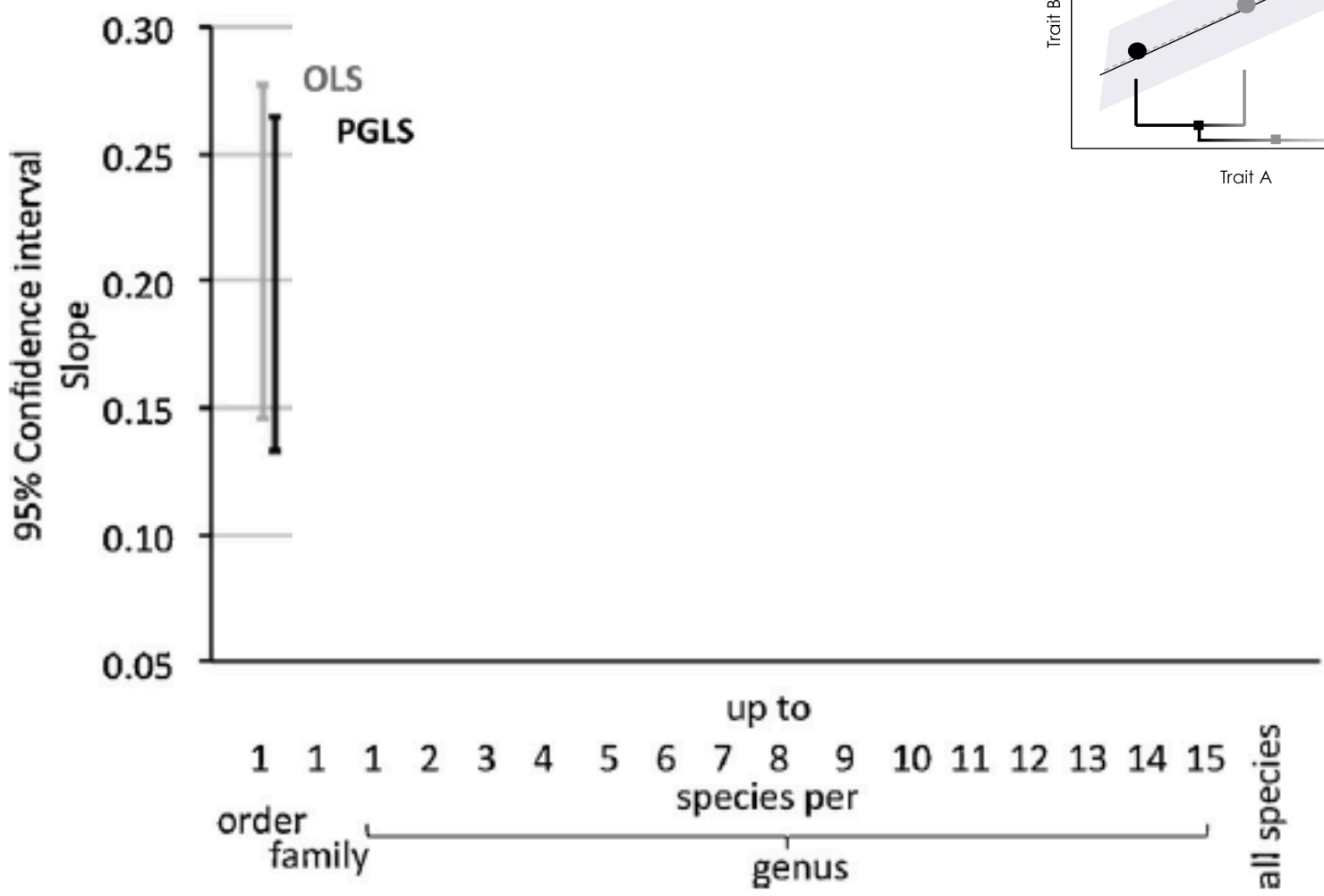




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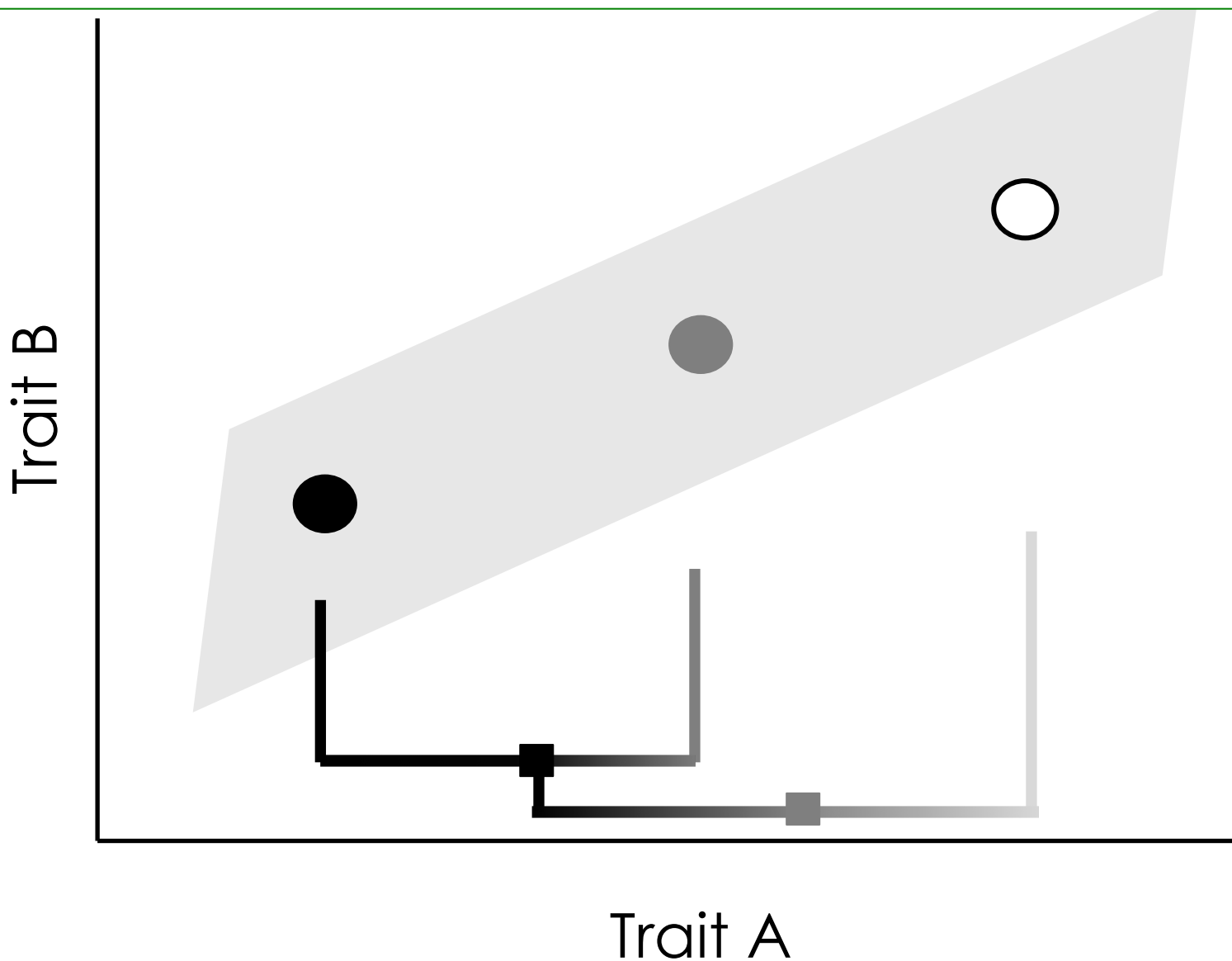
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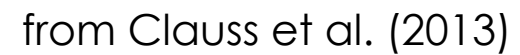




Accounting for phylogeny

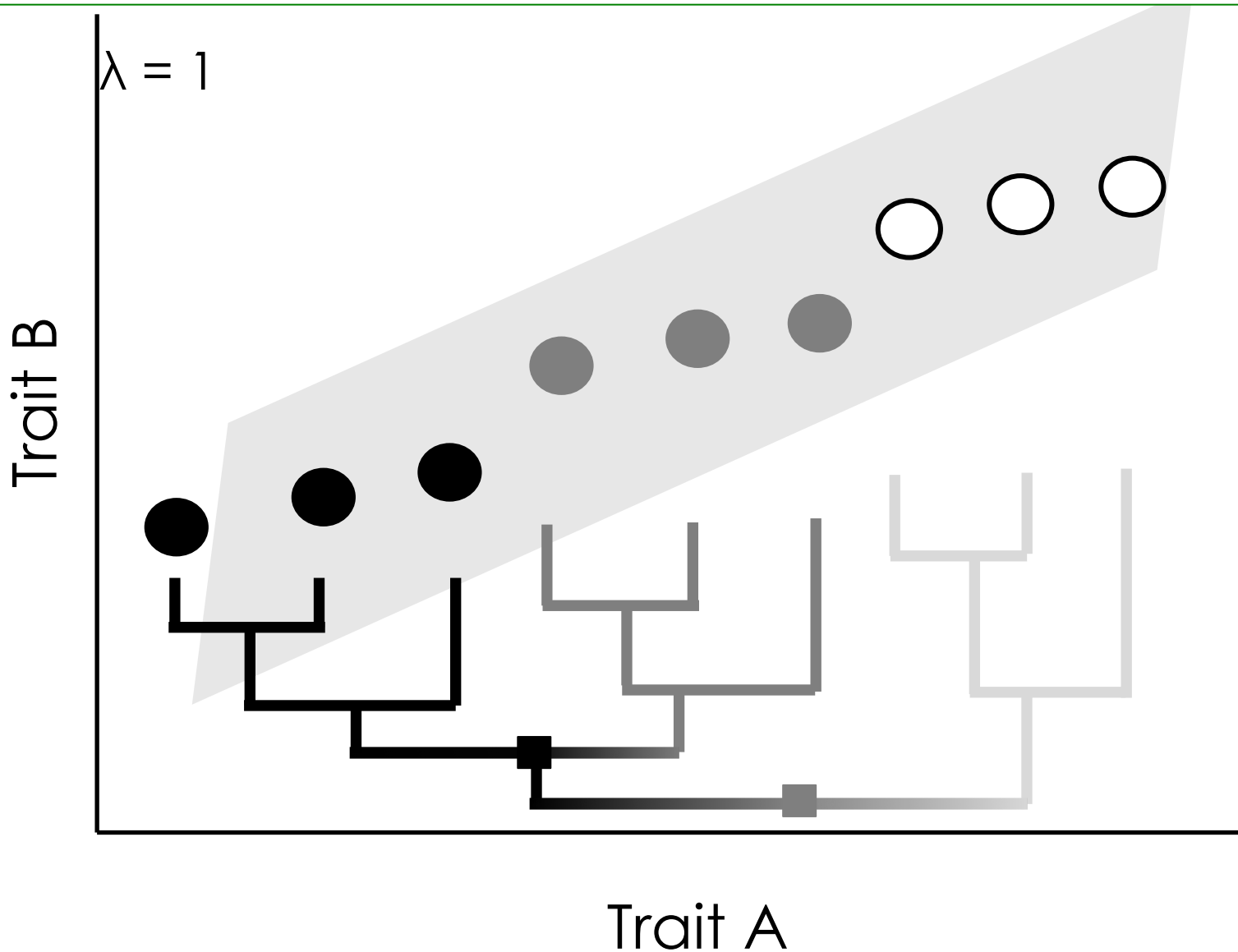


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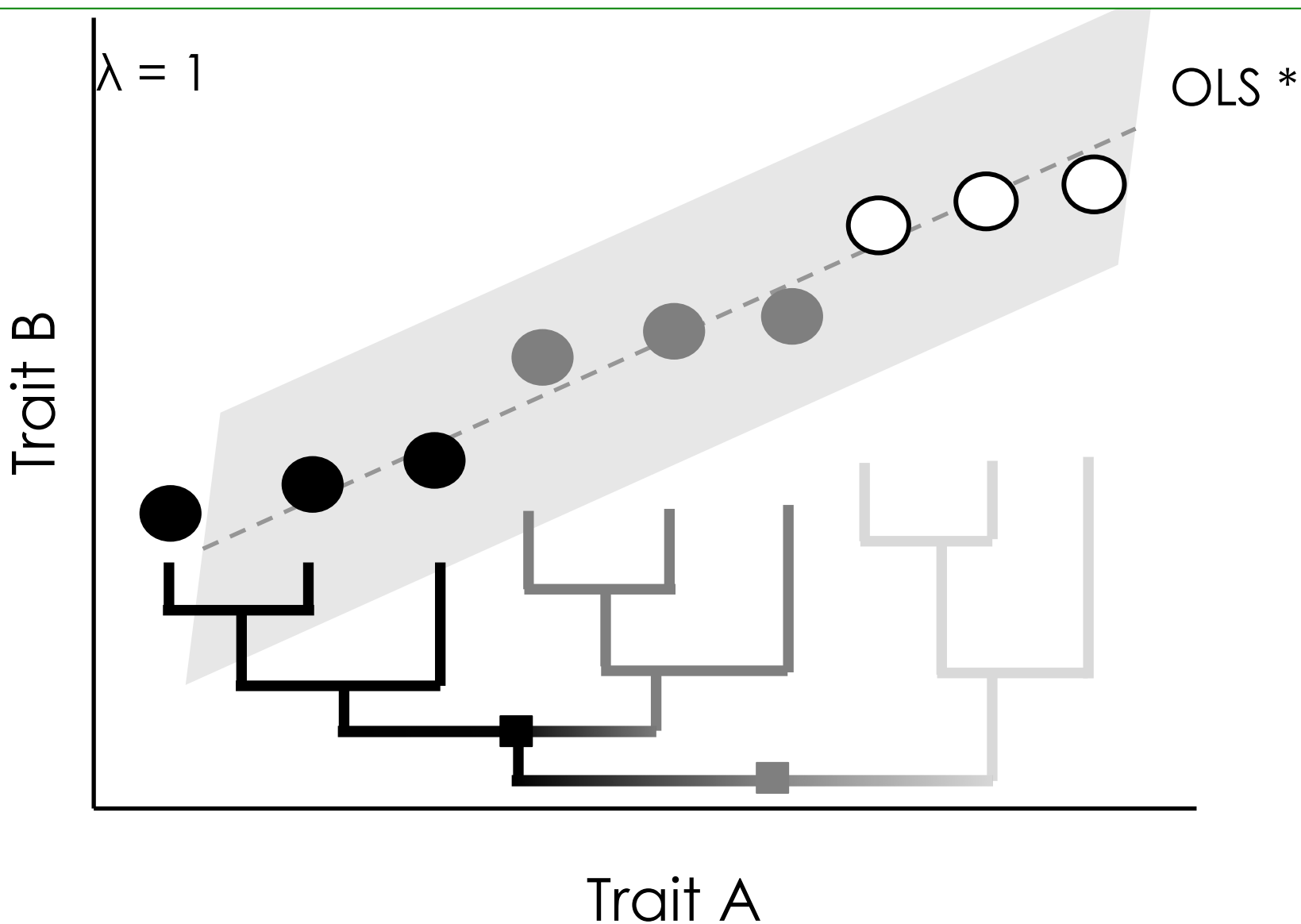
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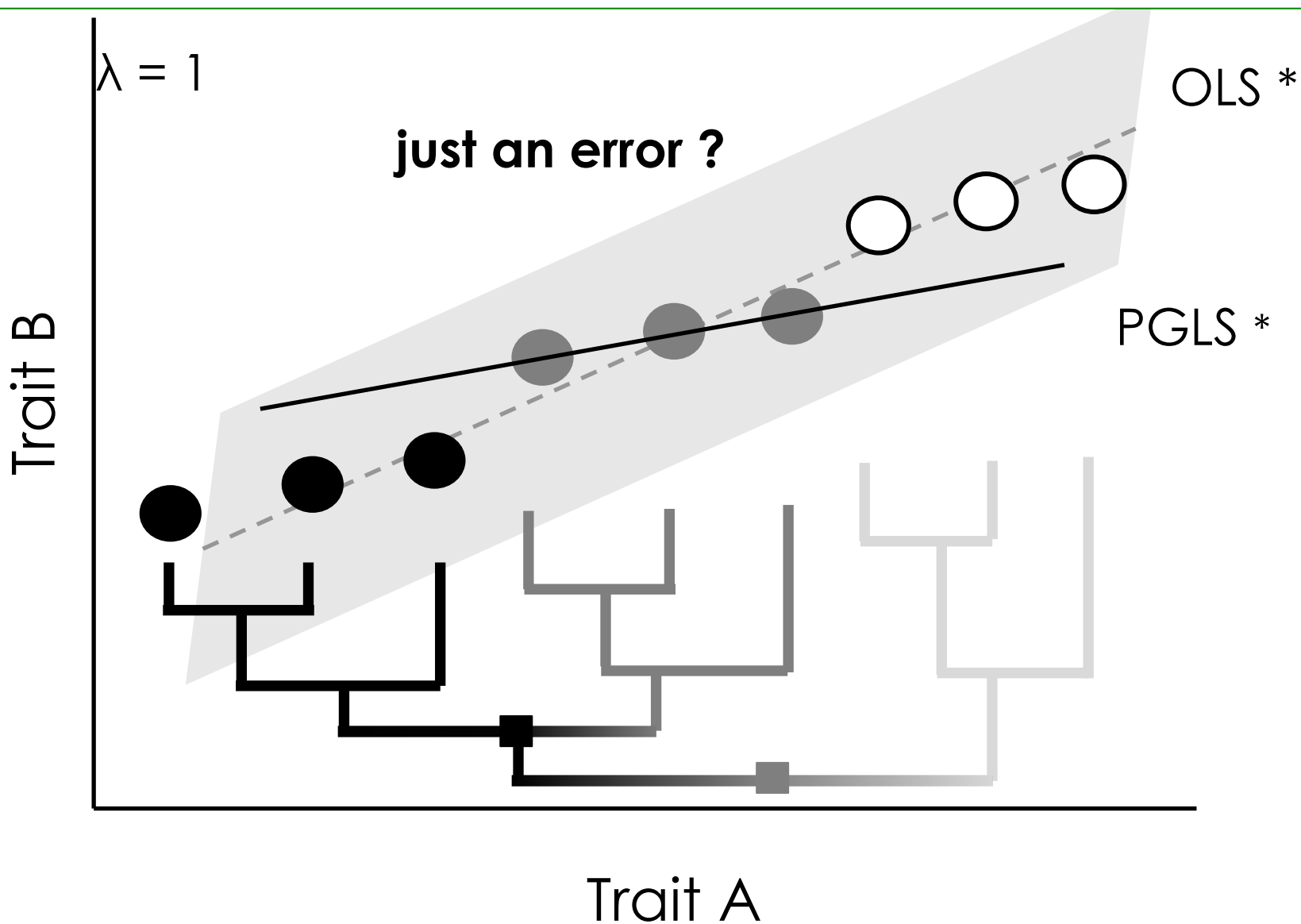


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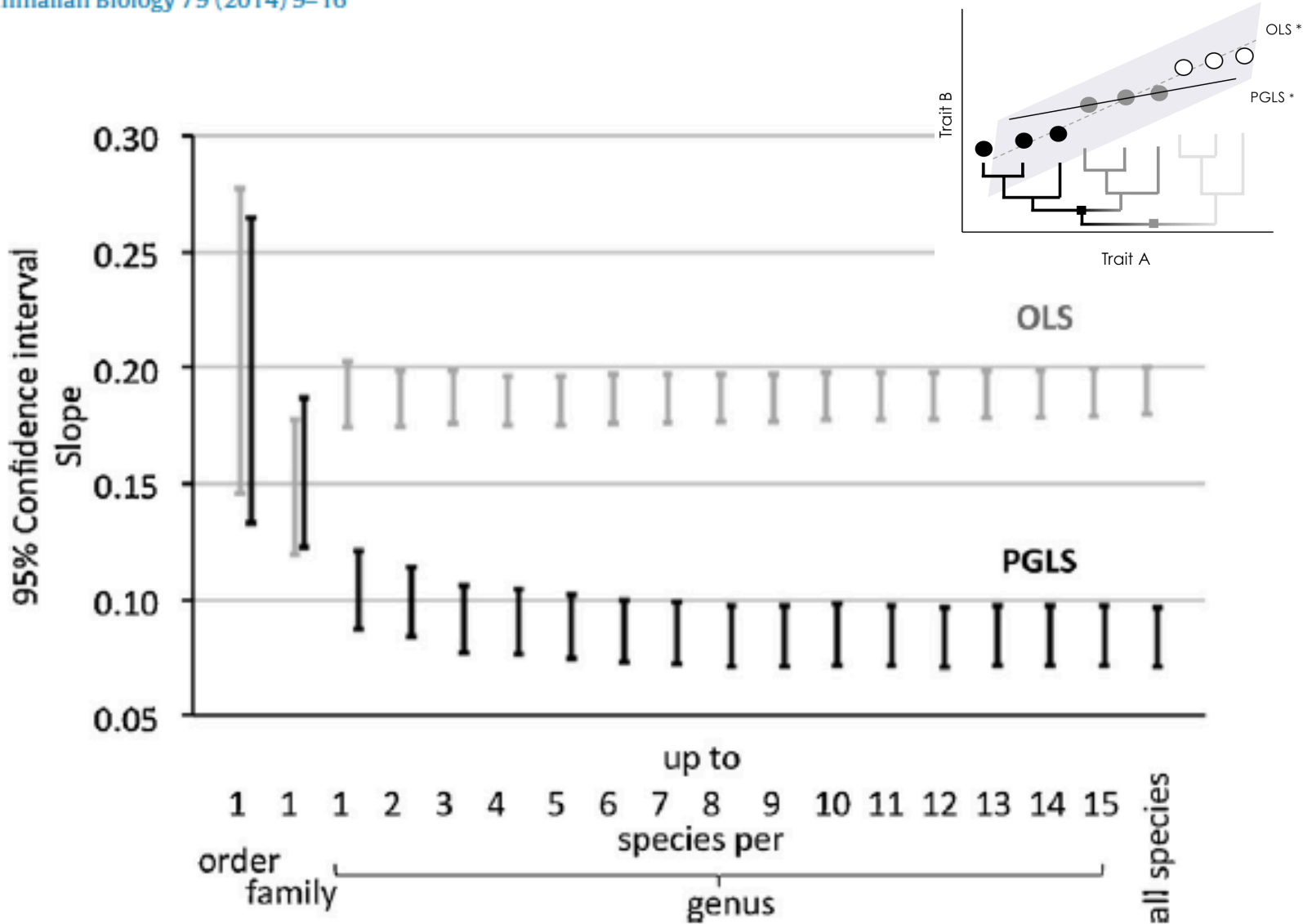




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A deadly sin ?

REVIEW

The seven deadly sins of comparative analysis

R. P. FRECKLETON

J. EVOL. BIOL. **22** (2009) 1367–1375

Reporting both PI and PC analyses

Frequently, both across-species and phylogenetically corrected analyses of the same data are reported simultaneously. This is despite the fact that the two forms of analysis make very different assumptions about the distribution of the data.



A deadly sin ? – **No !**

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~~Frequently, both across-species and phylogenetically corrected analyses of the same data are reported simultaneously. This is despite the fact that the two forms of analysis make very different assumptions about the distribution of the data.~~

A comparison of OLS and PGLS results is an important tool for understanding the structure of the data! (irrespective of which is the ‘correct’ one) **because the two make very different assumptions about the data**



Directionality in Evolution: Allometries as snapshots in evolutionary time

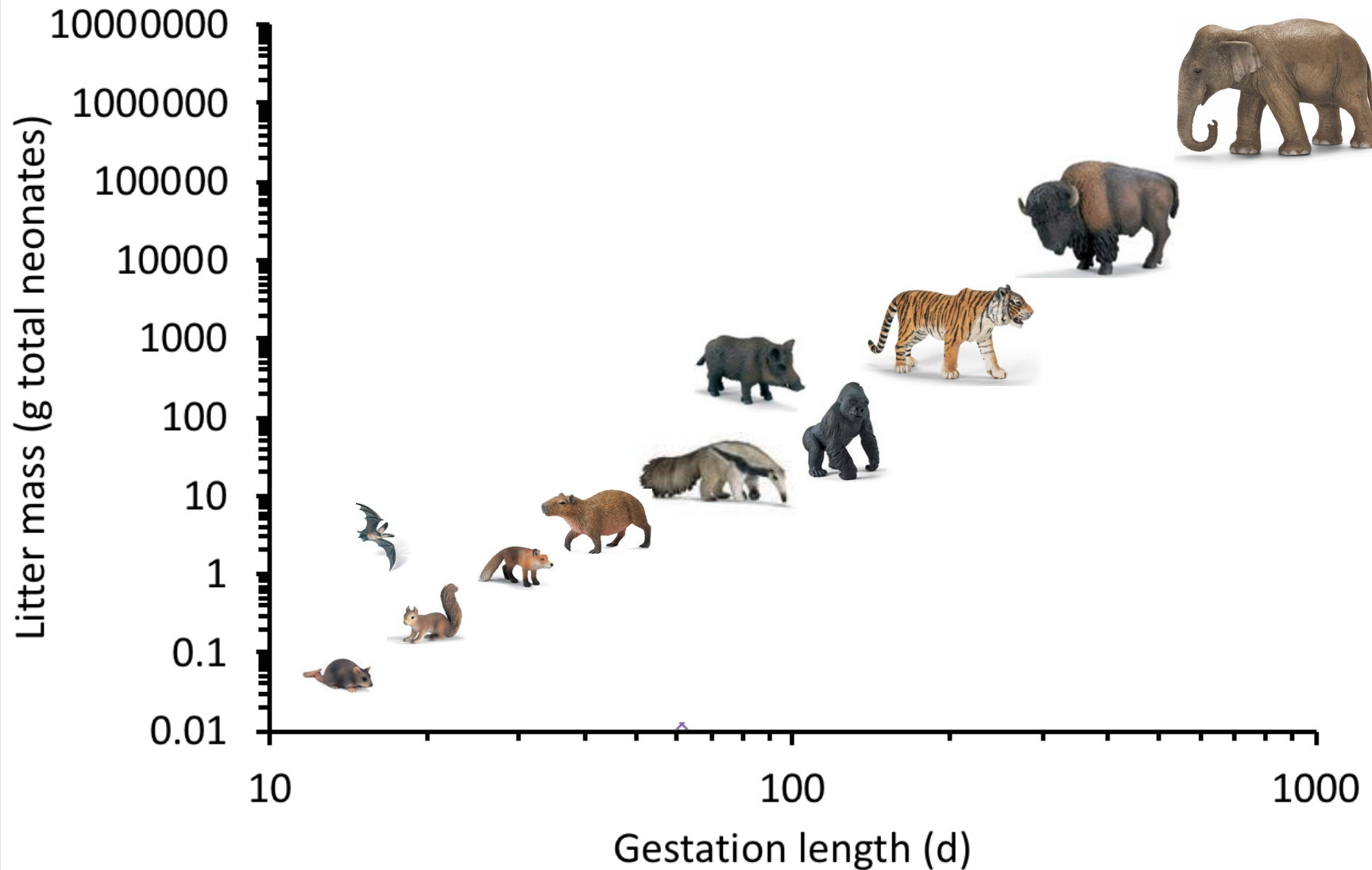


Directionality in Evolution: beware of the *natural fallacy*



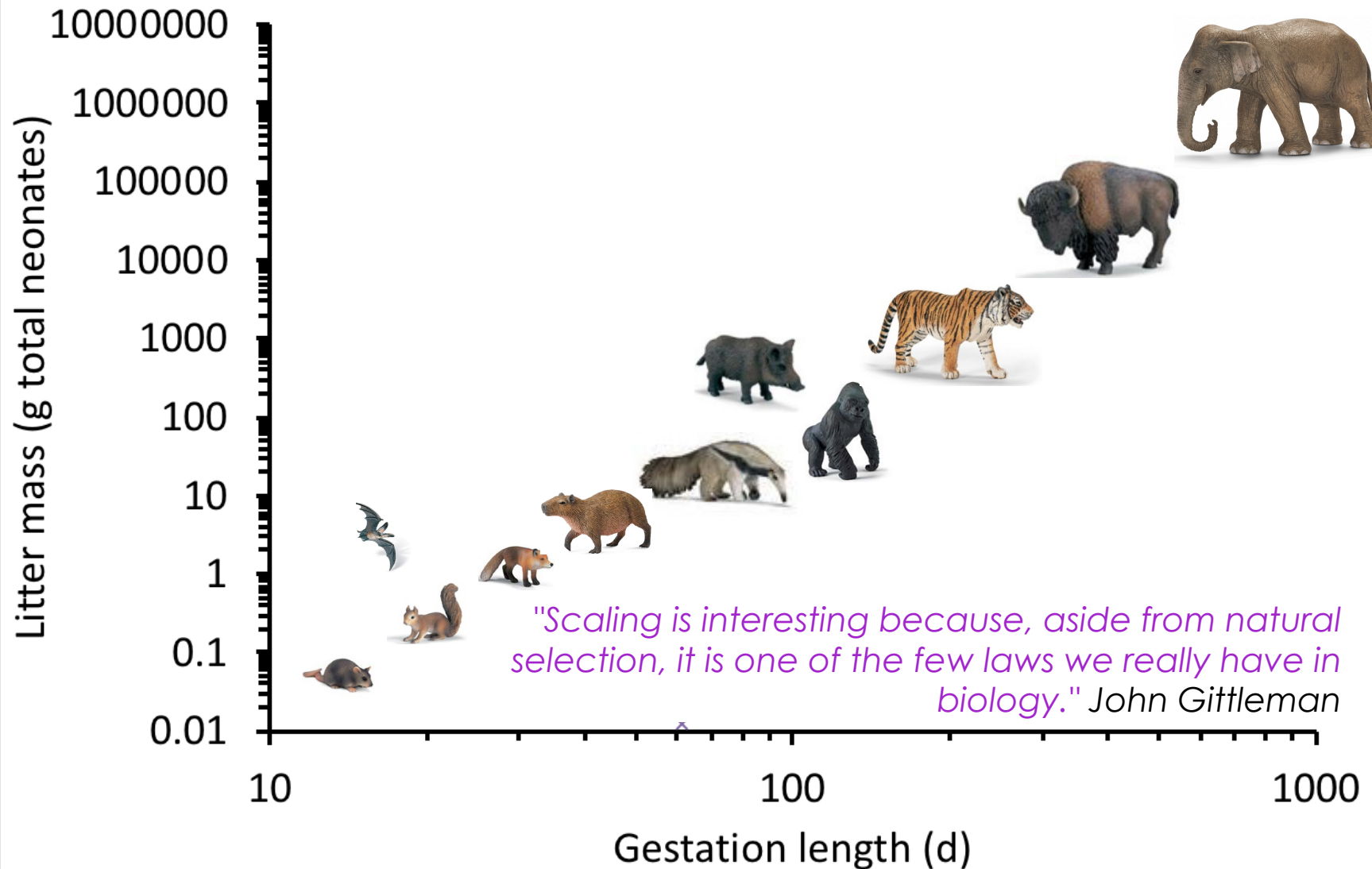


Biology: fixed laws ?



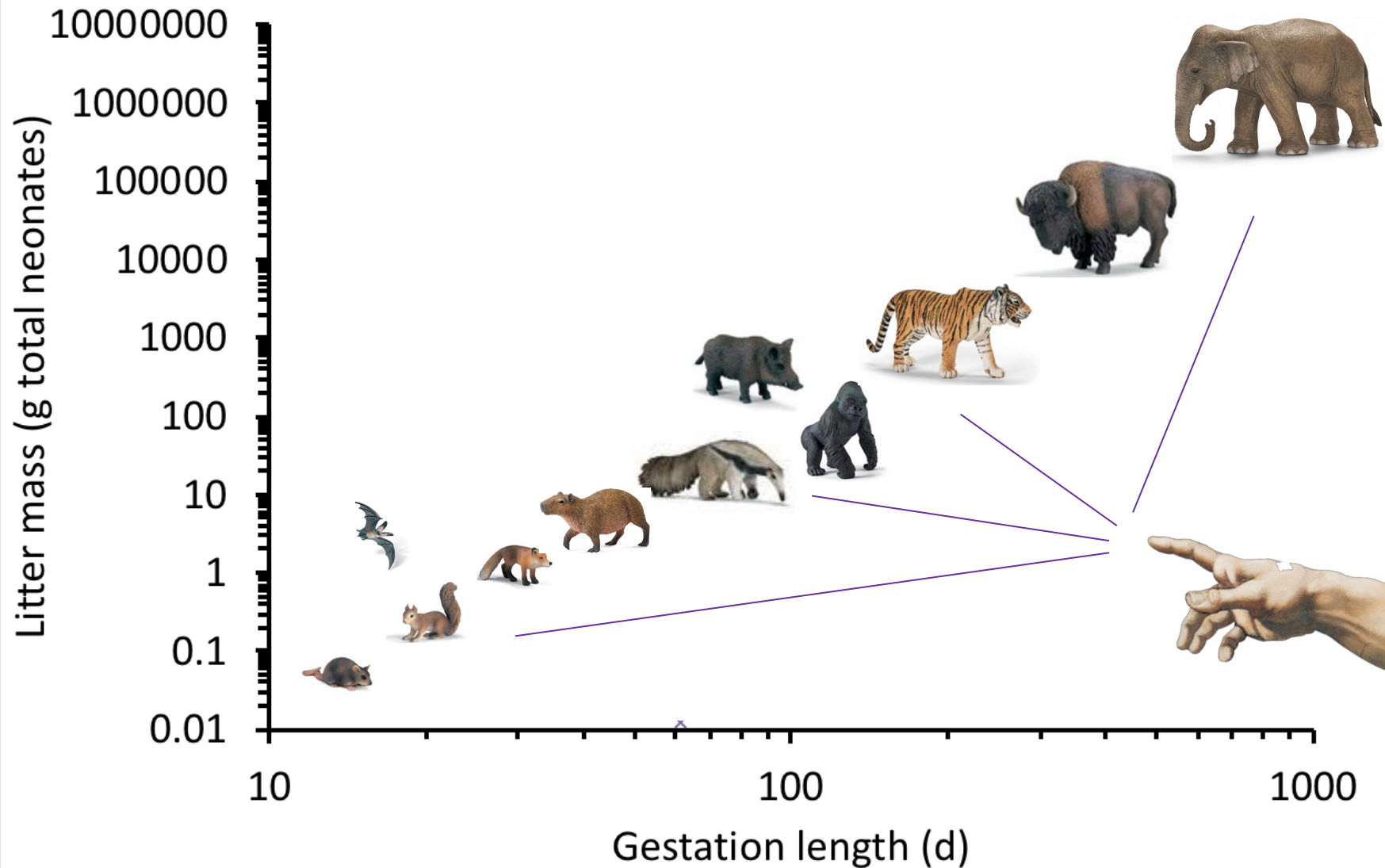


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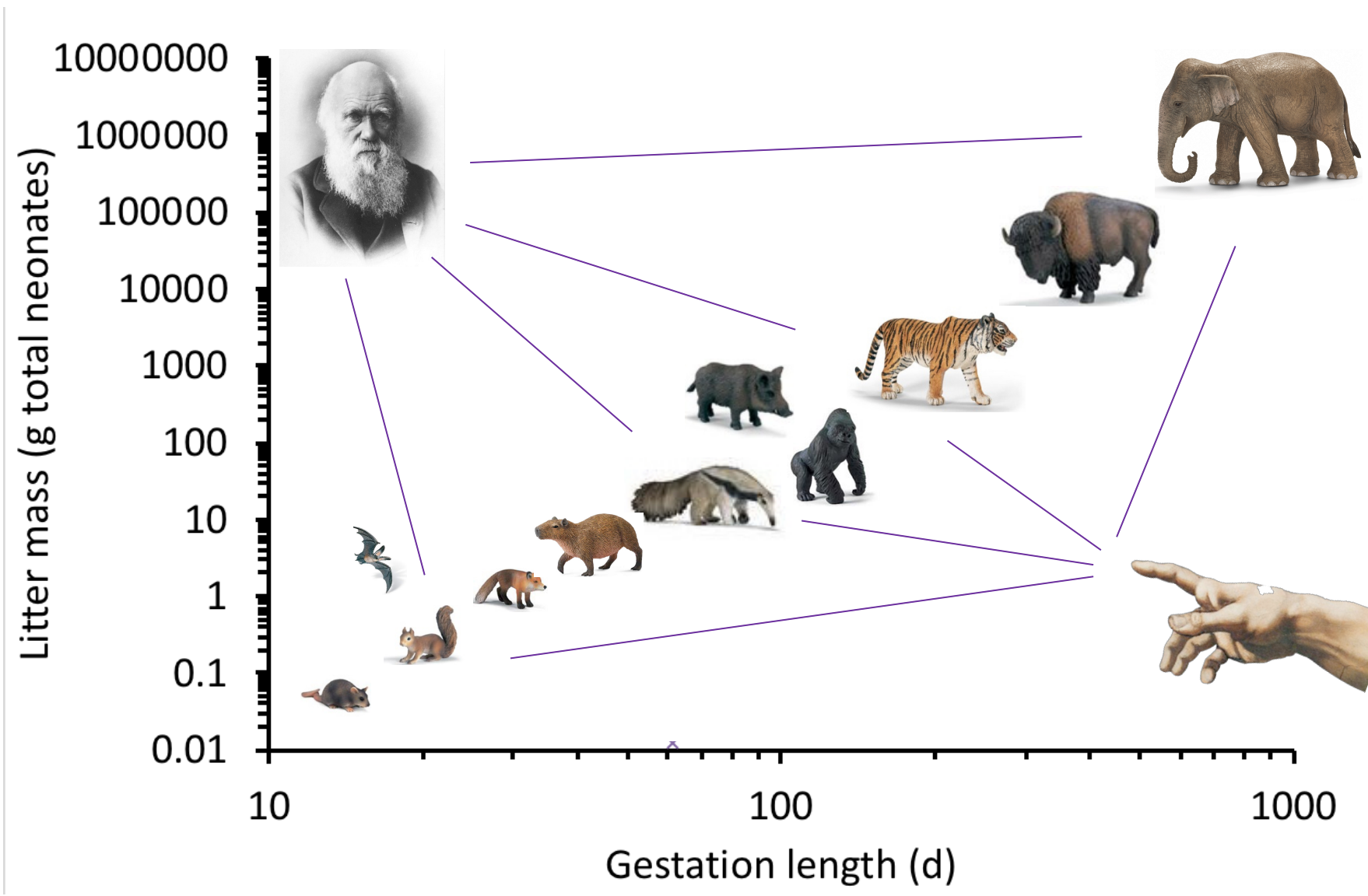


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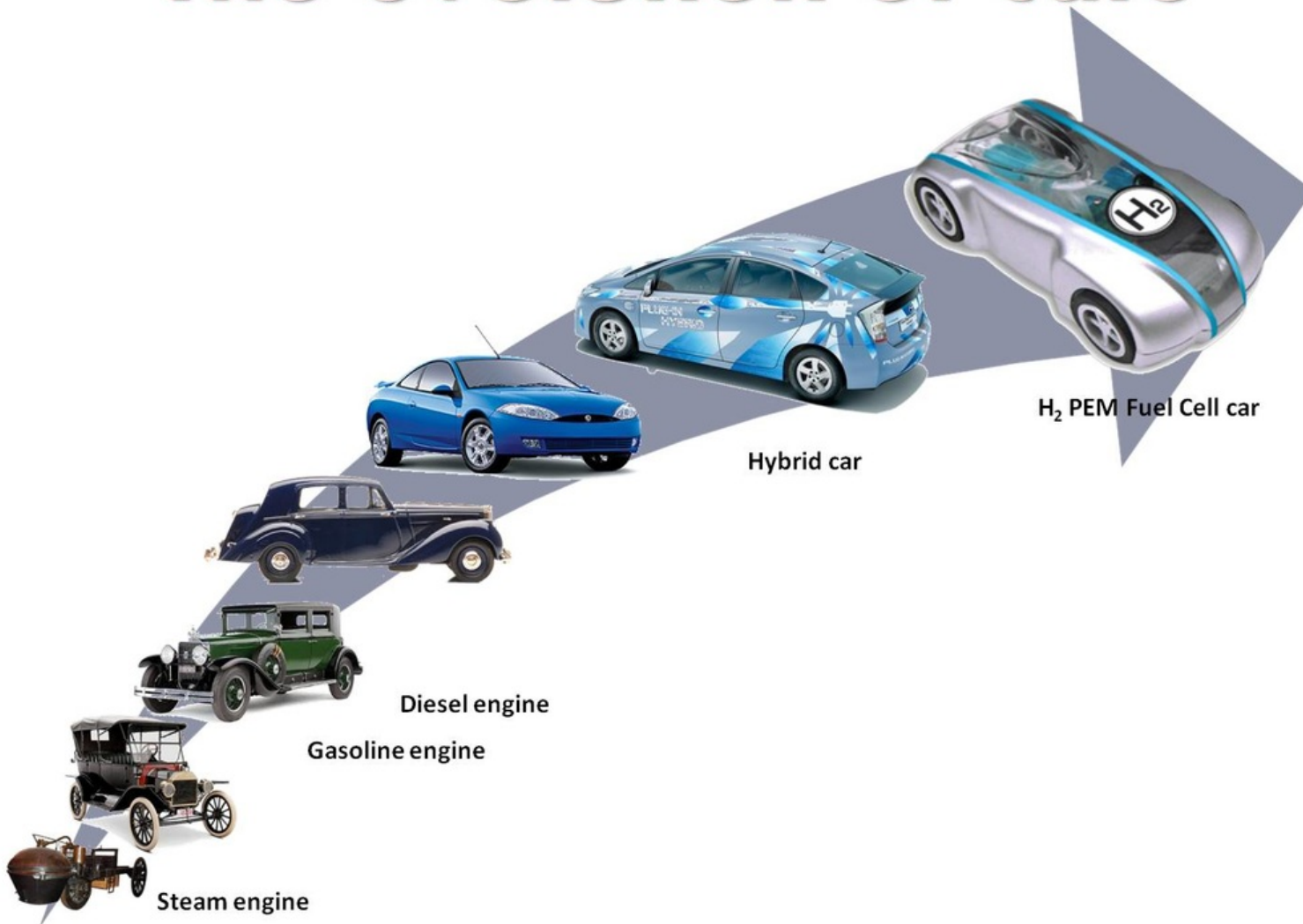


Biology: fixed laws ?





The evolution of cars



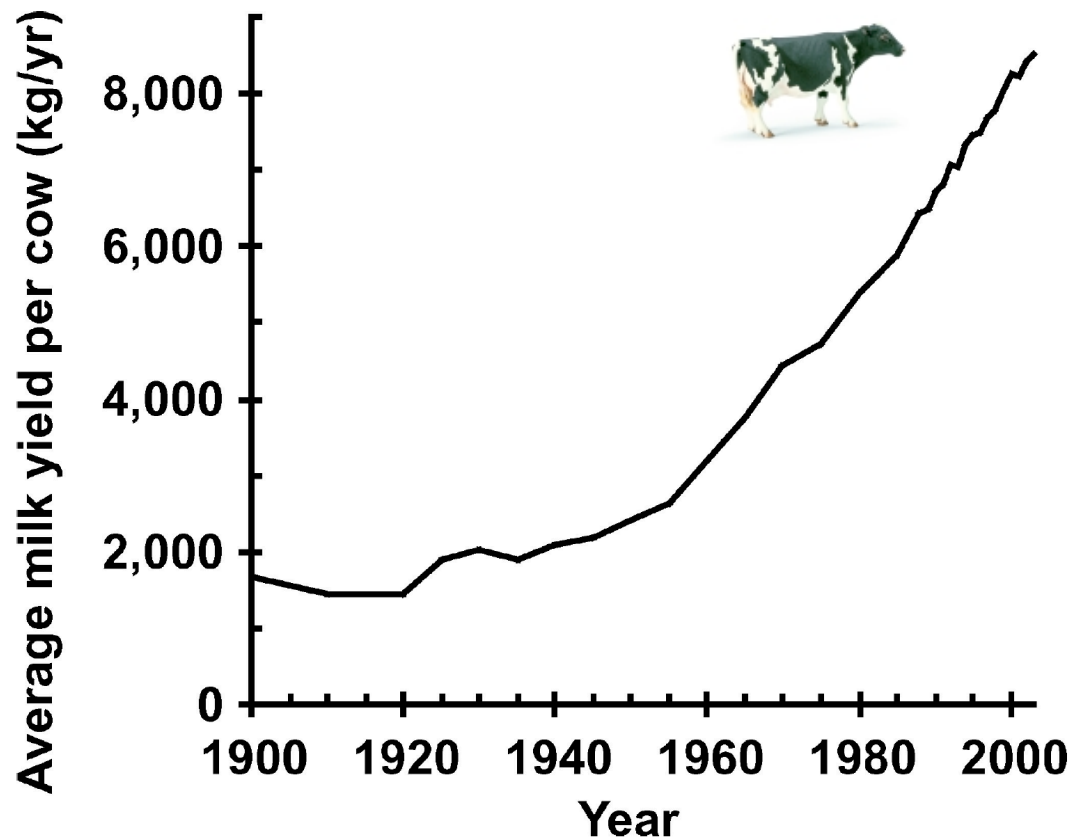


J. Dairy Sci. 89:1280–1291

© American Dairy Science Association, 2006.

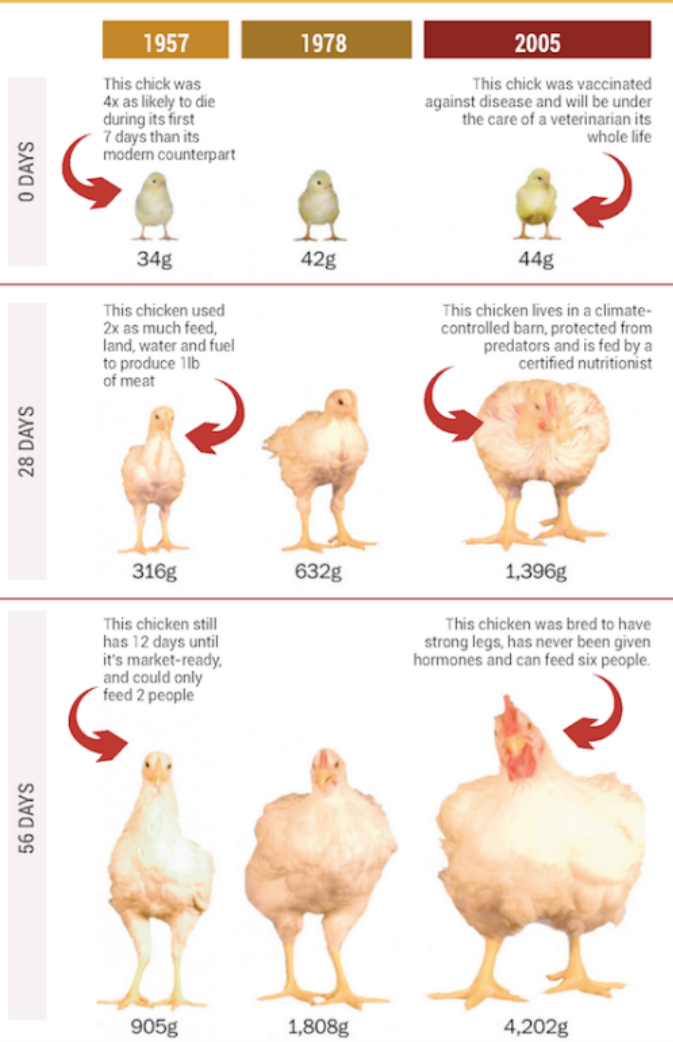
Major Advances in Nutrition: Relevance to the Sustainability of the Dairy Industry

M. J. VandeHaar*¹ and N. St-Pierret†





YEP, CHICKENS ARE BIGGER TODAY



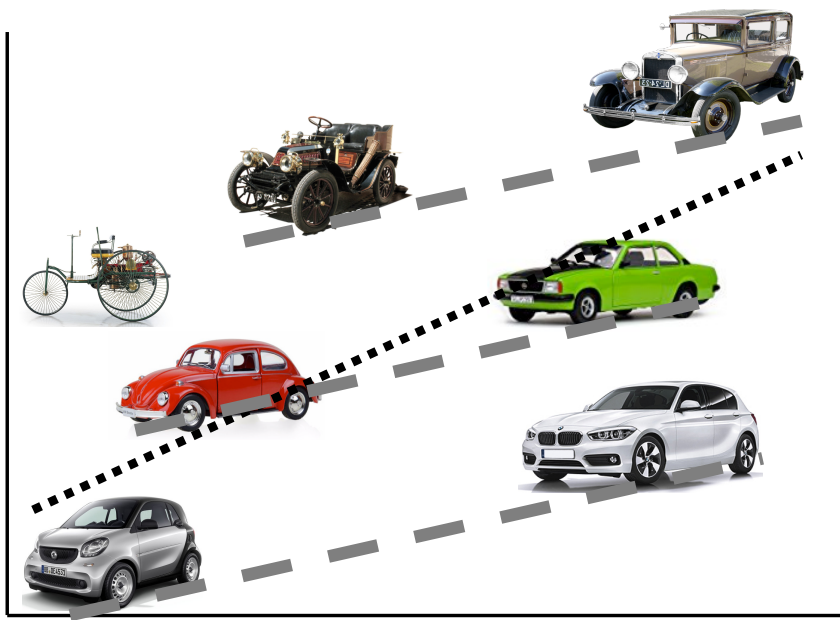
It's no secret that today's chickens are bigger than in years past. They're also the healthiest they've ever been. Find out how at chickencheck.in



Note: 1,000 grams equals 2.2 pounds
Source: University of Alabama Meat Control
Image Credit: <https://www.washingtonpost.com/news/energy/wp/2015/07/04/the-unbelievable-growth-of-american-food-bodies-faster-and-easier-visualized/>

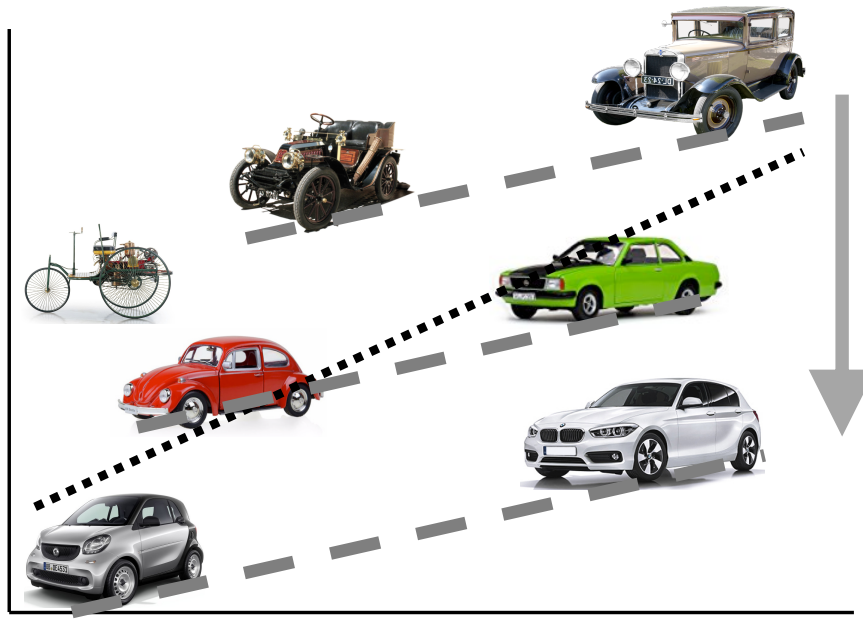


Energy per km





Energy per km

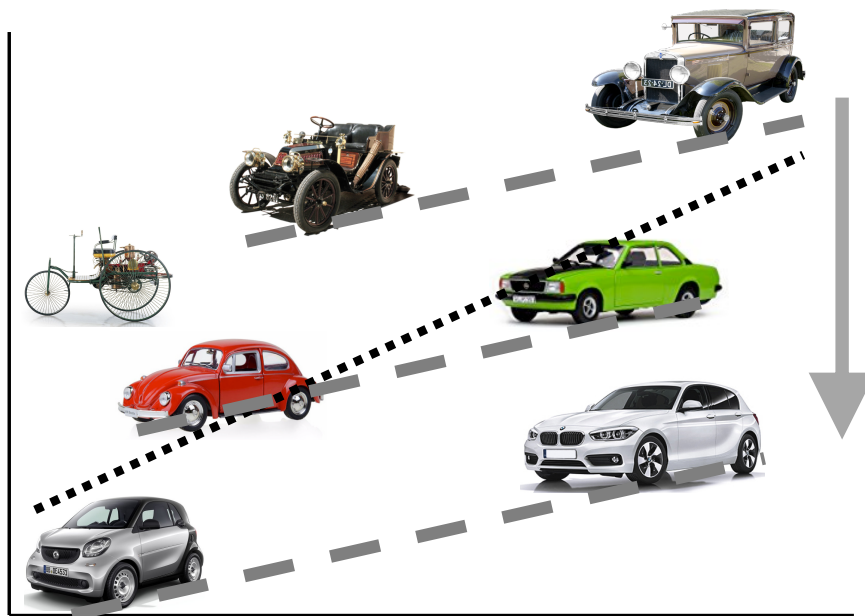


Mass

You would not consider the overall pattern a fixed law, but consider it with respect to technical progress.



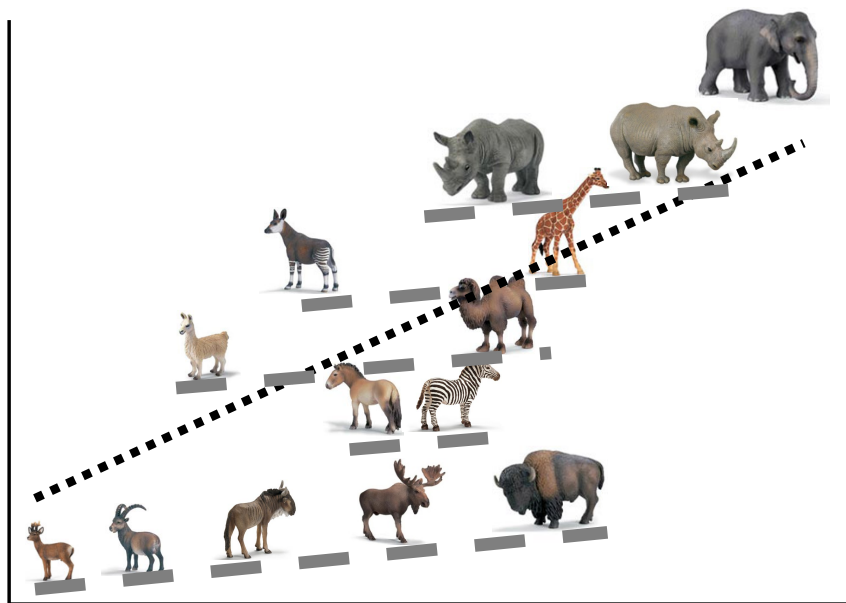
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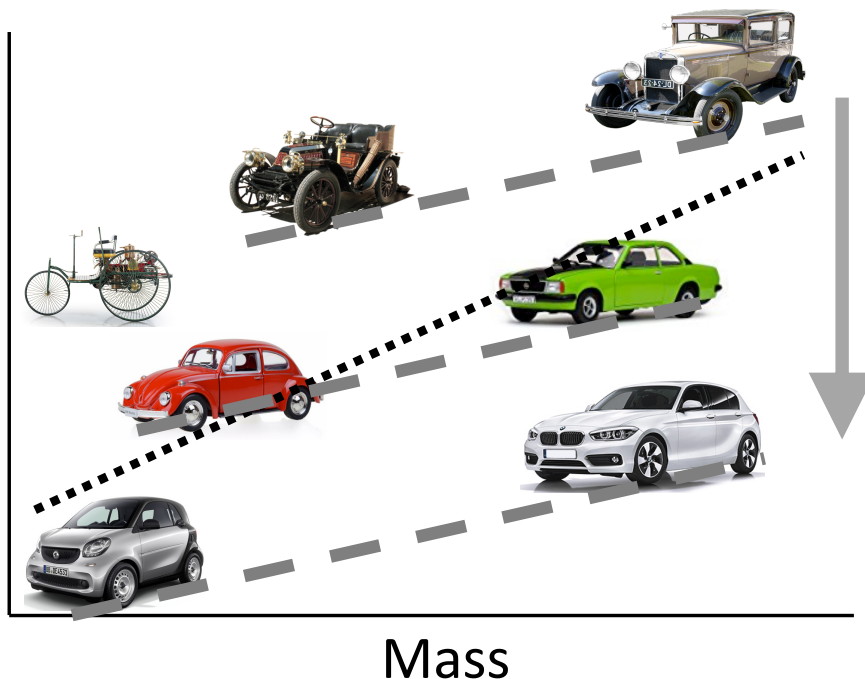
Time per offspring



Mass

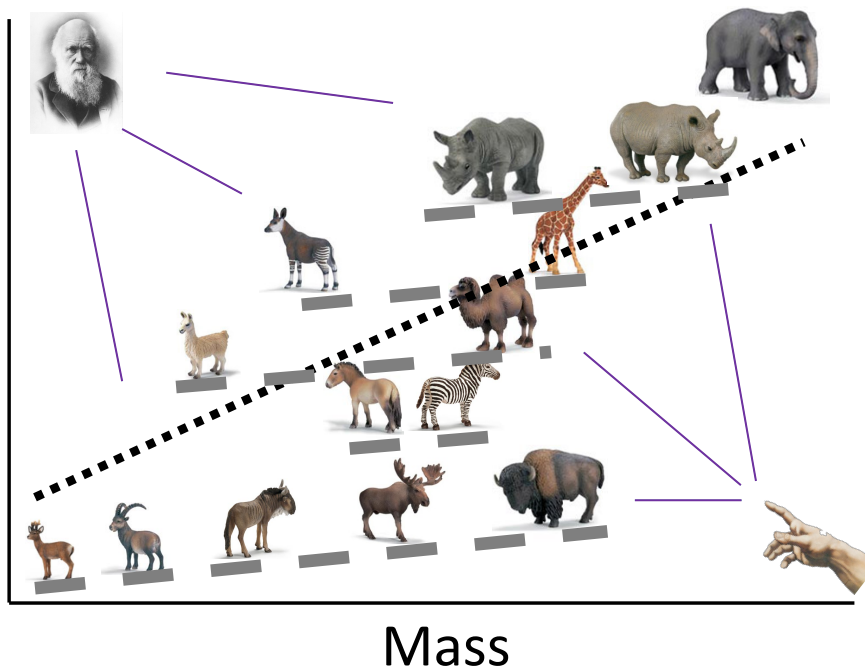


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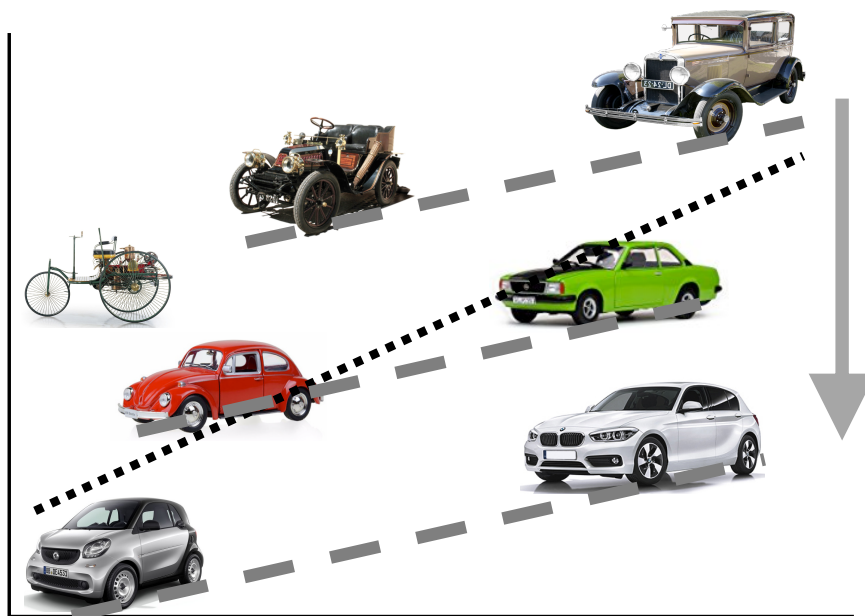
Time per offspring



Why would you consider this a pattern due to fixed life history tradeoff laws?



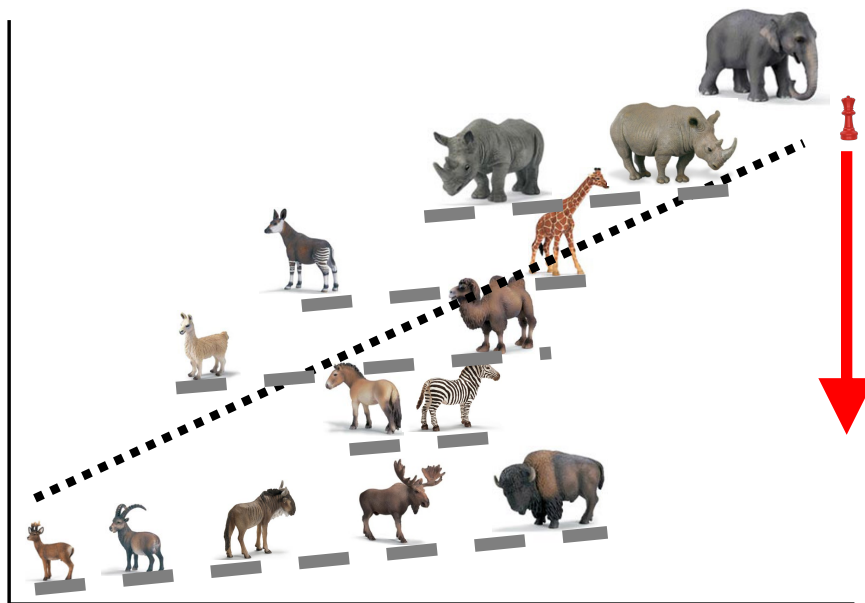
Energy per km



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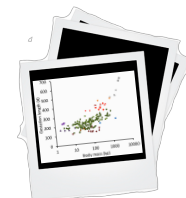
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Time per offspring



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Why would you consider this a pattern due to fixed life history tradeoff laws, and not rather a **snapshot** in a process of optimization?





Some simple *a priori* assumptions and their consequences



A priori conditions and their consequences



A priori conditions and their consequences

Life requires input of resources.



A priori conditions and their consequences

Life requires input of resources.

Life starts simple (non-complex).



A priori conditions and their consequences

Life requires input of resources.

Life starts simple (non-complex).

Life means reproduction.



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- spontaneously occurring yet heritable variability





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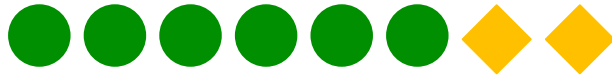
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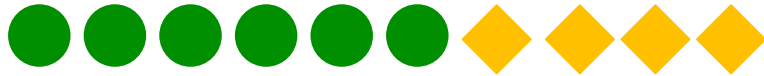
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Probabilistic directionality I: towards non-stasis



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Probabilistic directionality I: towards non-stasis

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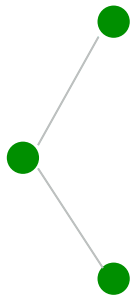
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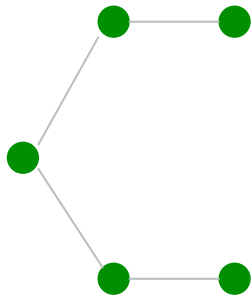
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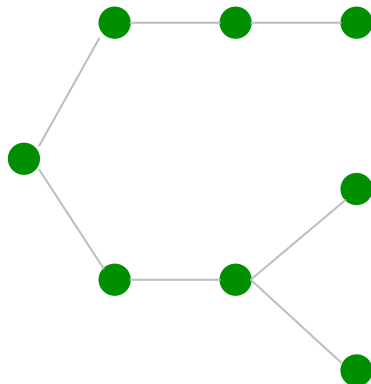
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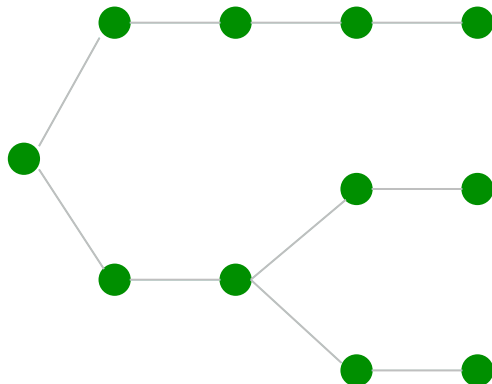
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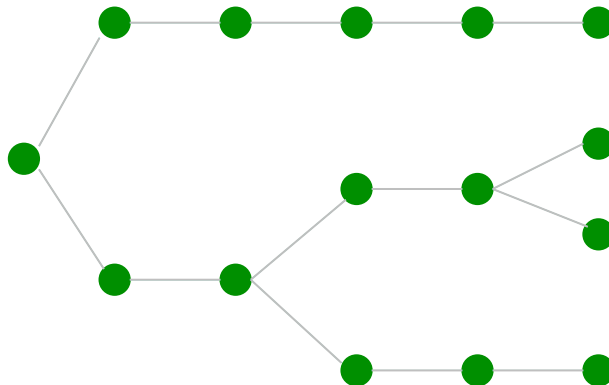
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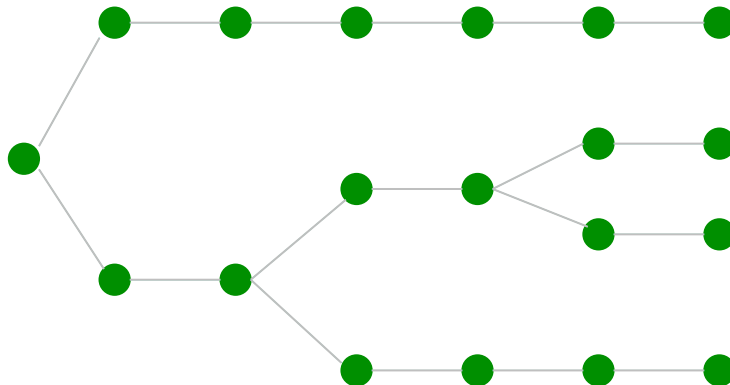
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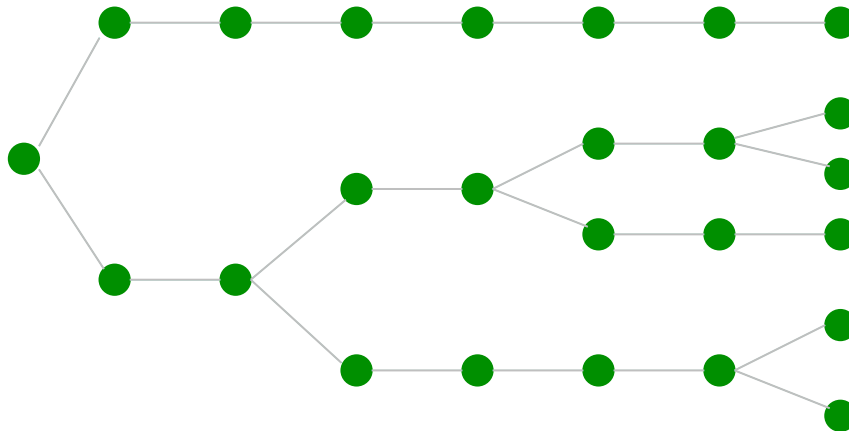
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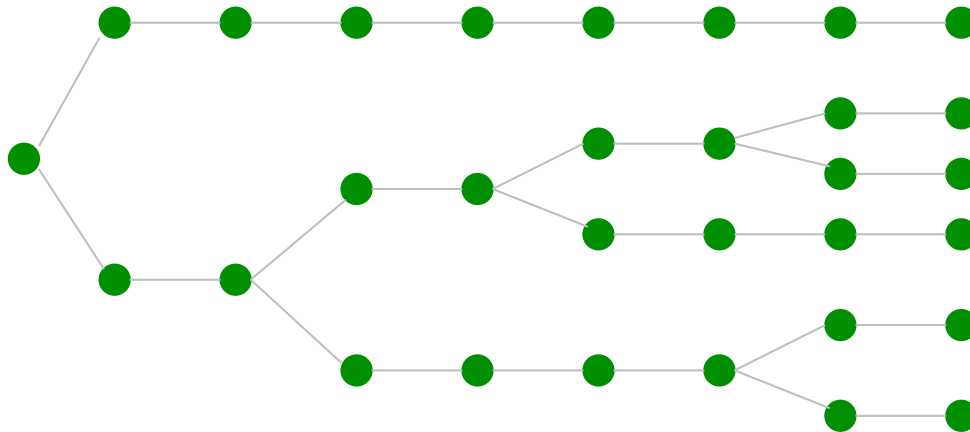
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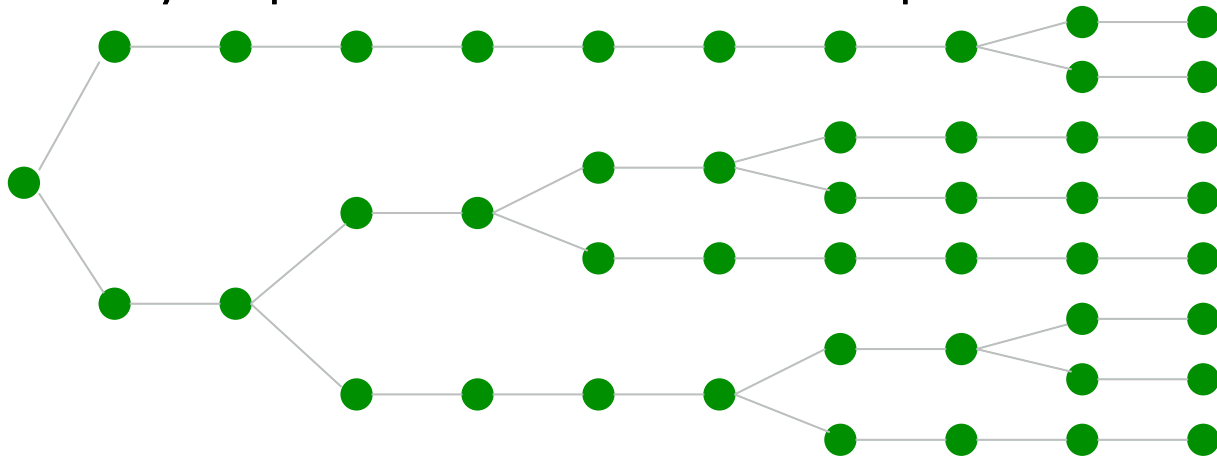
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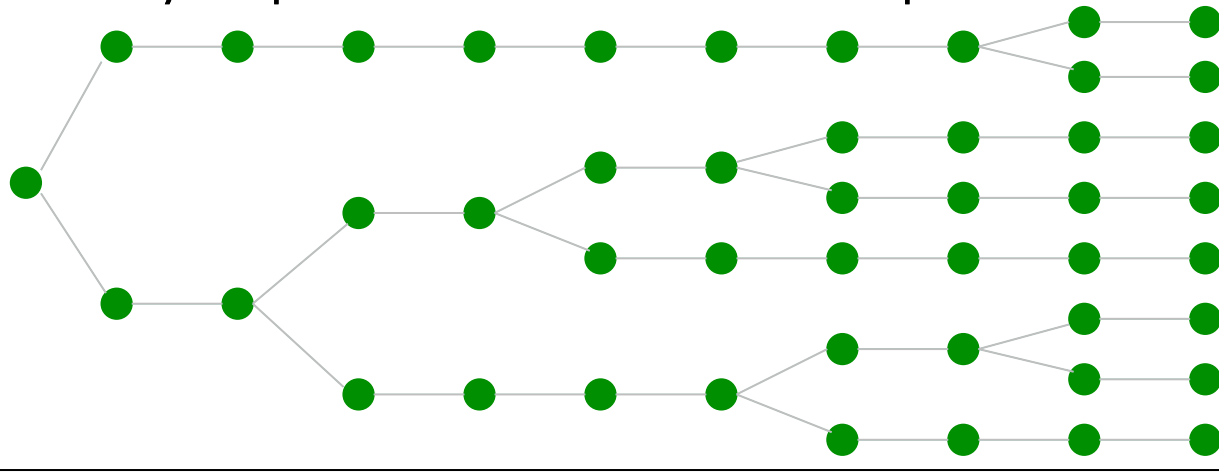
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Probabilistic directionality I: towards non-stasis

- not only replacement but multiplication



Probabilistic directionality II: more



Life starts simple (non-complex).

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- not only replacement but multiplication





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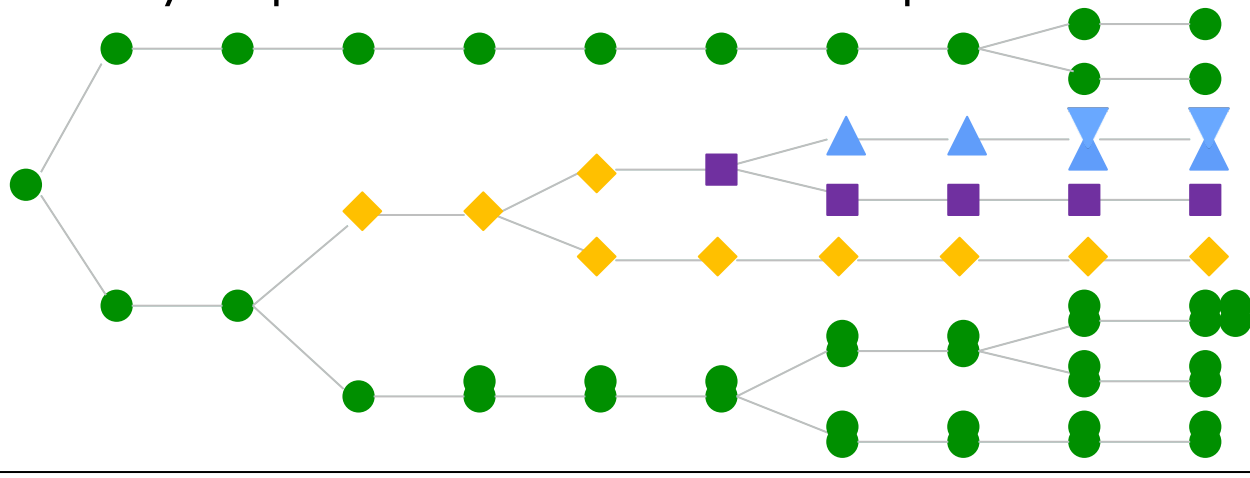
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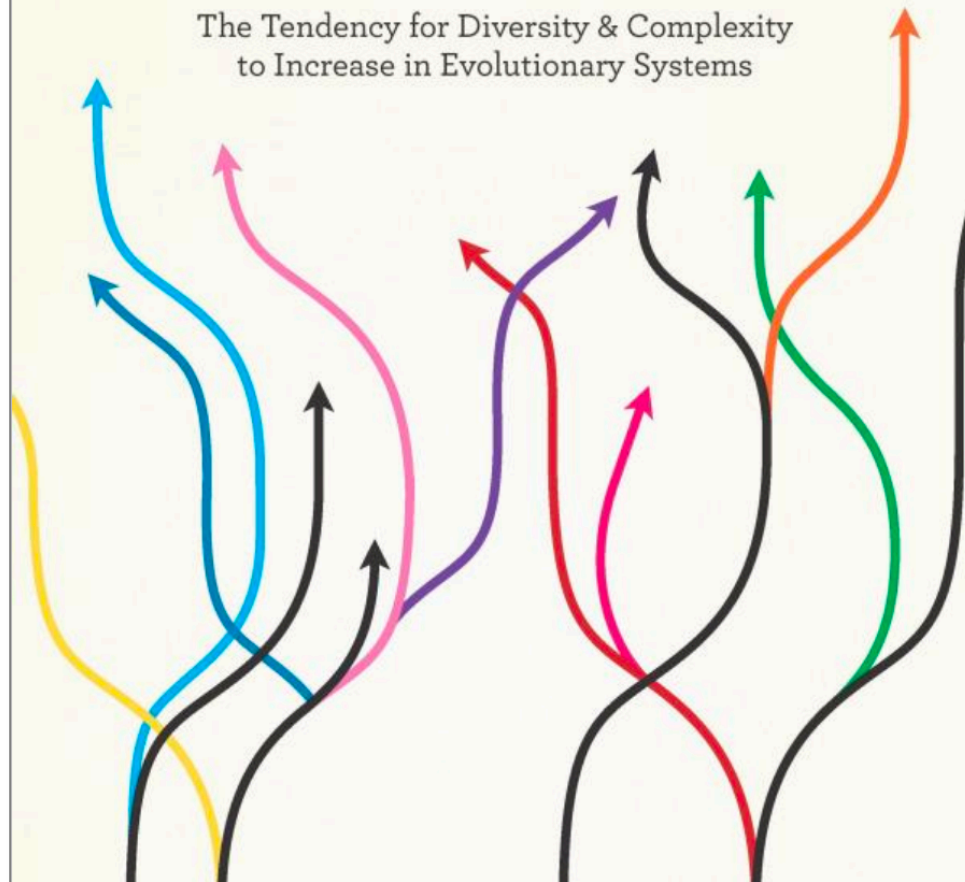
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Probabilistic directionality II: more diversity & complexity



The Tendency for Diversity & Complexity
to Increase in Evolutionary Systems



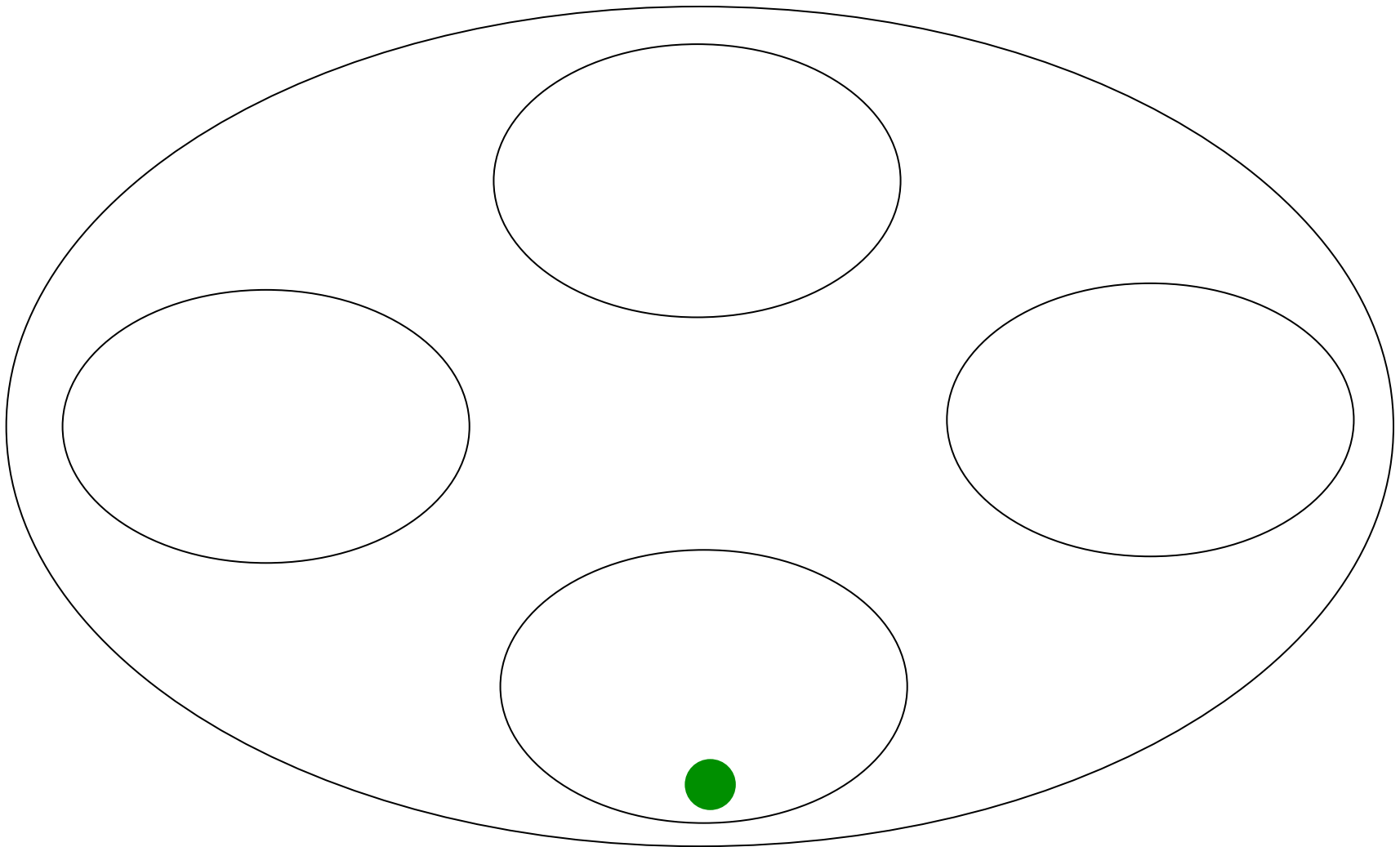
BIOLOGY'S FIRST LAW

DANIEL W. McSHEA & ROBERT N. BRANDON



A priori conditions and their consequences

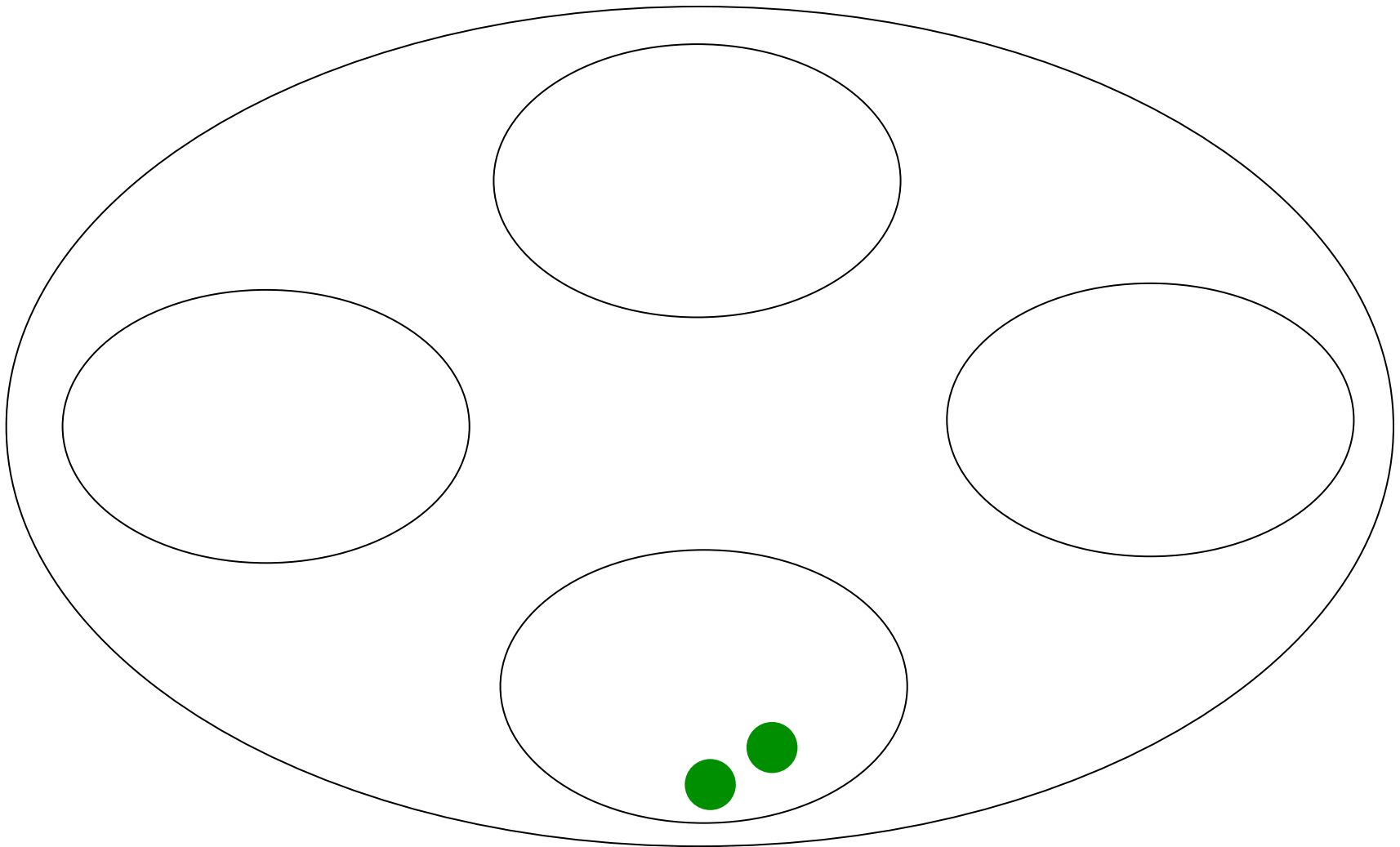
Resources are finite.





A priori conditions and their consequences

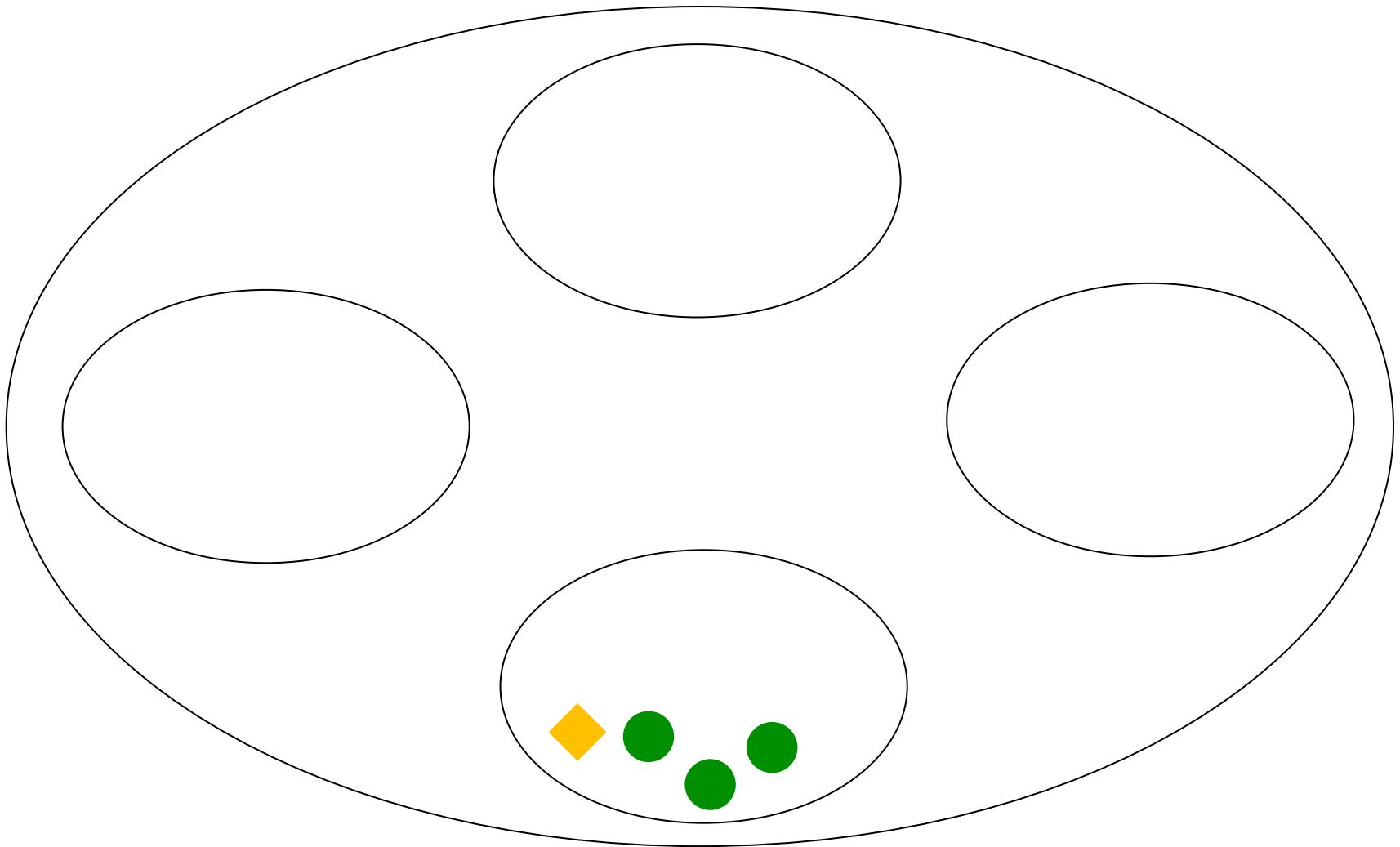
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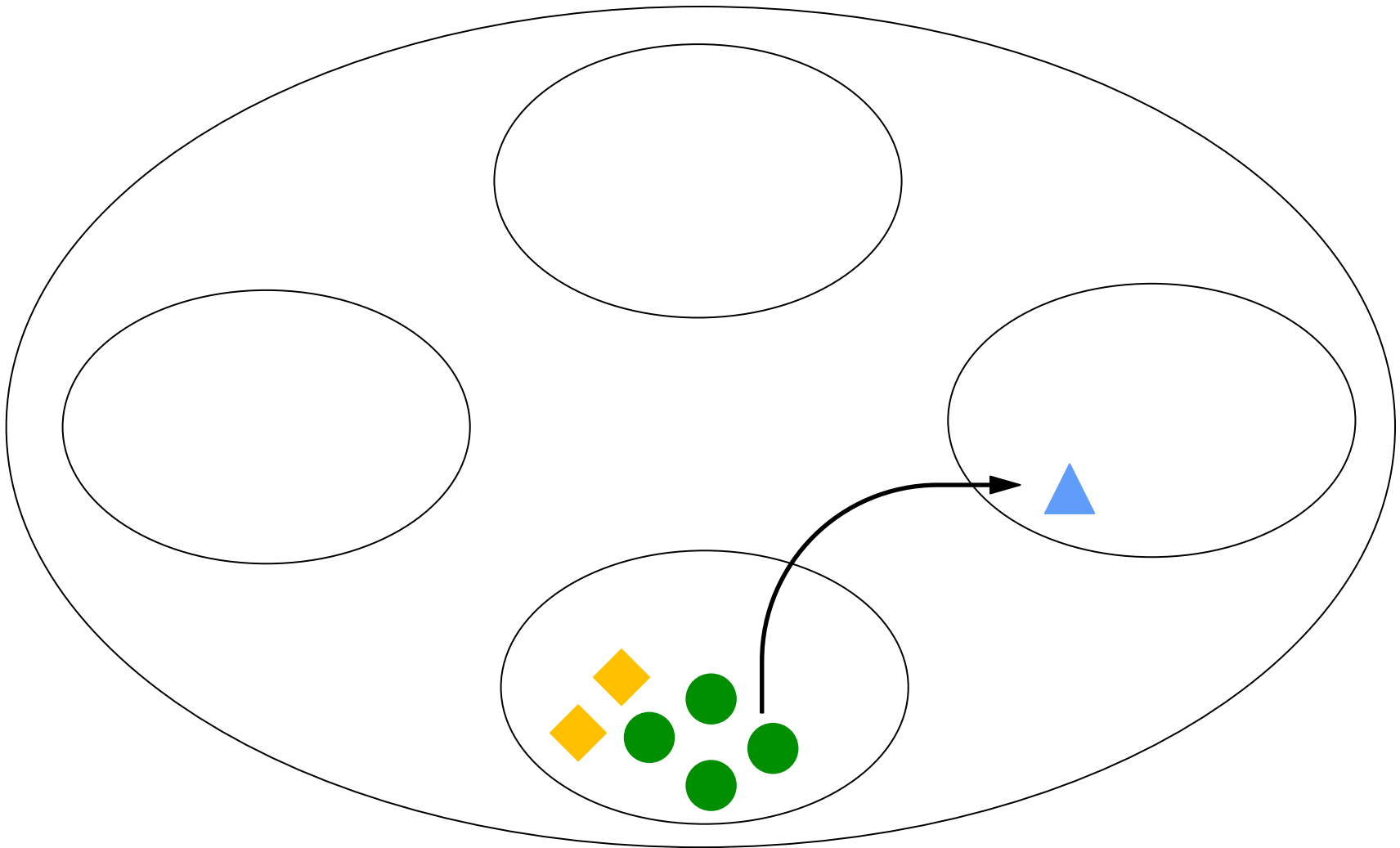
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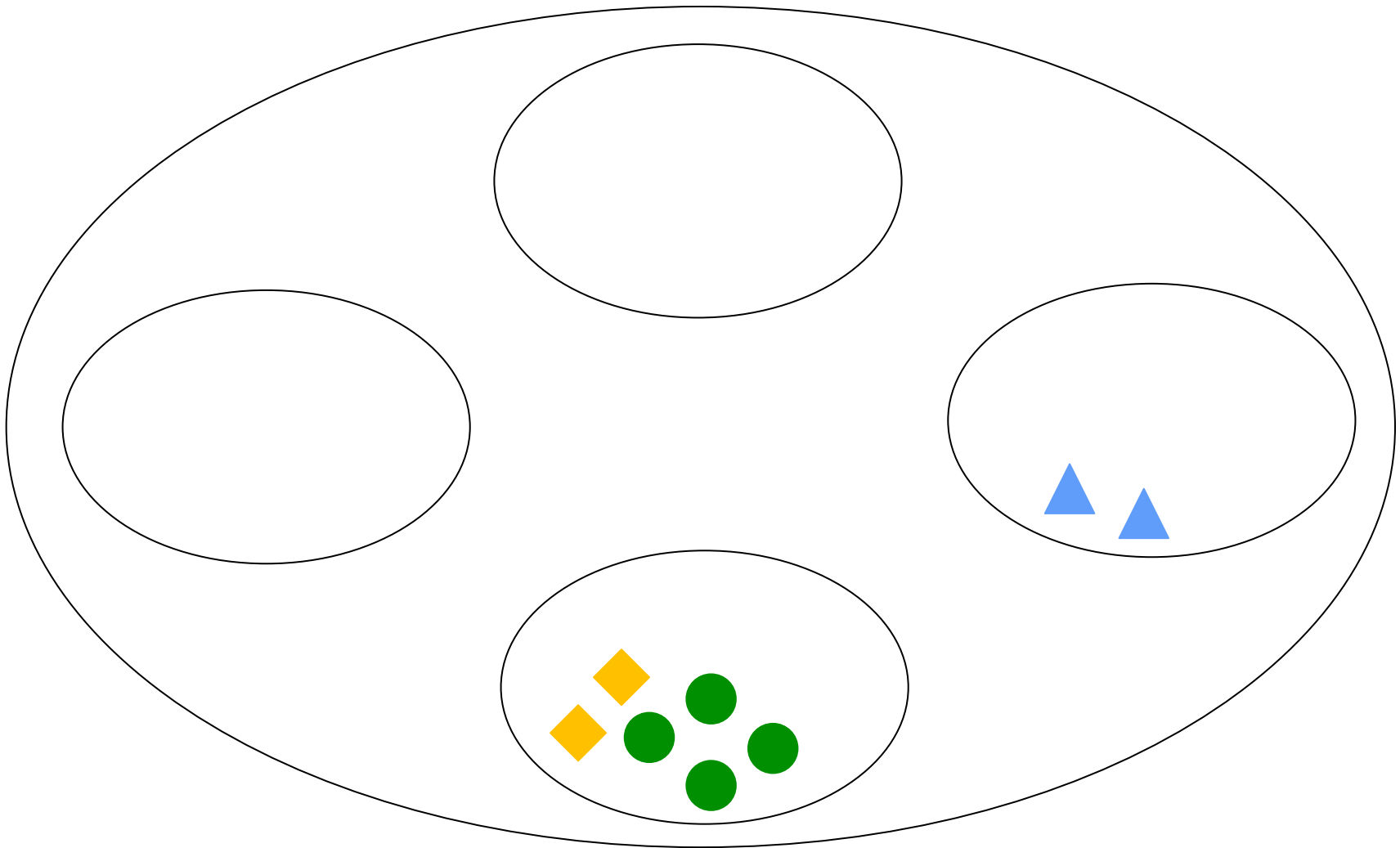
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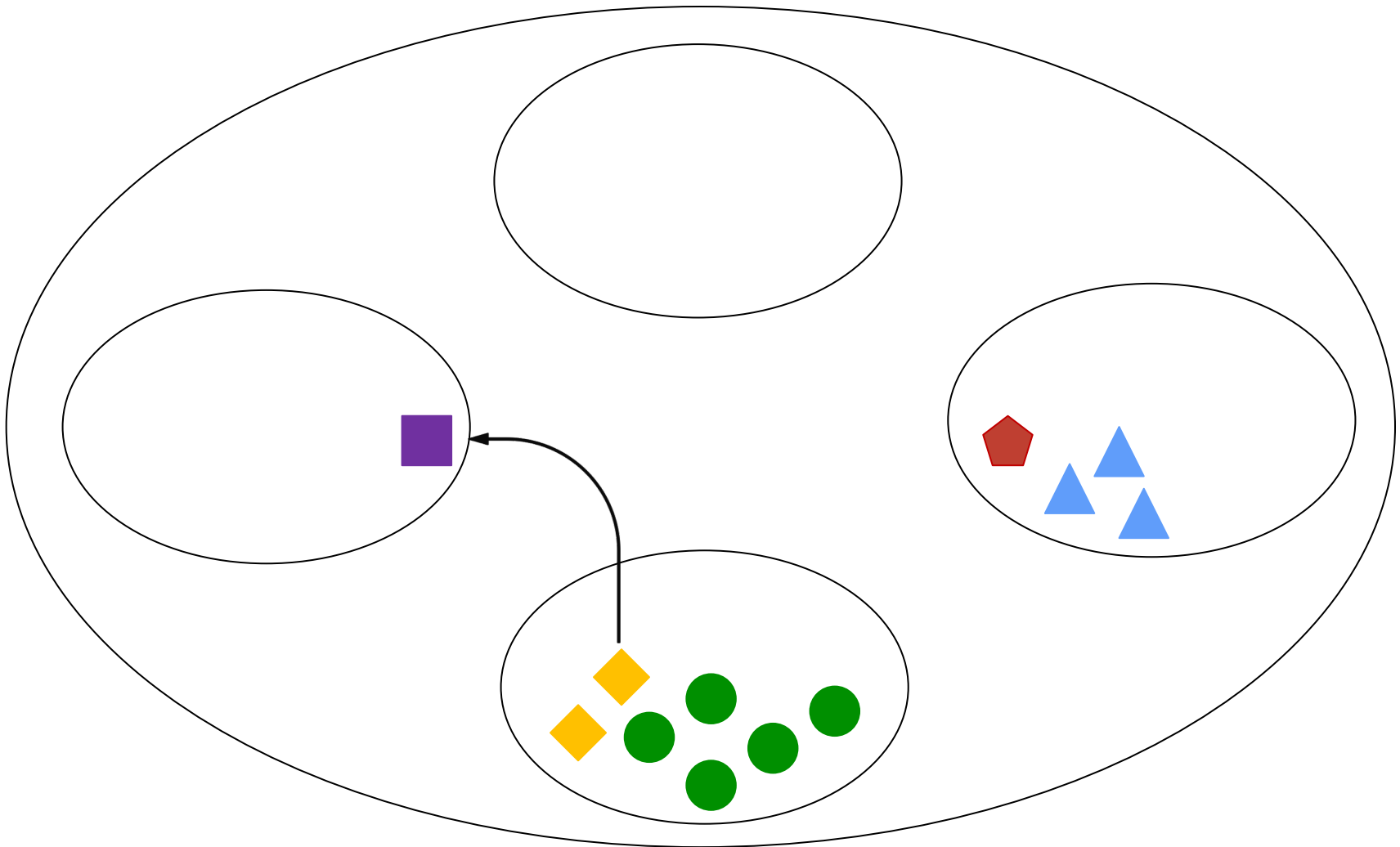
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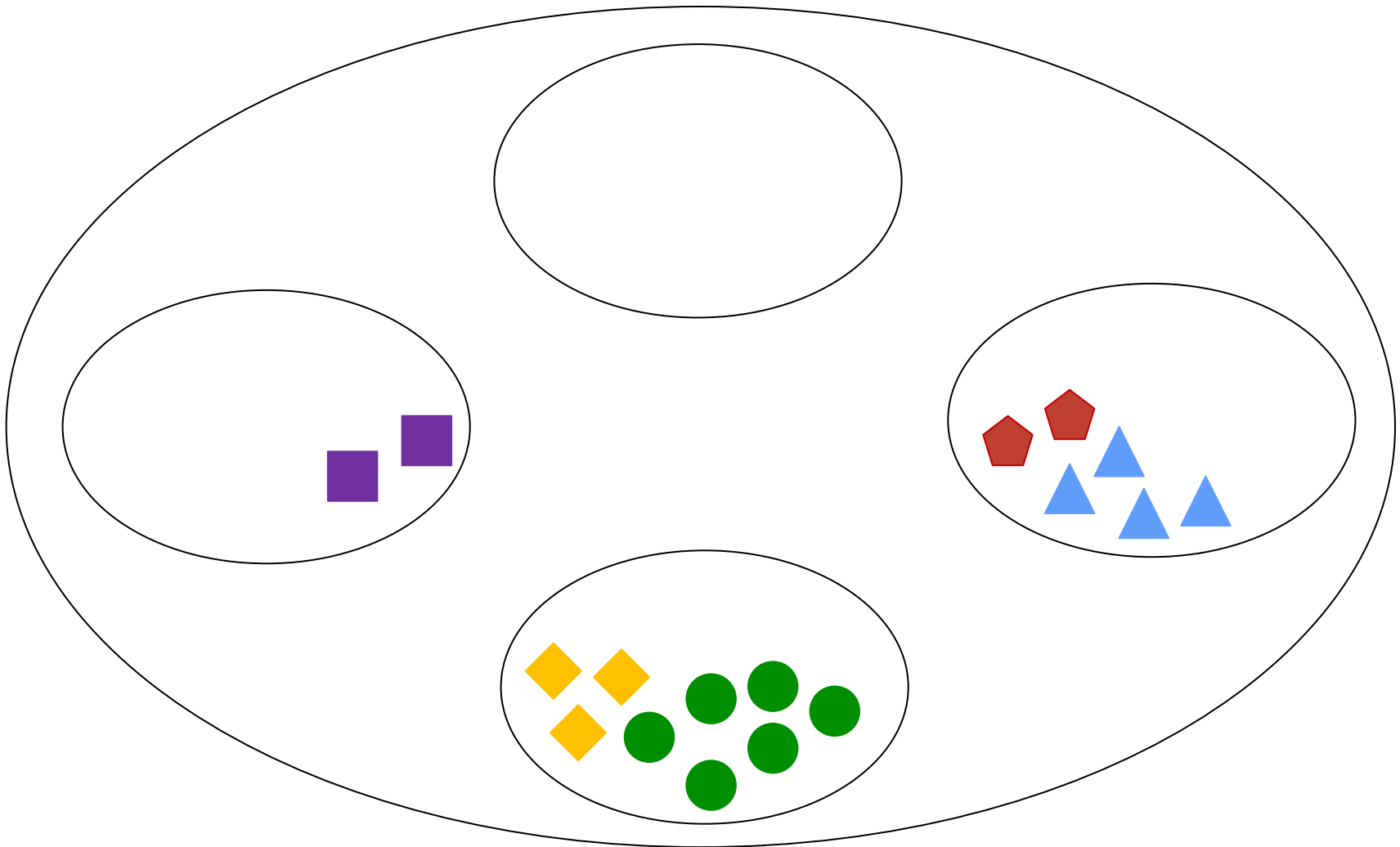
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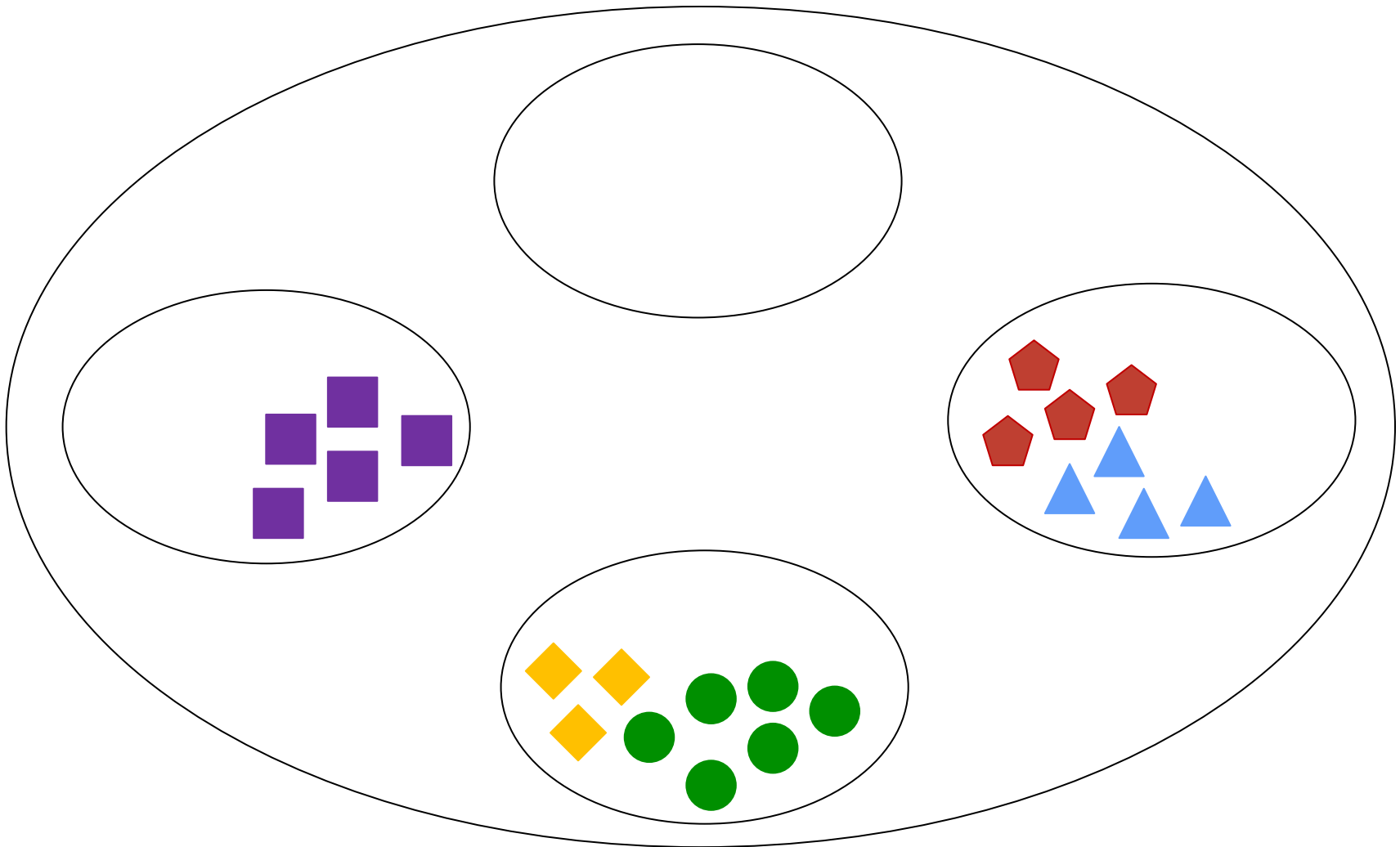
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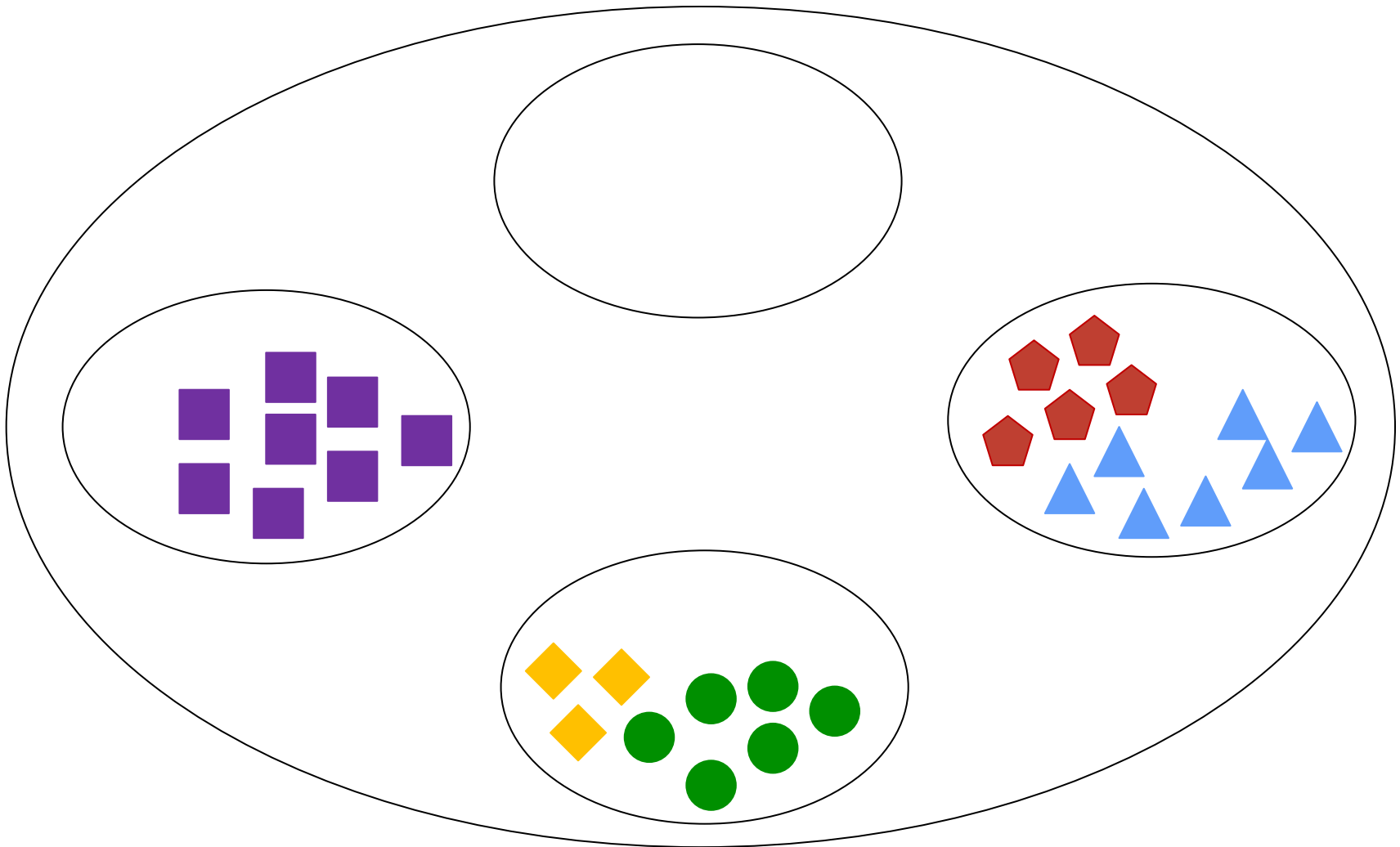
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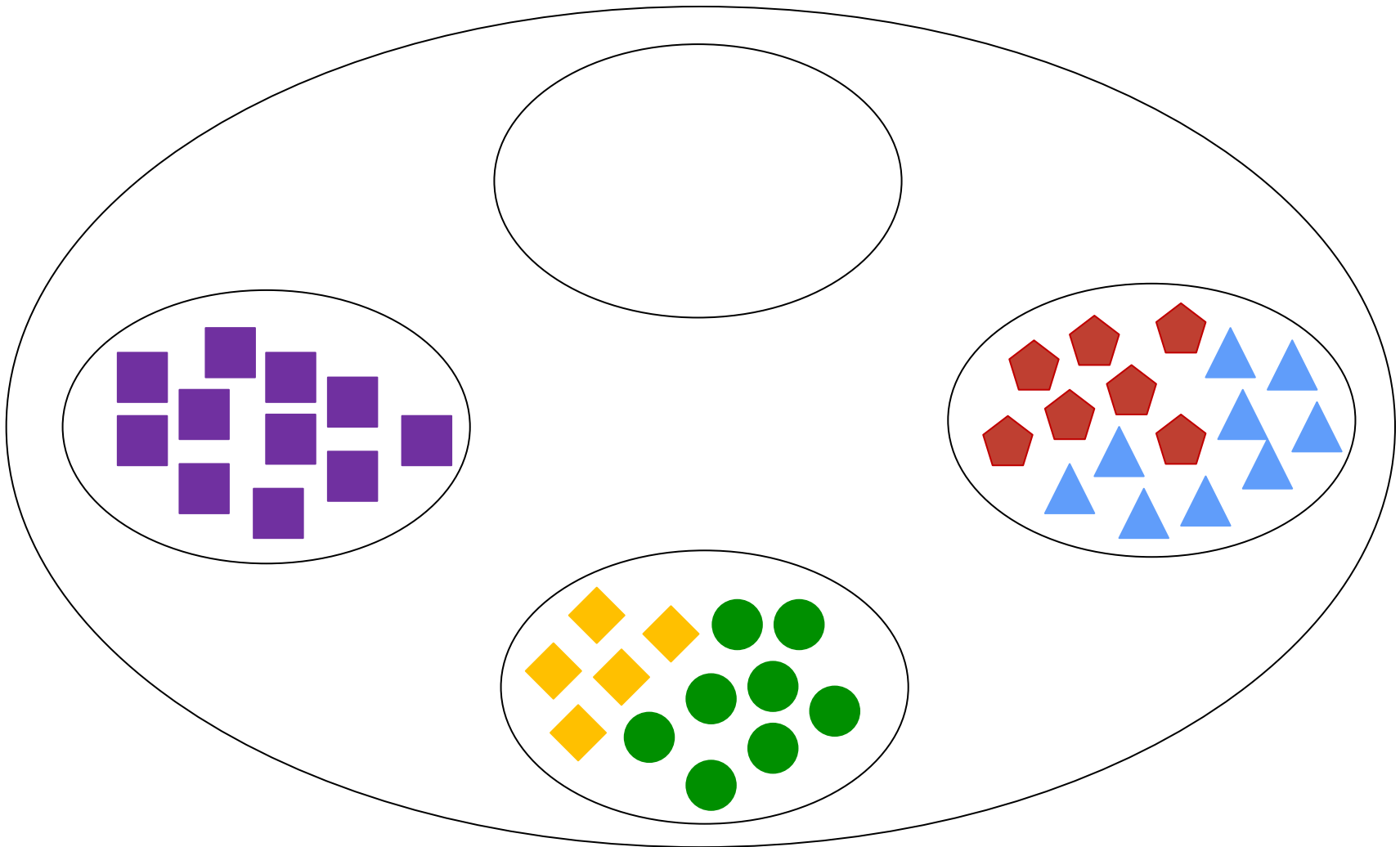
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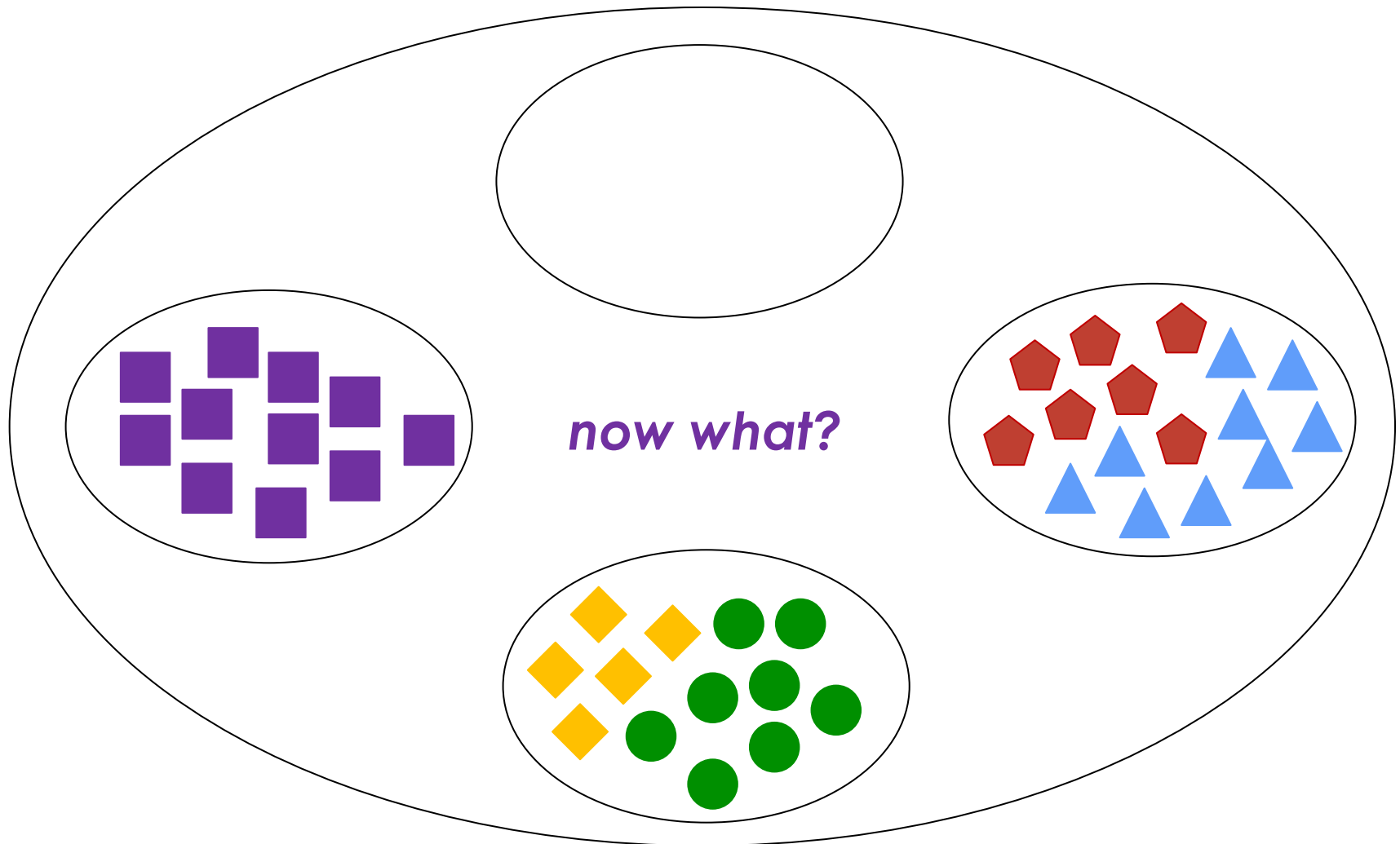
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A priori conditions and their consequences

Resources are finite.





A NEW EVOLUTIONARY LAW

Leigh Van Valen

The Red Queen's Hypothesis (32)

Evol. Theory 1:1-30 (July 1973)





A NEW EVOLUTIONARY LAW

Leigh Van Valen

The Red Queen's Hypothesis (32)

(32). "Now here, you see, it takes all the running you can do, to keep in the same place." (L. Carroll, Through the Looking Glass.)

Evol. Theory 1:1-30 (July 1973)





‘Evolutionary progress’ – directional evolution

Proc. R. Soc. Lond. B **205**, 489–511 (1979)

489

Printed in Great Britain

Arms races between and within species

BY R. DAWKINS AND J. R. KREBS

reverse as to continue the previous one. But in fact consistent directionality is introduced because the environment of any one evolving lineage includes other evolving lineages. Above all, it is because adaptations in one lineage call forth counter-adaptations in others, setting in motion the unstable evolutionary progressions we call arms races.





‘Evolutionary progress’ – directional evolution



The crab is the natural predator of the snail.



Natural selection favors snails with thicker shells and spines.



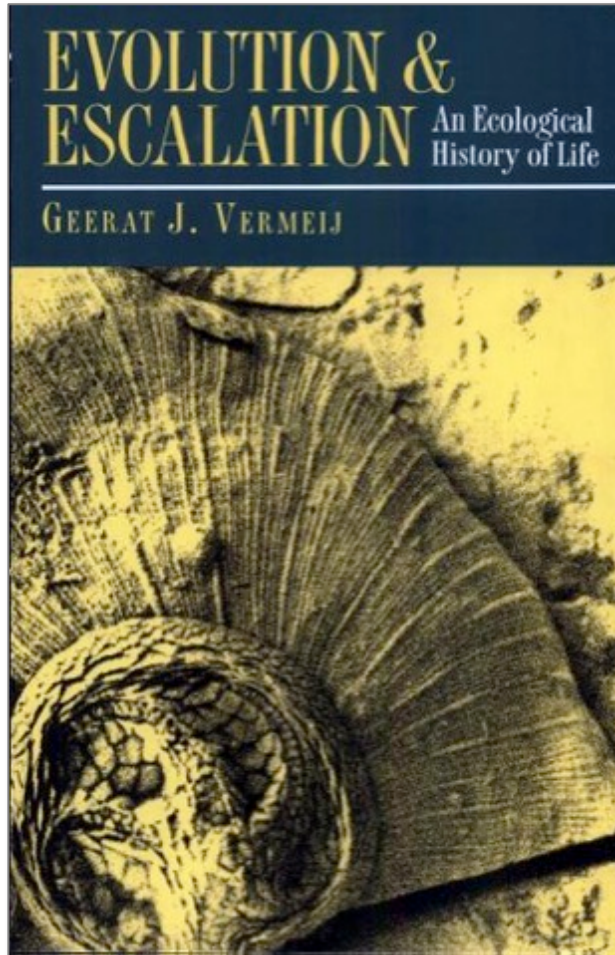
Through natural selection, crabs evolve more powerful claws that can pierce the snails' thick, spiny shells.

In response, natural selection favors snails with even thicker shells and spines.





‘Evolutionary progress’ – directional evolution



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On Escalation

Geerat J. Vermeij

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Keywords

Red Queen, coevolution, Phanerozoic, enabling factors, predation, competition

Abstract

Organisms have been important agents of selection throughout the history of life. The processes and outcomes of this selection are the subject of this review. Among these, escalation is the most widespread. The primary selective agents are powerful competitors and consumers, which together push many populations toward higher performance in acquiring and defending resources while relegating less competitive species to physiologically marginal settings, where escalation also ensues. The extent to which performance standards rise depends on enabling factors, which control availability of and access to resources. By establishing positive feedbacks between species and enabling factors, effective competitors regulate and enhance resource supply. The pace of escalation toward greater power and reach is dictated by geological factors as well as by growing interdependencies between species and their resources. Evolutionary events on land related to the production of oxygen may have been instrumental in triggering the major episodes of escalation.



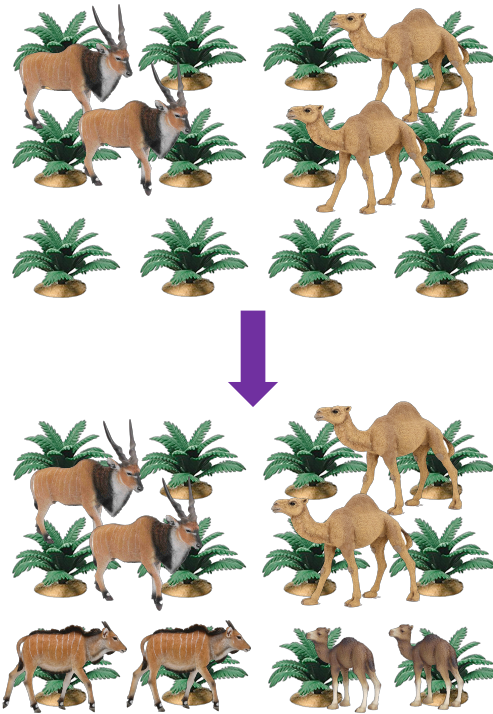


Equivalent use of limited resources





Stasis





Stasis





Competition for limited resources



Competition for limited resources



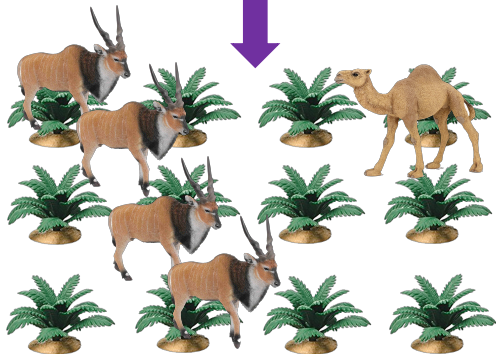


Competition for limited resources



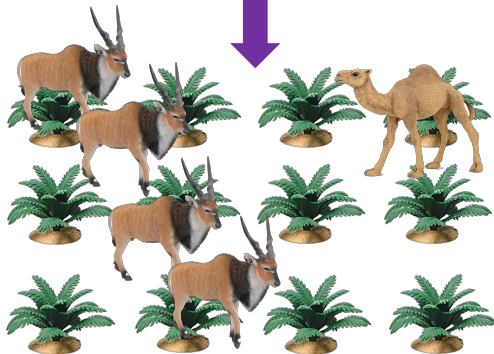


Competition for limited resources



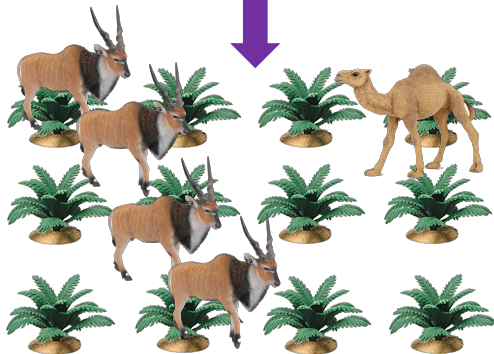


Competition for limited resources



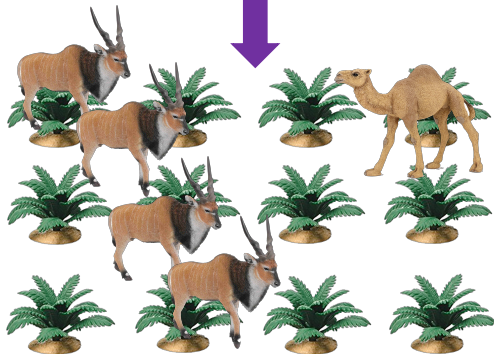


Competition for limited resources





Competition for limited resources





Are conditions stable enough so that the direction of a Darwinian Demon is always the same?





The Red Queen and the Court Jester: Species Diversity and the Role of Biotic and Abiotic Factors Through Time

Michael J. Benton

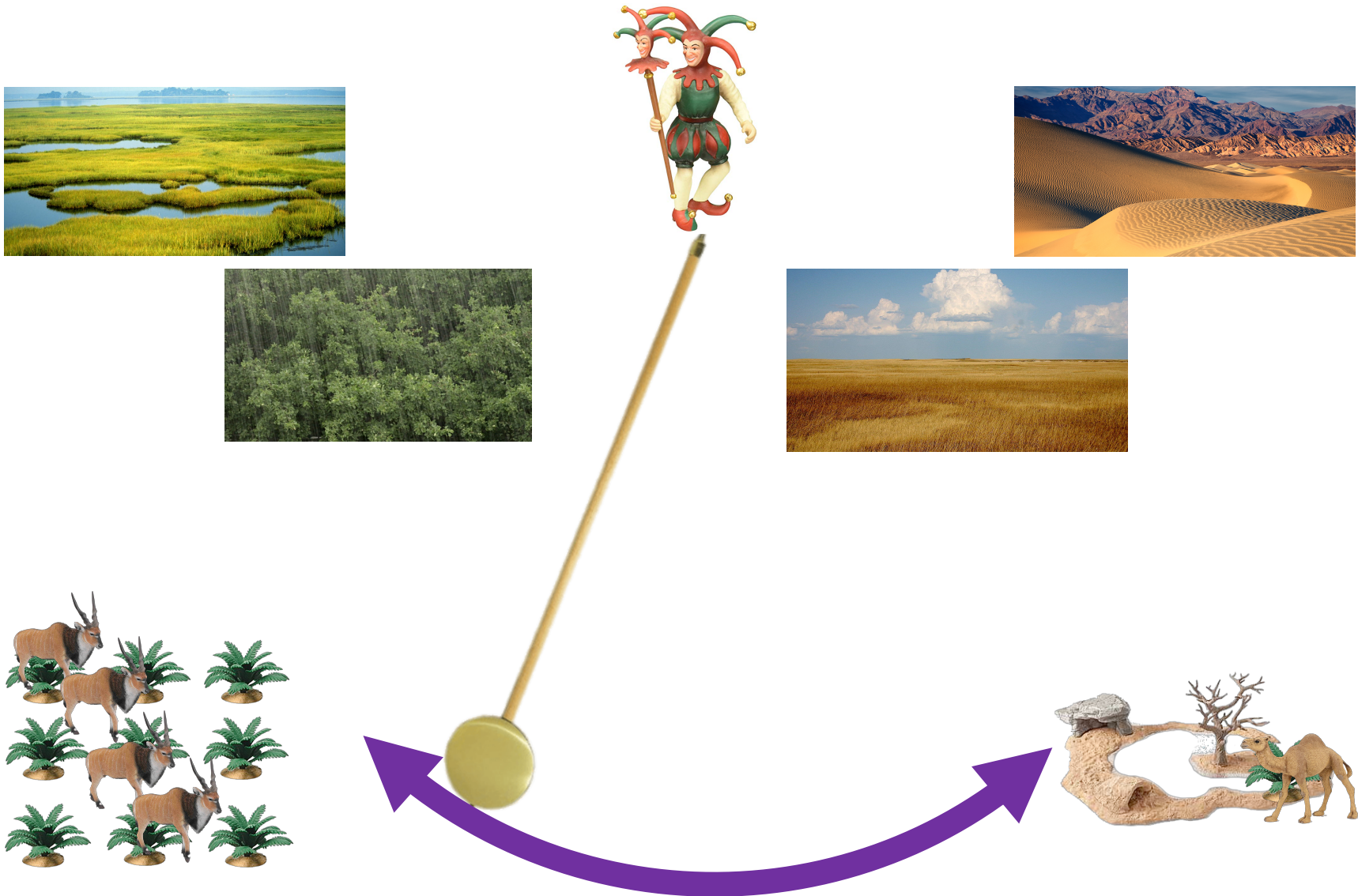
FEBRUARY 2009 VOL 323 SCIENCE



The Court Jester's pendulum



The Court Jester's pendulum



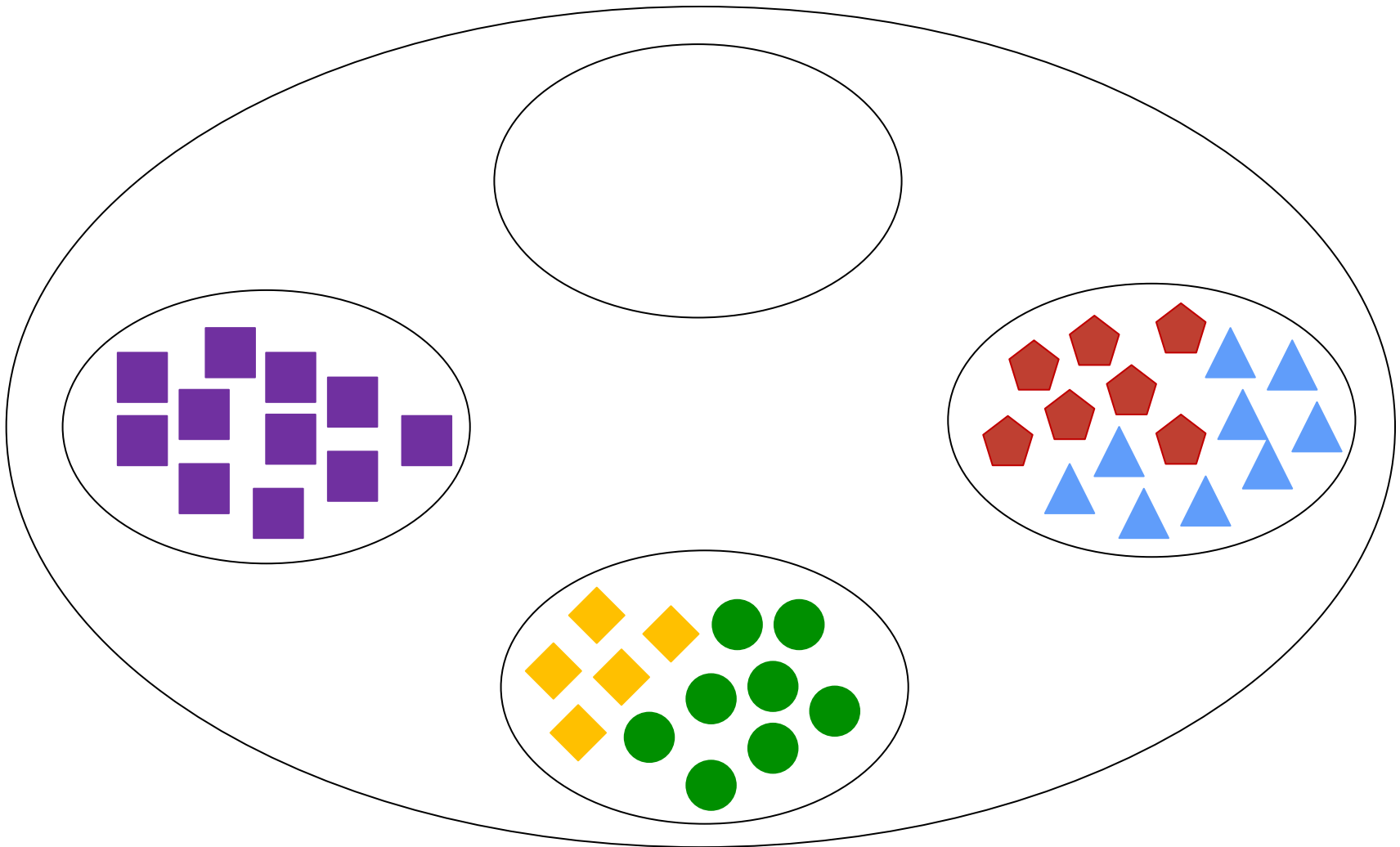
The Court Jester's pendulum





A priori conditions and their consequences

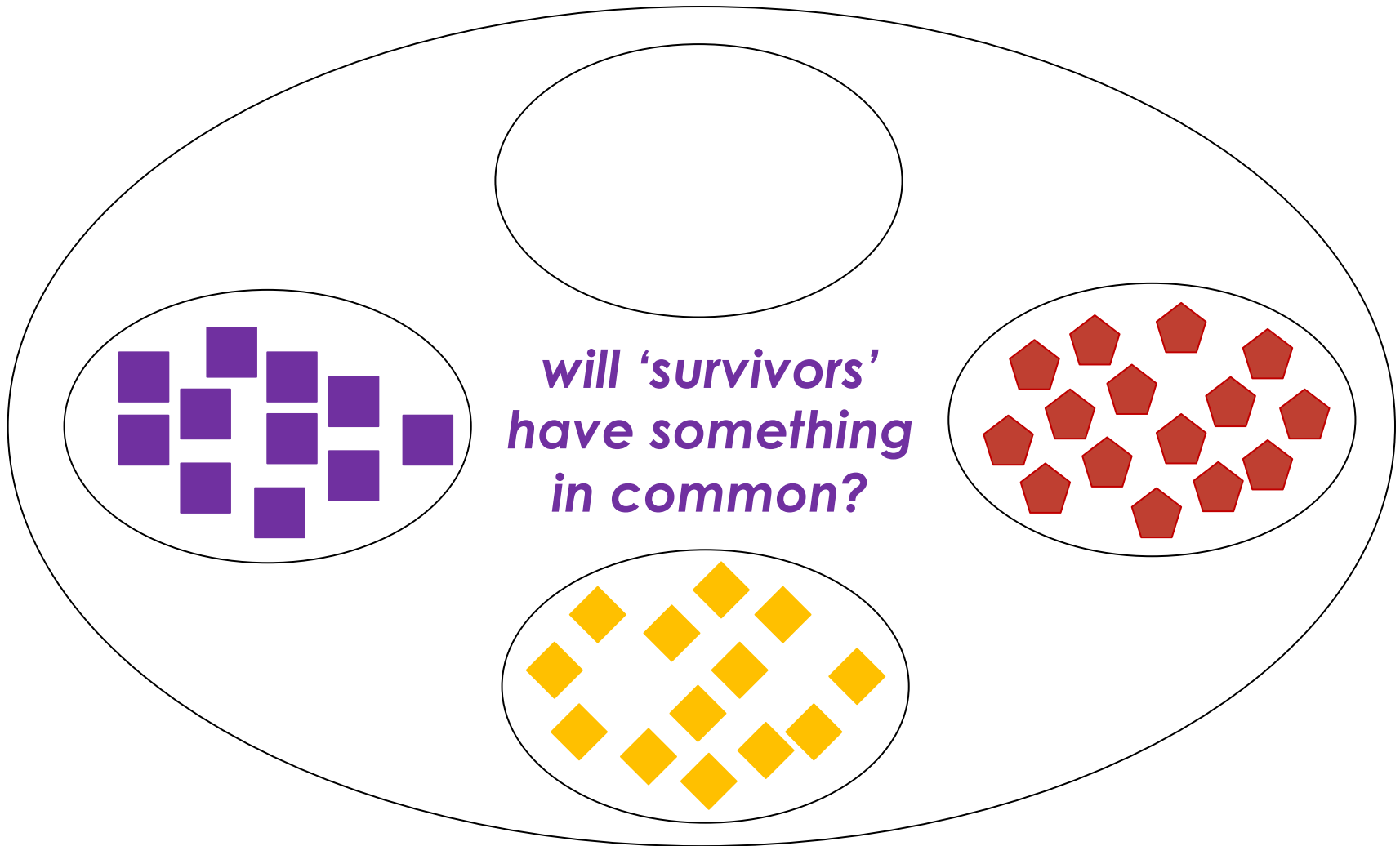
Resources are finite.





A priori conditions and their consequences

Resources are finite.





'Evolutionary progress' – directional evolution

Biol. Rev. (1987), **62**, pp. 305–338

PROGRESS AND COMPETITION IN MACROEVOLUTION

By MICHAEL J. BENTON

It is merely a tautology to identify the later animal (the 'winner') as a 'superior competitor' in the absence of any other evidence (Schopf, 1979).

it is hard to envisage a constant competitive advantage that lasted so long and persistently favoured all of the species of one large taxon against all of the species of another in all environments.



Darwinian evolution is a set of rules where the one constant (demographic) selective pressure is to outreproduce competitors.

*In the presence of competitors, Darwinian selection should always go in the **direction** of a 'Darwinian Demon'.*





Darwinian demon



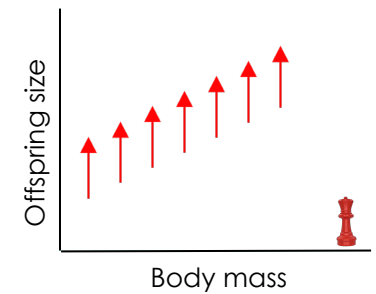
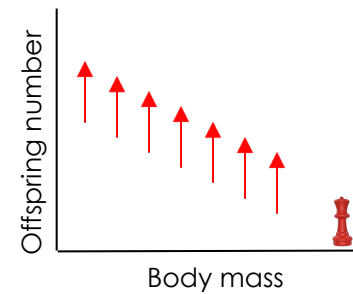
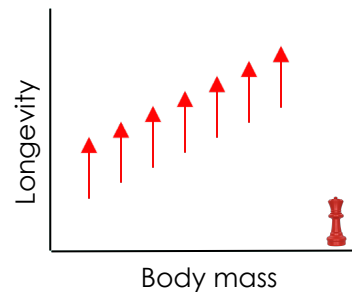
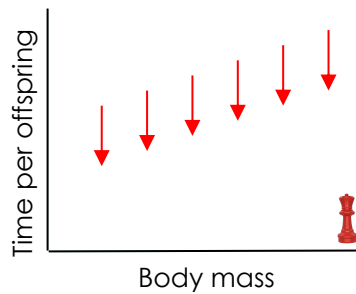
An organism that starts reproducing directly after birth, producing a large number of surviving offspring at extreme speed without ever dying.



Darwinian demon



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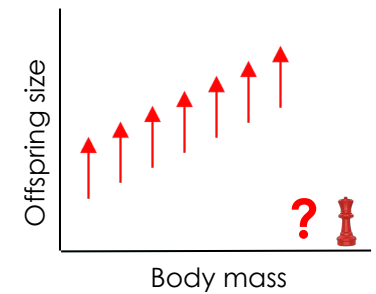
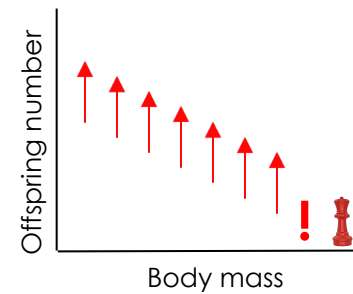
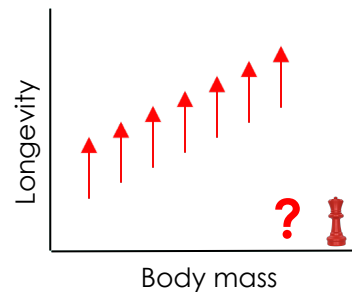
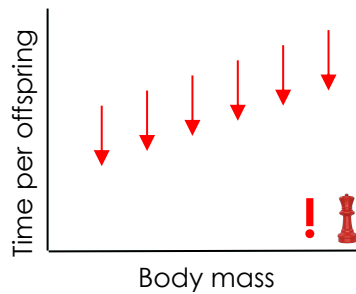




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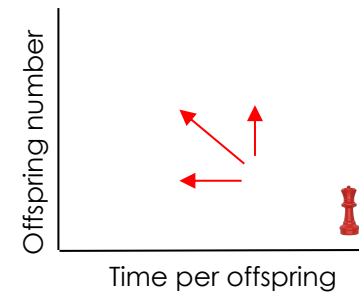
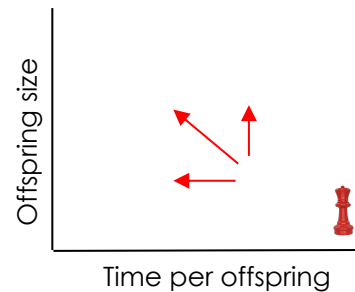
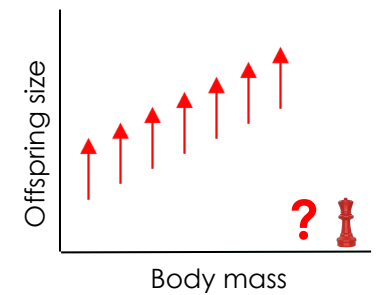
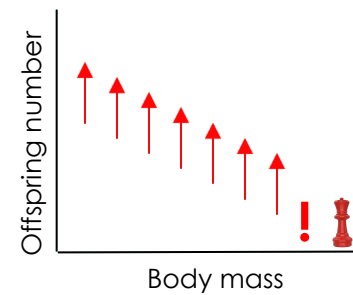
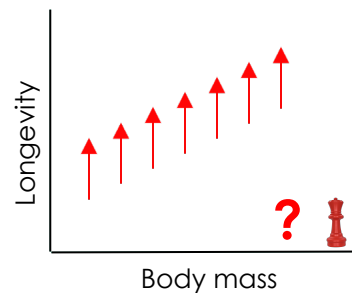
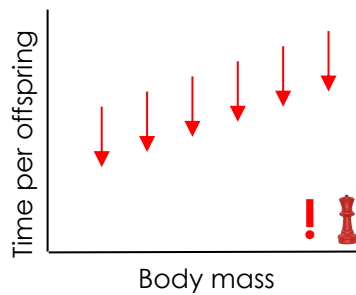




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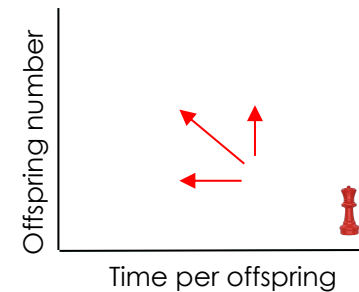
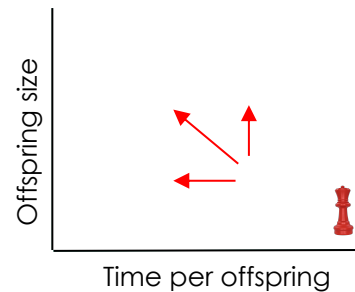
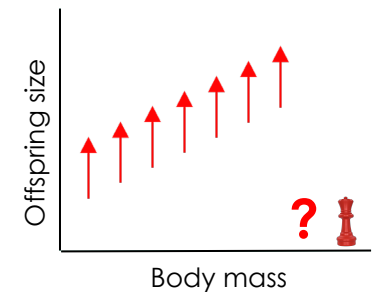
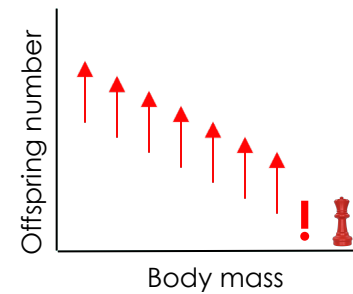
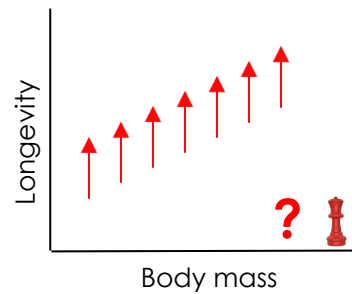
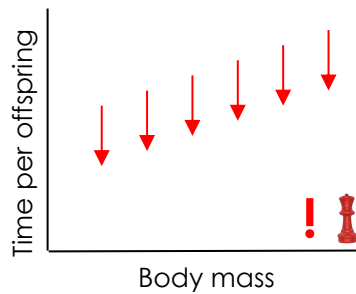




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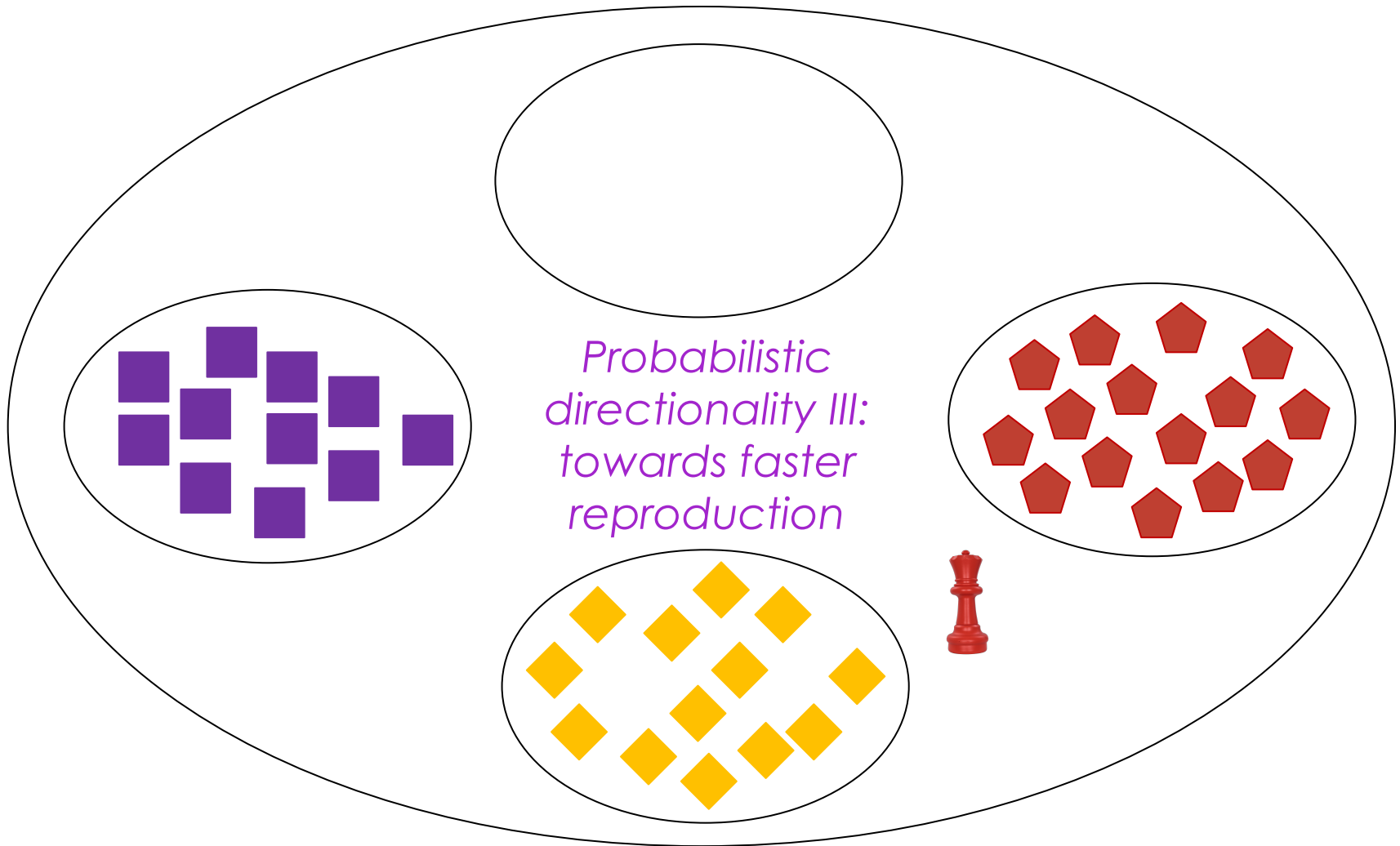
FASTER





A priori conditions and their consequences

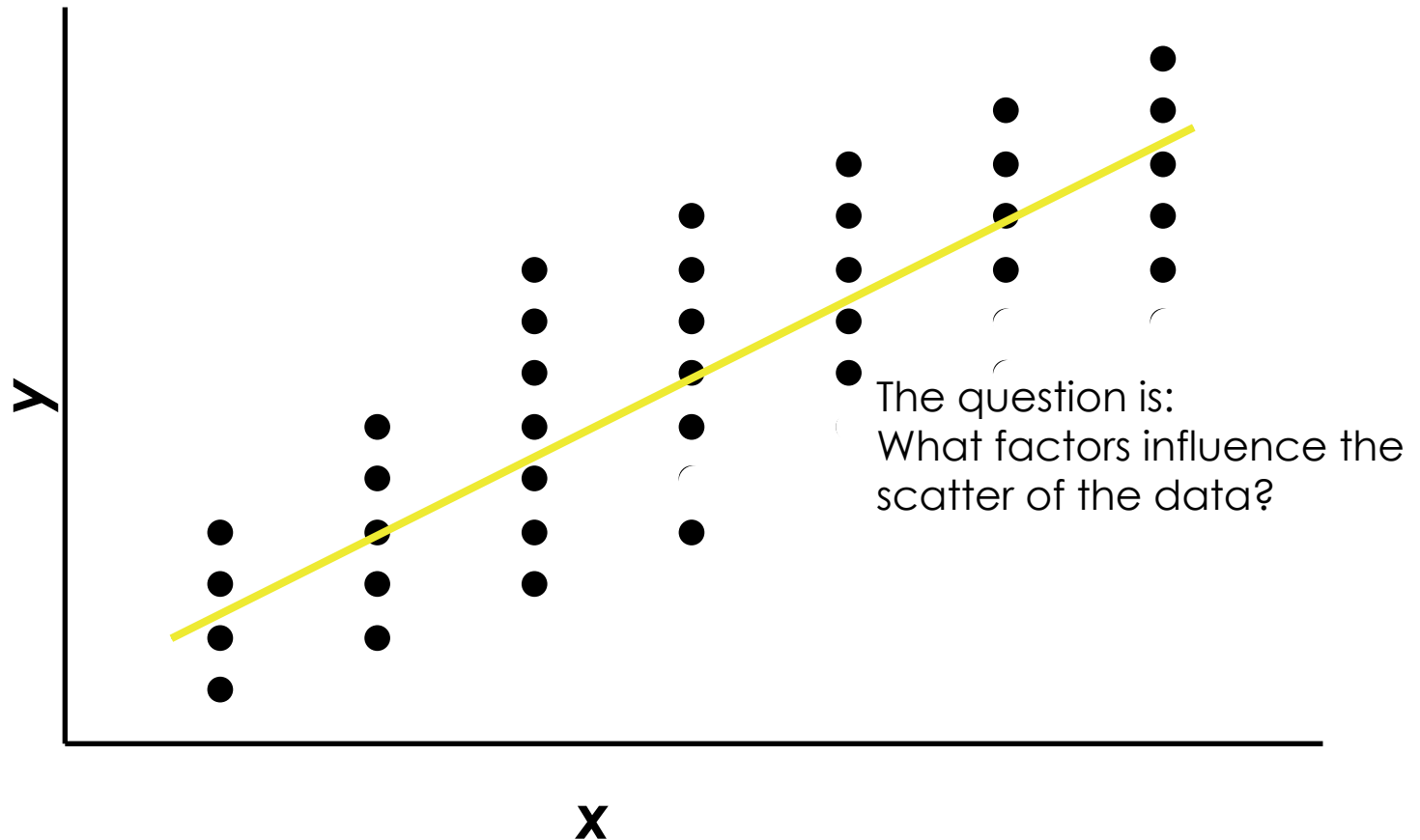
Resources are finite.





Interpreting scaling

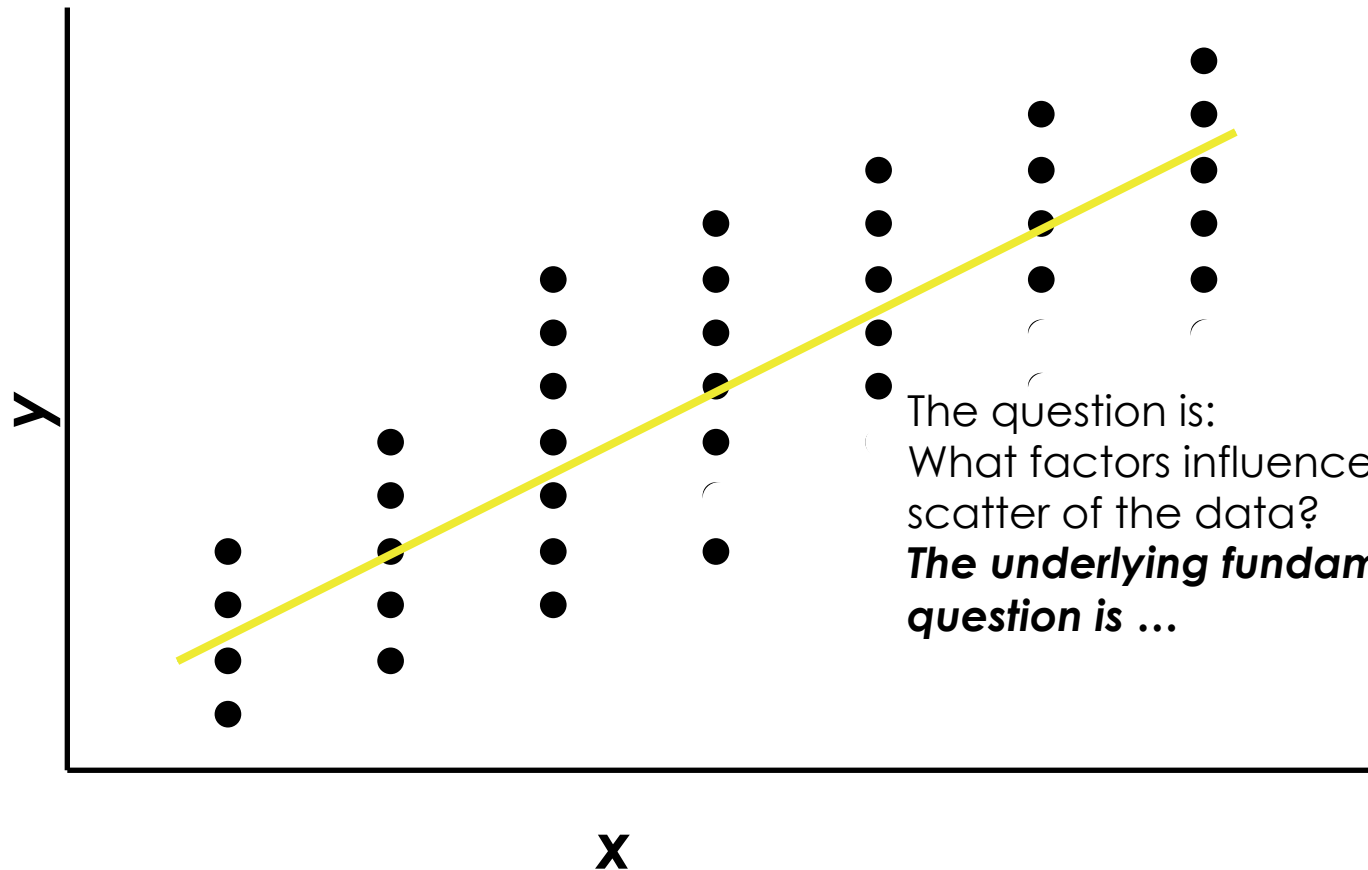
Morphological, physiological and life history variables scale.





Interpreting scaling

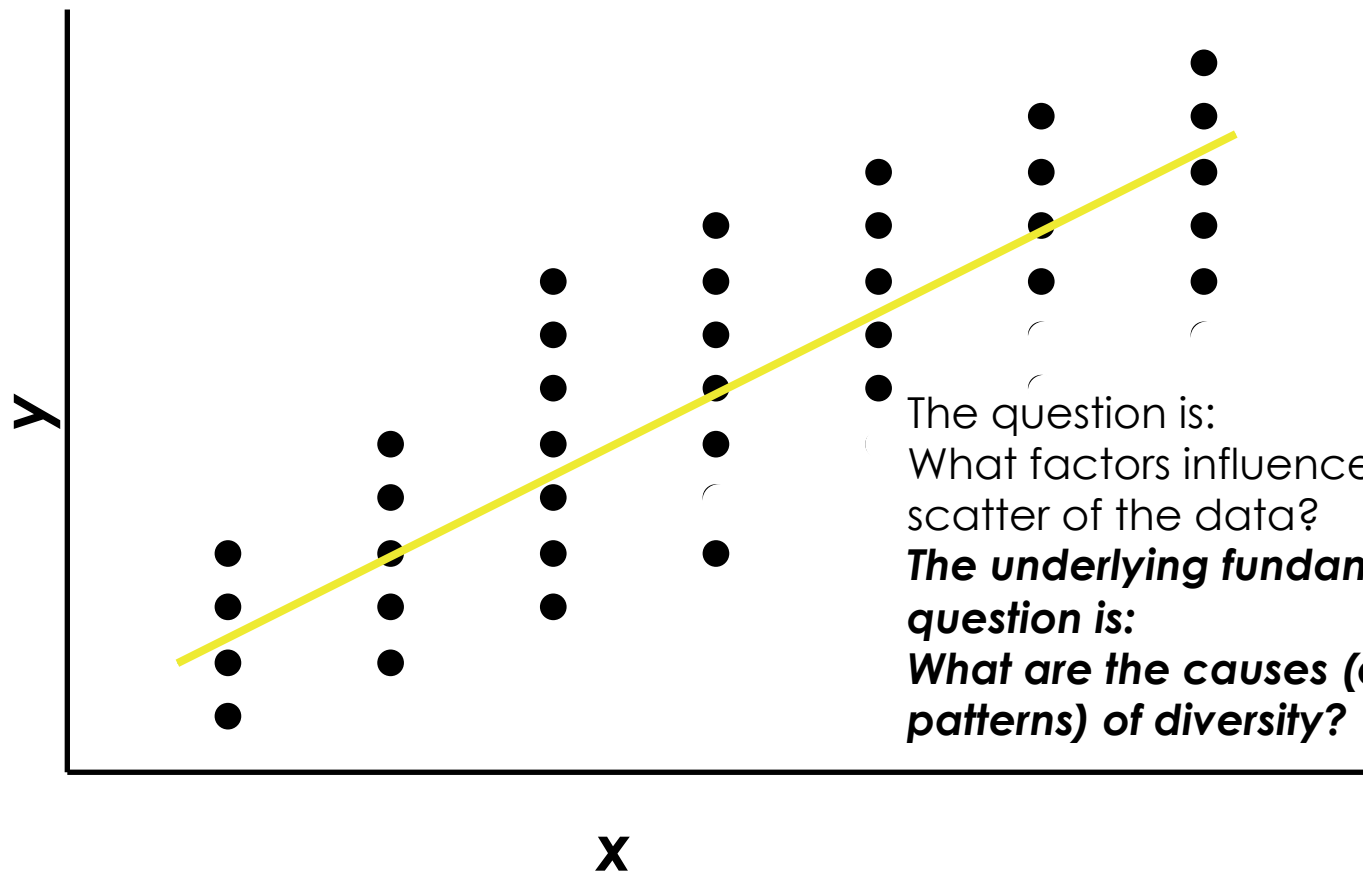
Morphological, physiological and life history variables scale.





Interpreting scaling

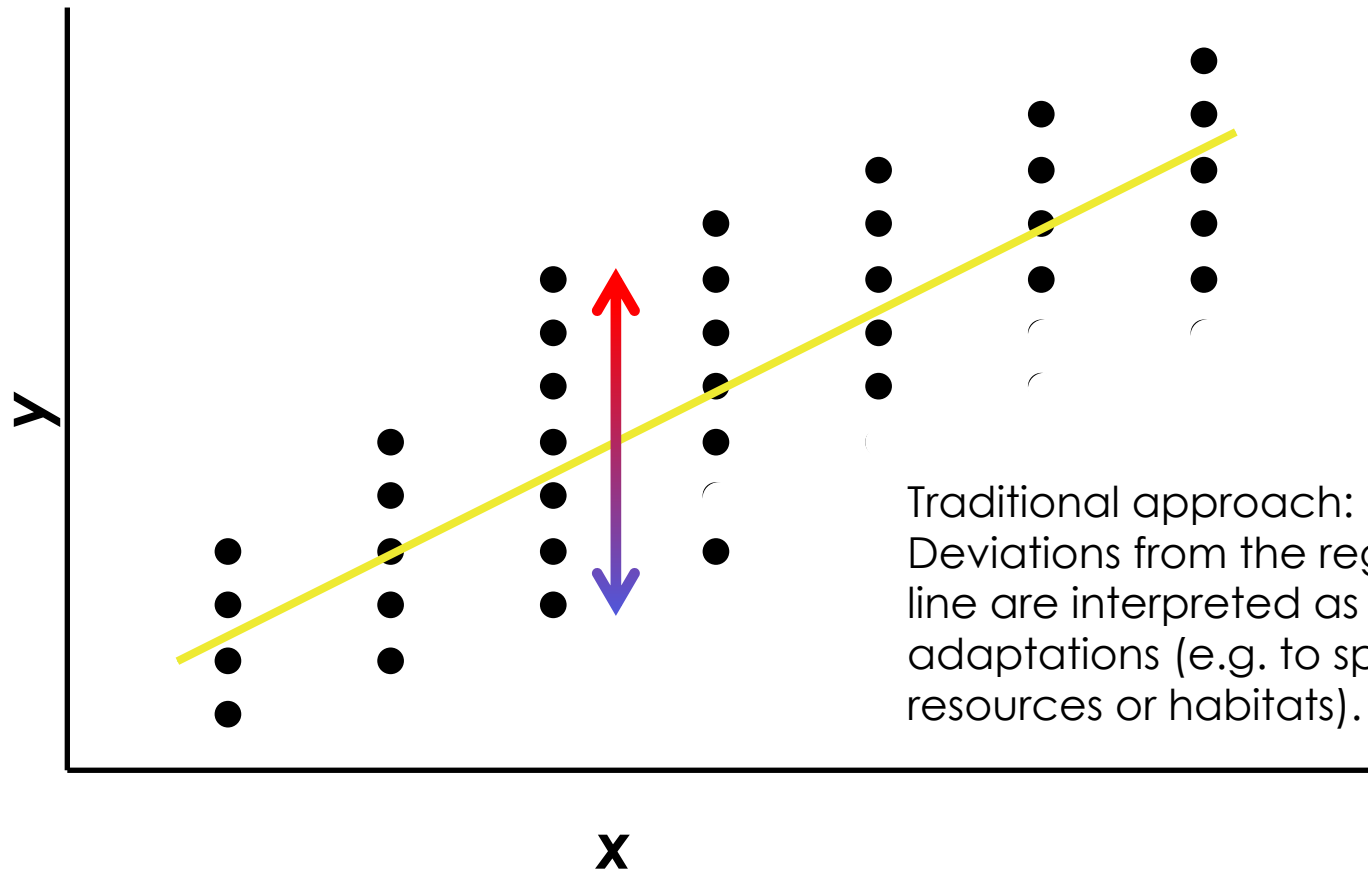
Morphological, physiological and life history variables scale.





Interpreting scaling

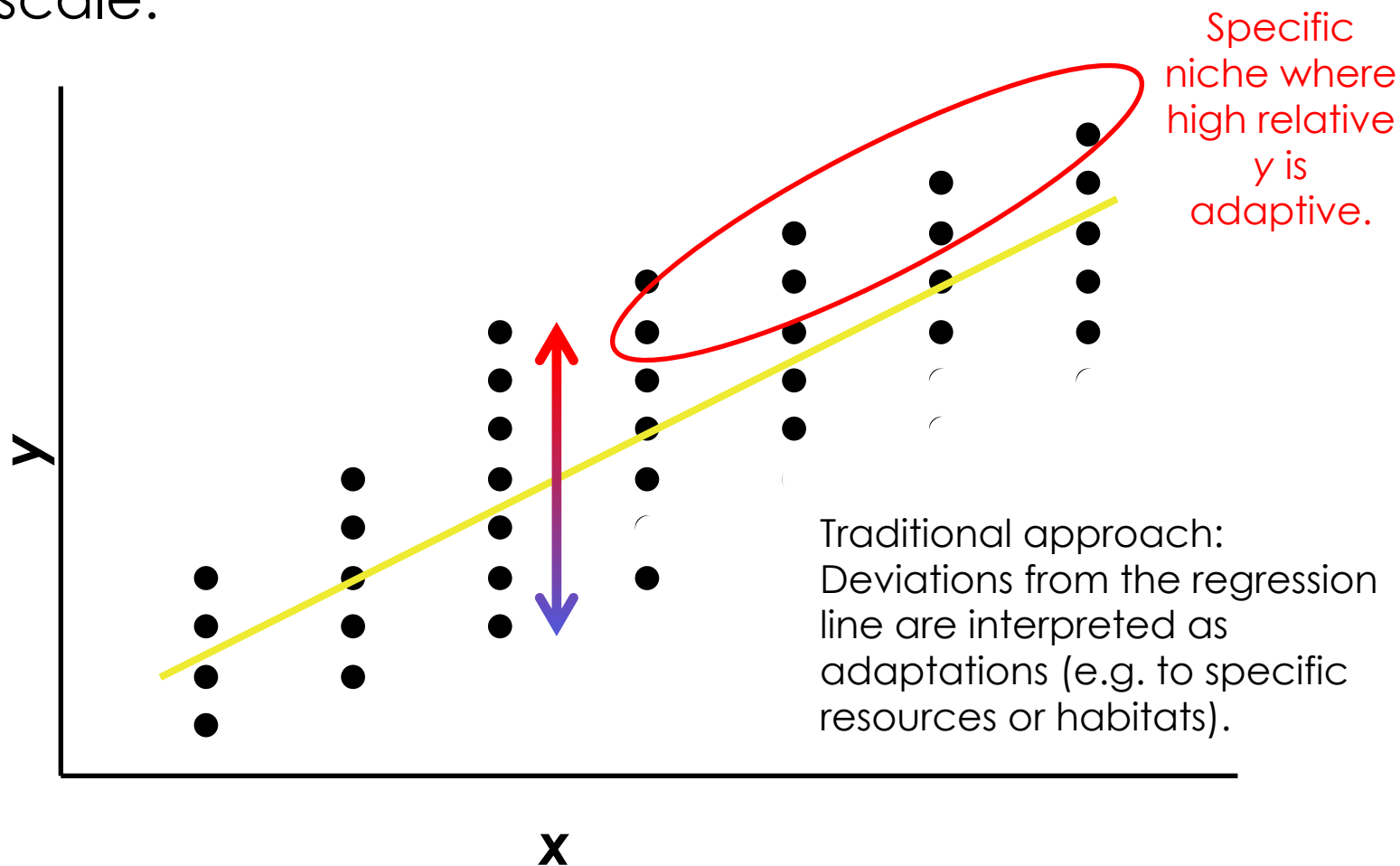
Morphological, physiological and life history variables scale.





Interpreting scaling

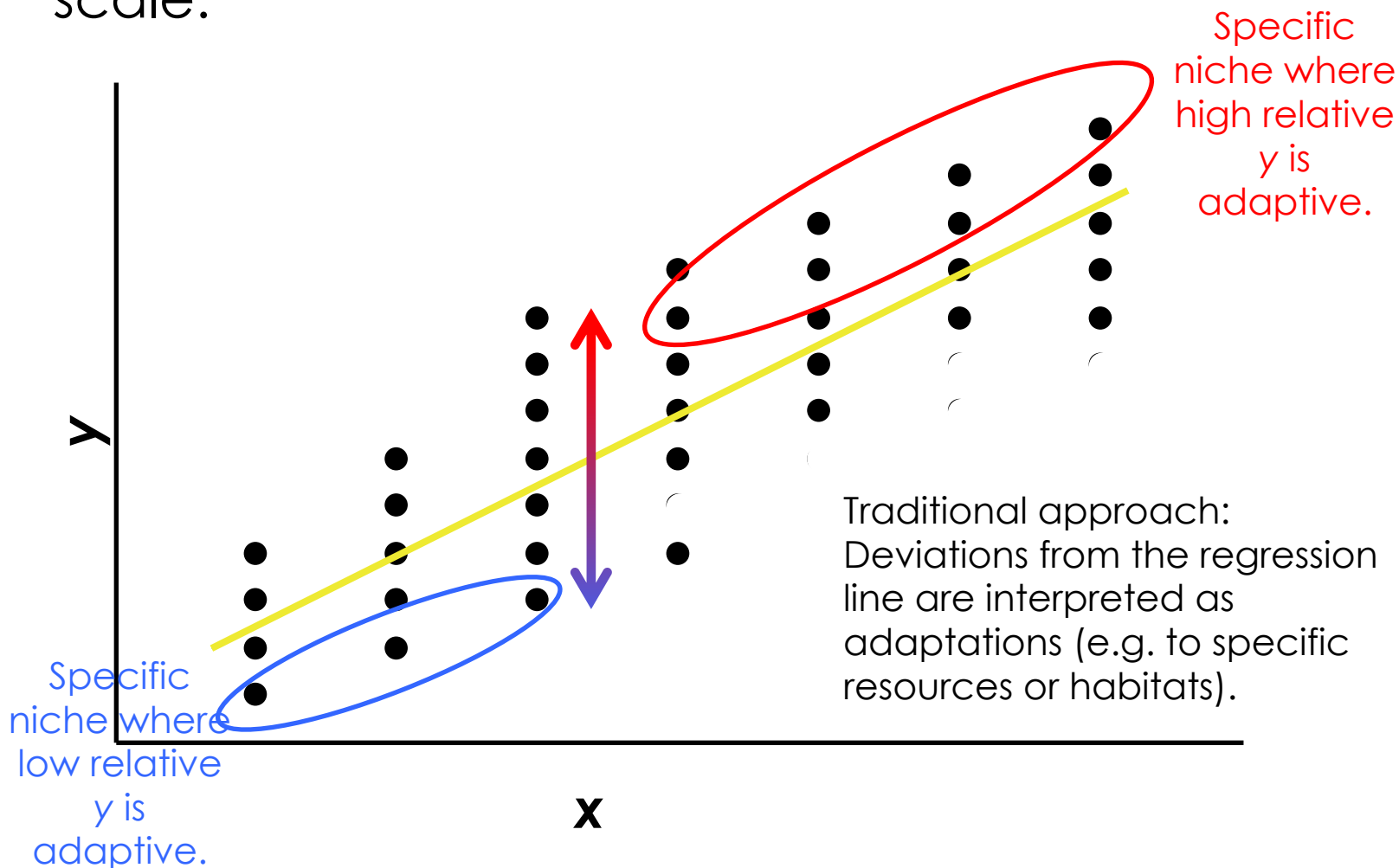
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Interpreting scaling

Morphological, physiological and life history variables scale.



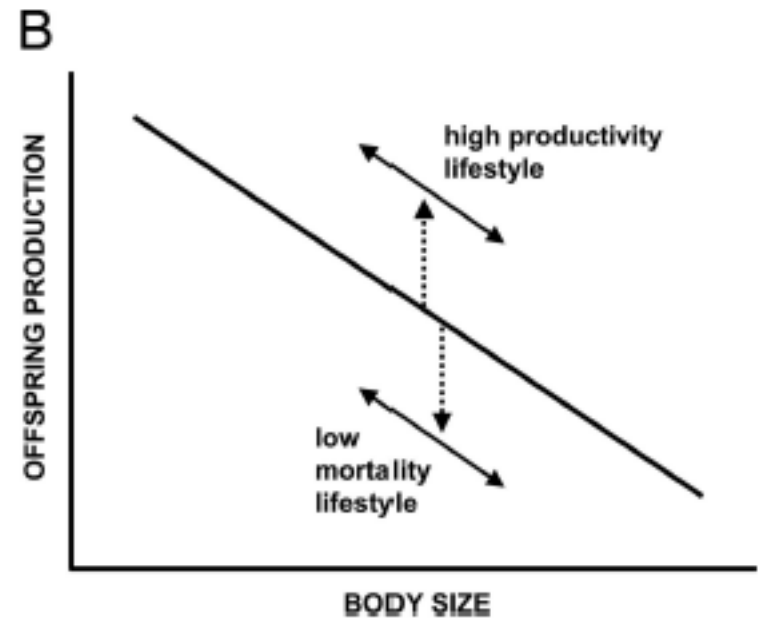
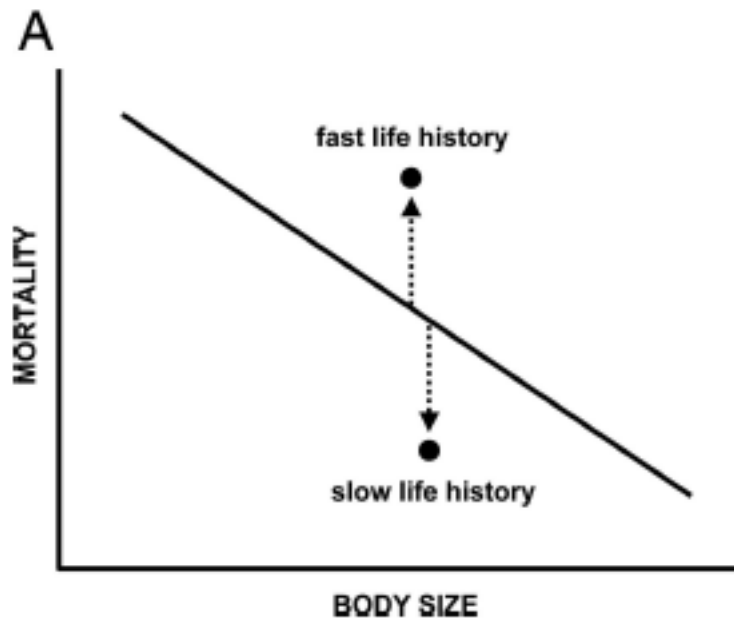


Life history scaling

A lifestyle view of life-history evolution

F. Stephen Dobson*

PNAS | November 6, 2007 | vol. 104 | no. 45 | 17565–17566





Effects of body size and lifestyle on evolution of mammal life histories

Richard M. Sibly^{*†‡} and James H. Brown^{‡§¶}

PNAS | November 6, 2007 | vol. 104 | no. 45 | 17707–17712

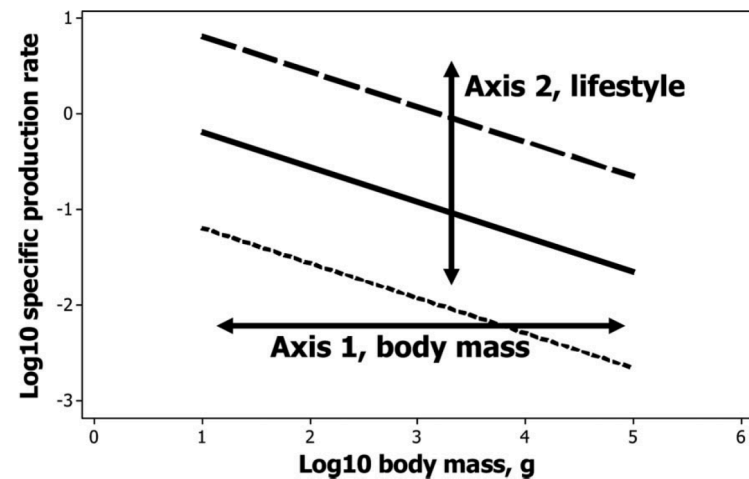
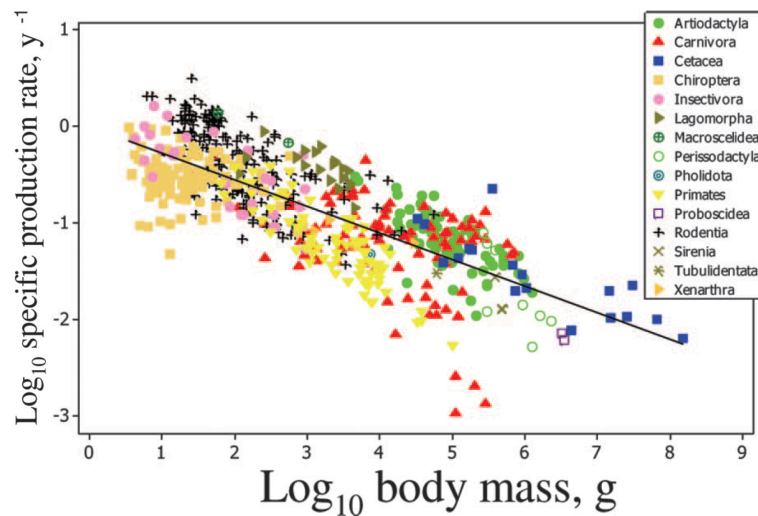
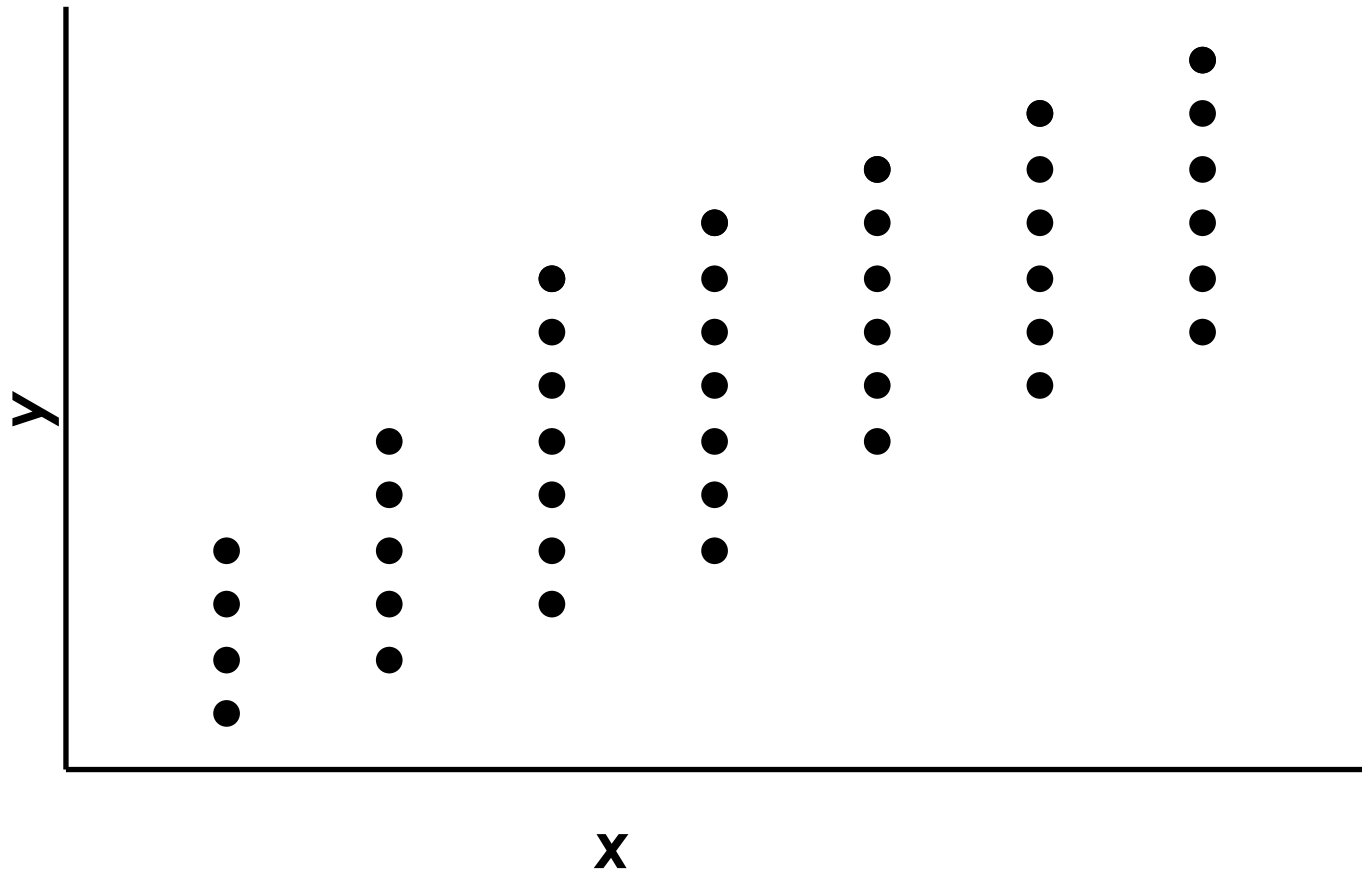


Fig. 4. The two major axes of the slow-fast life-history continuum, body mass, and lifestyle. To the well known axis of allometric variation due to body size, we have added a second orthogonal axis based on ecological lifestyle. Here the solid line represents an unspecialized ancestral condition, the dashed line depicts a more productive “live fast die young” lifestyle, and the dotted line shows a lifestyle with a lower death rate, slower life history, and consequently lower production.



Interpreting scaling

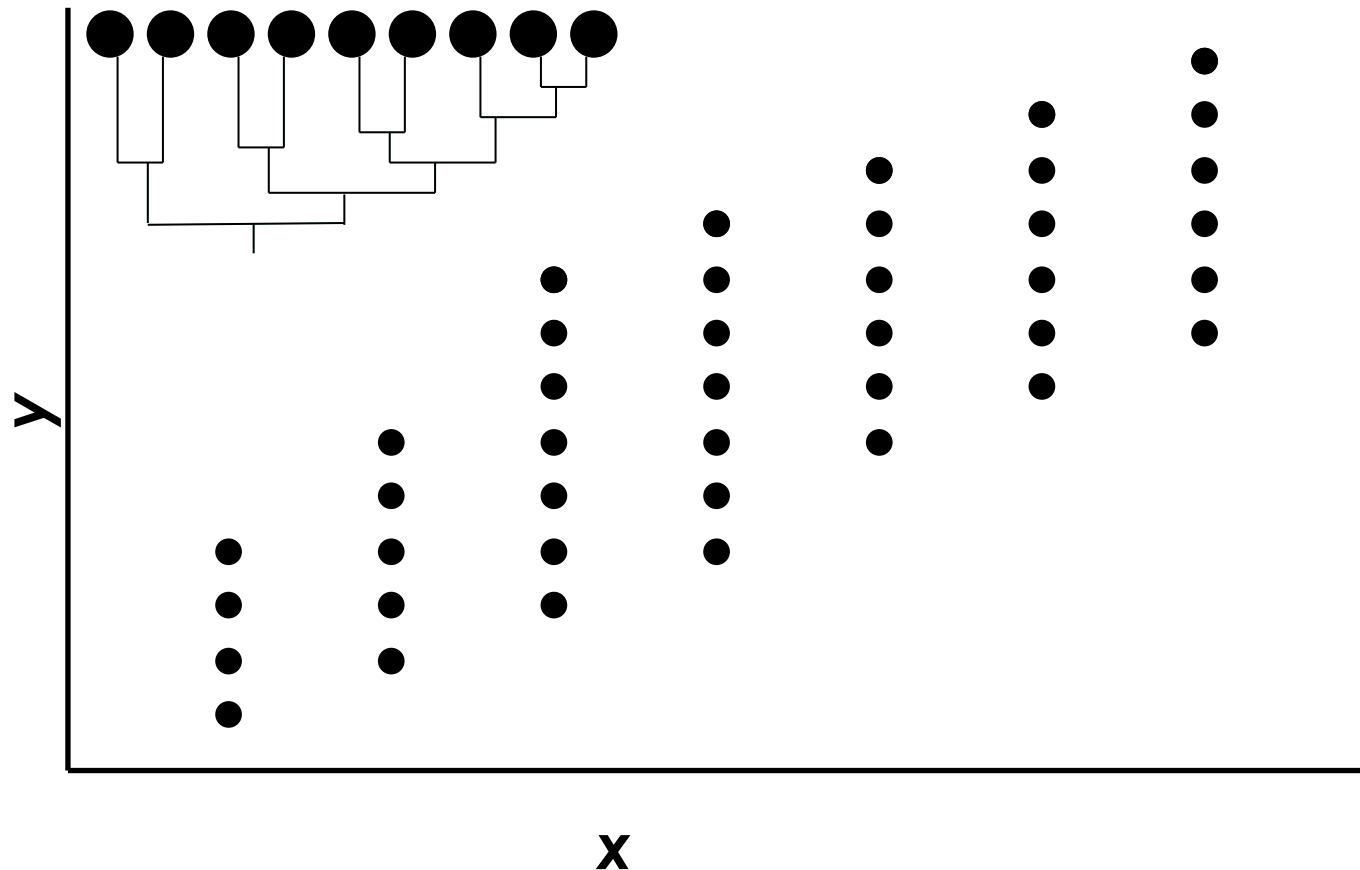
Is there a systematic phylogenetic structure in the dataset?





Interpreting scaling

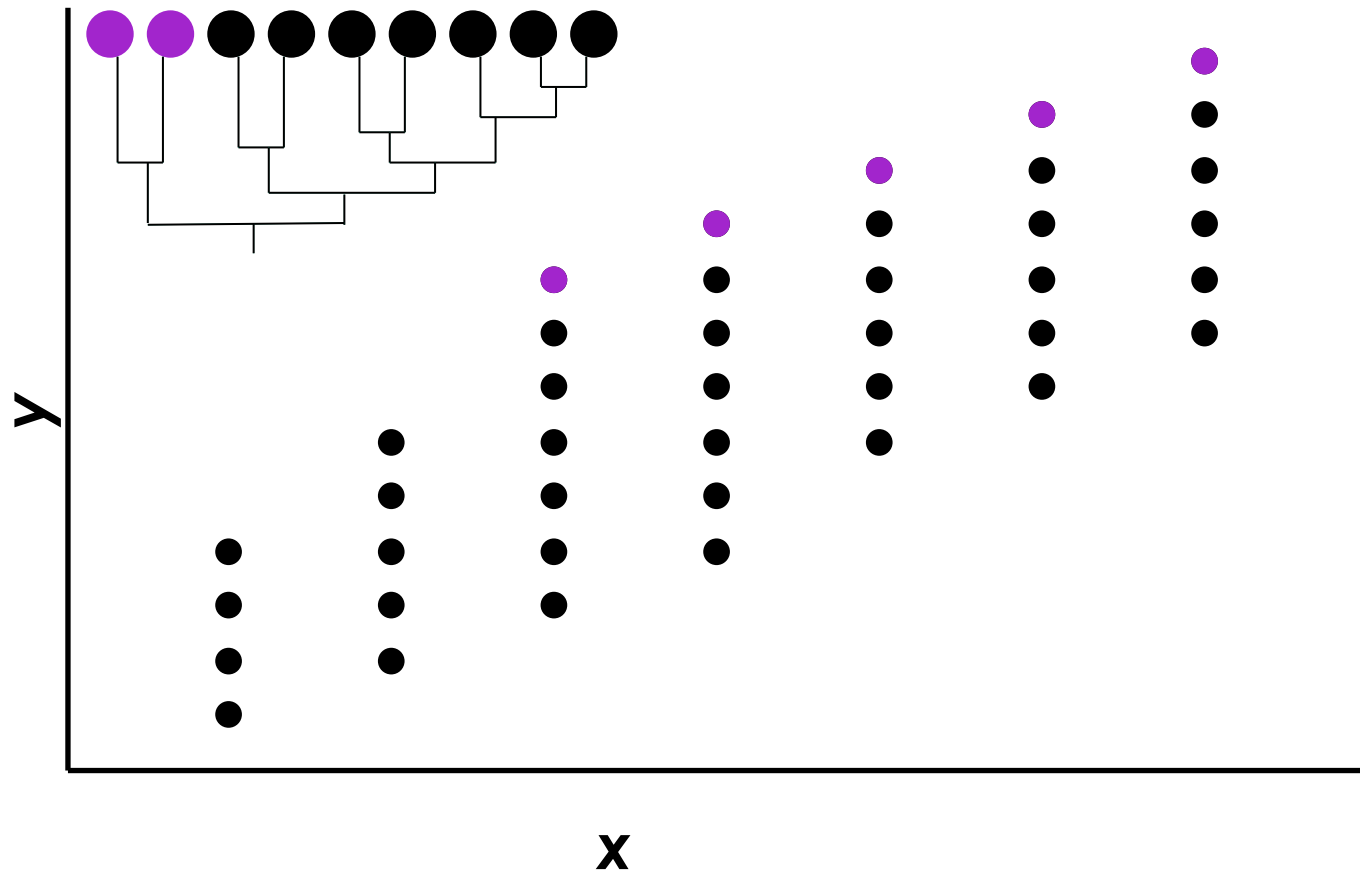
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Interpreting scaling

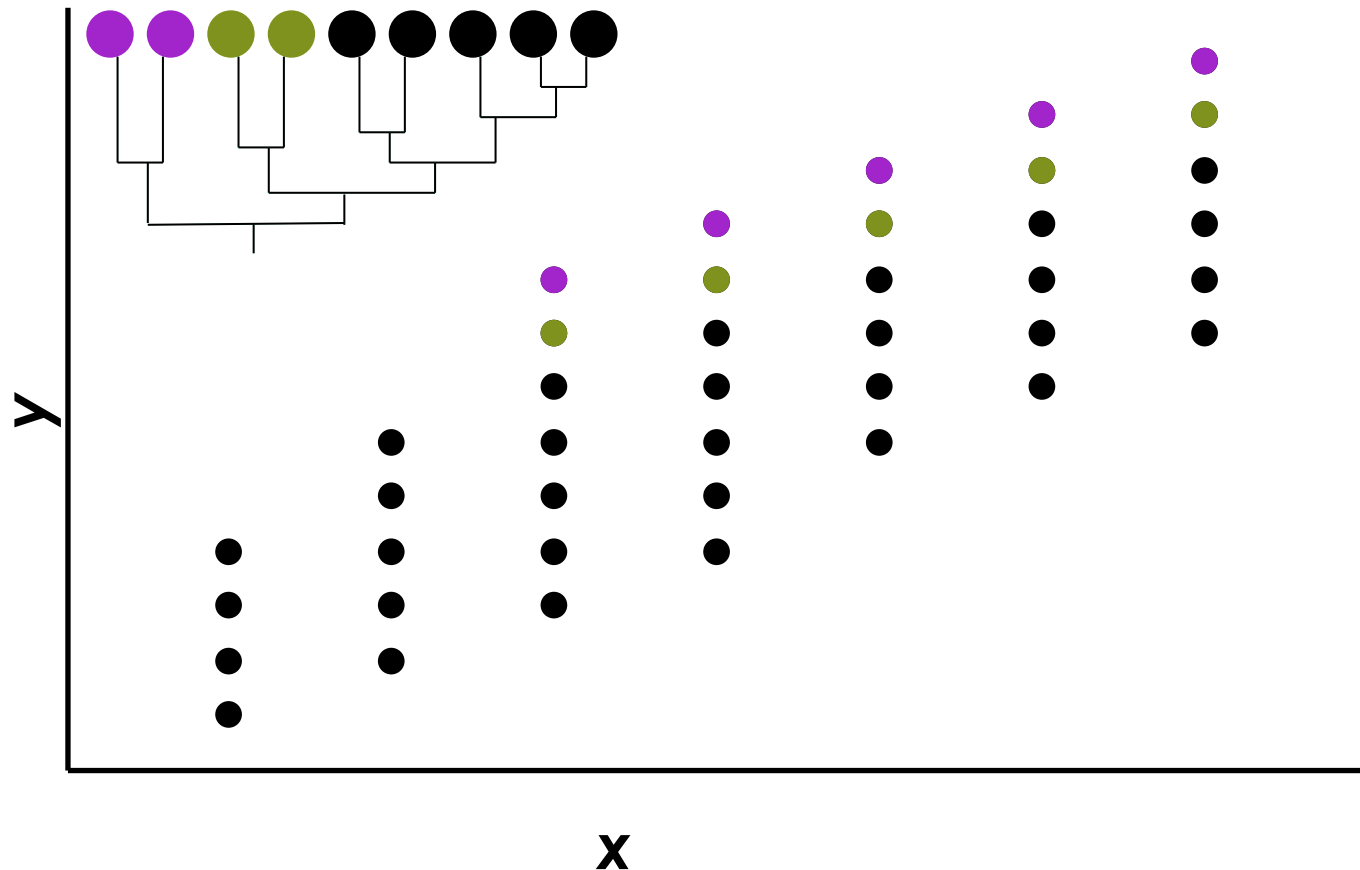
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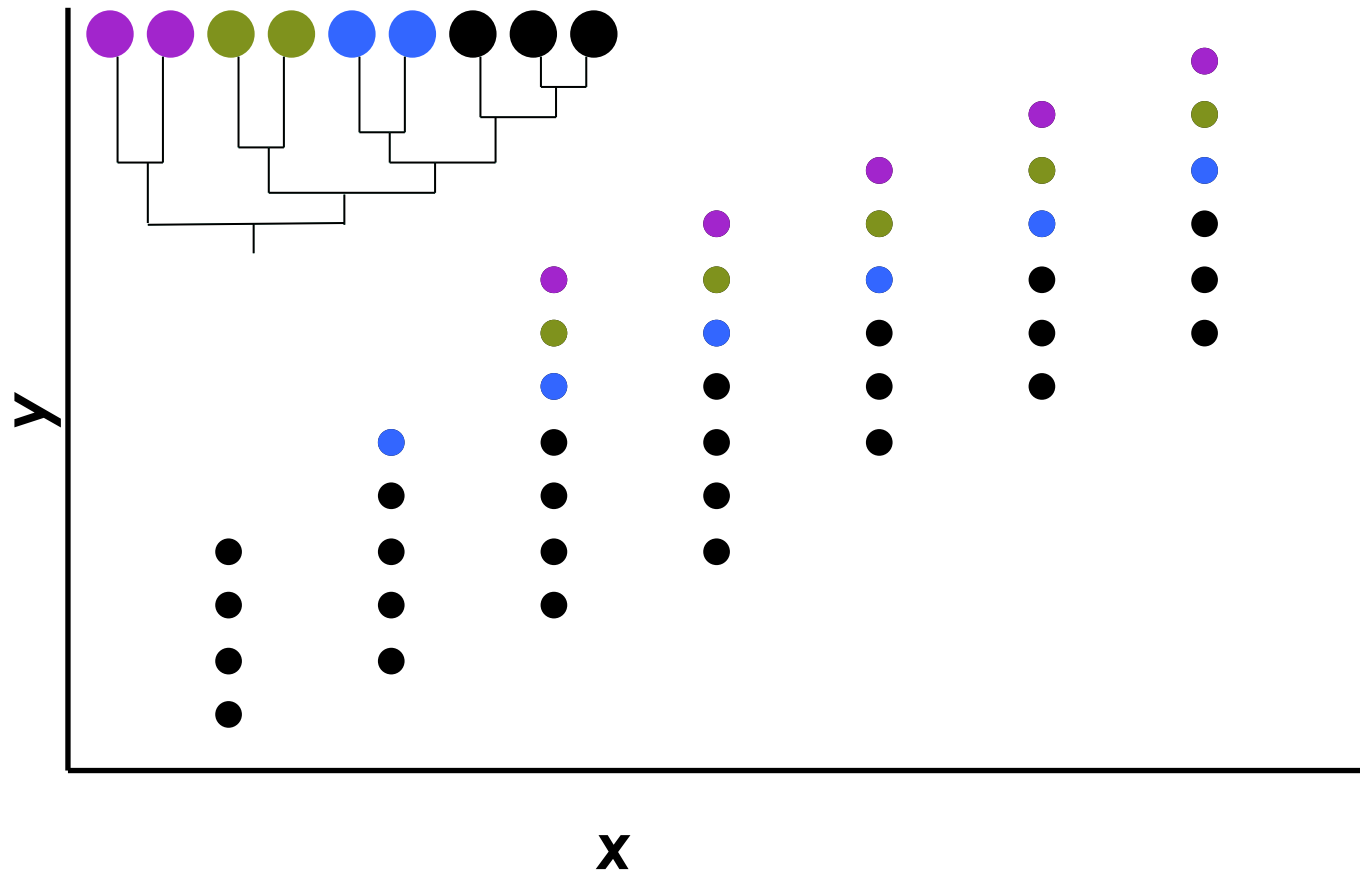
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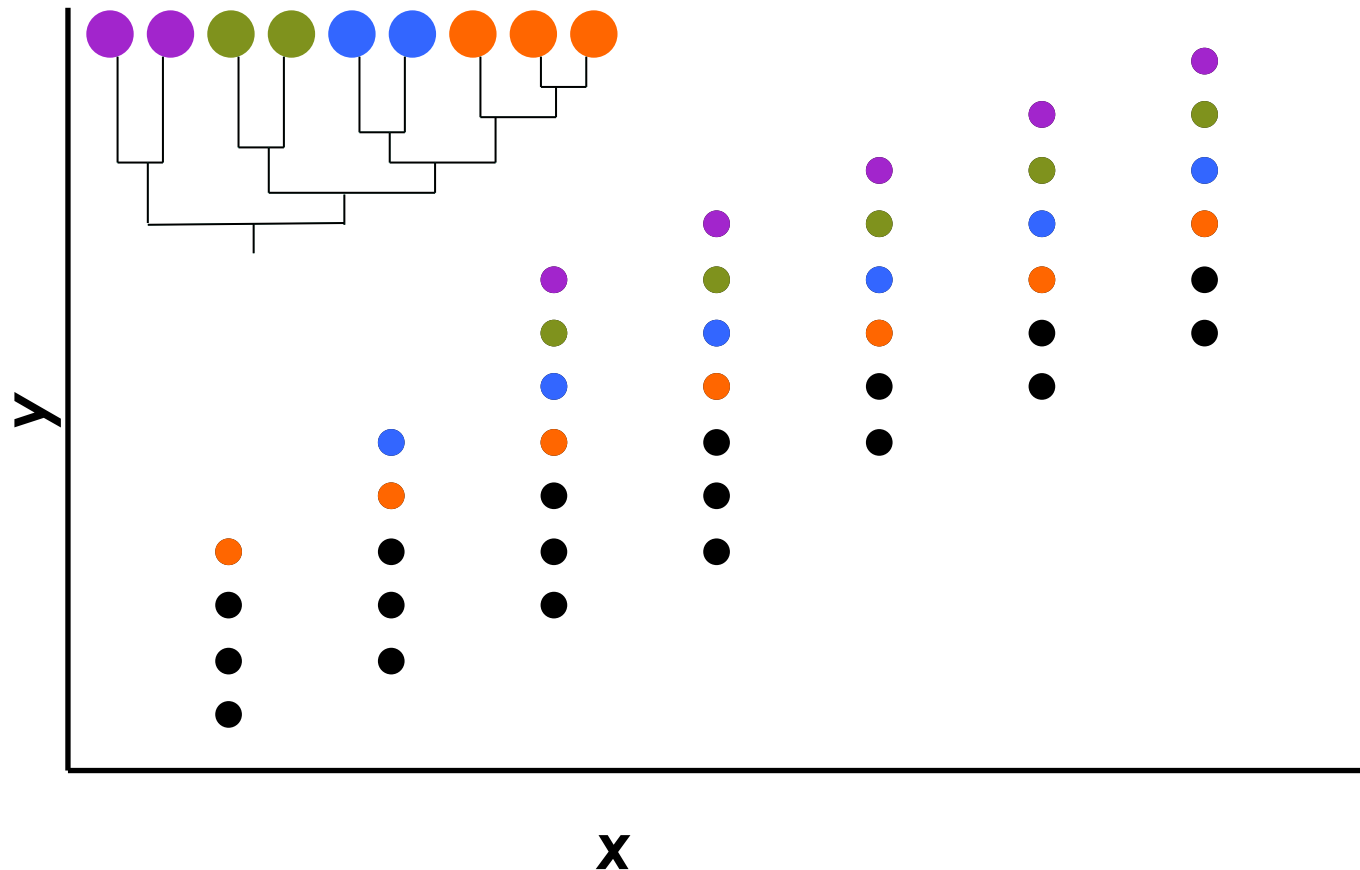
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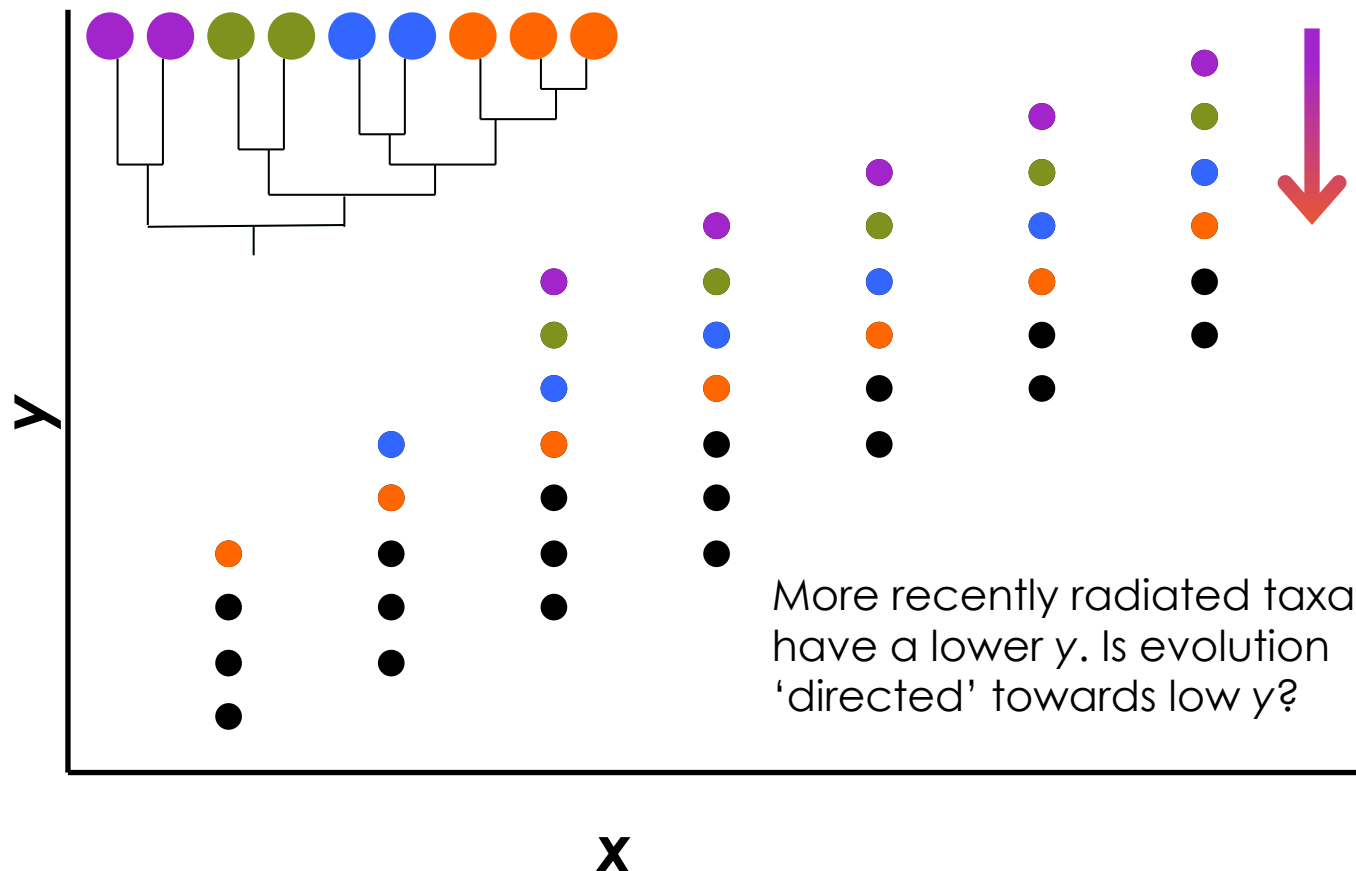
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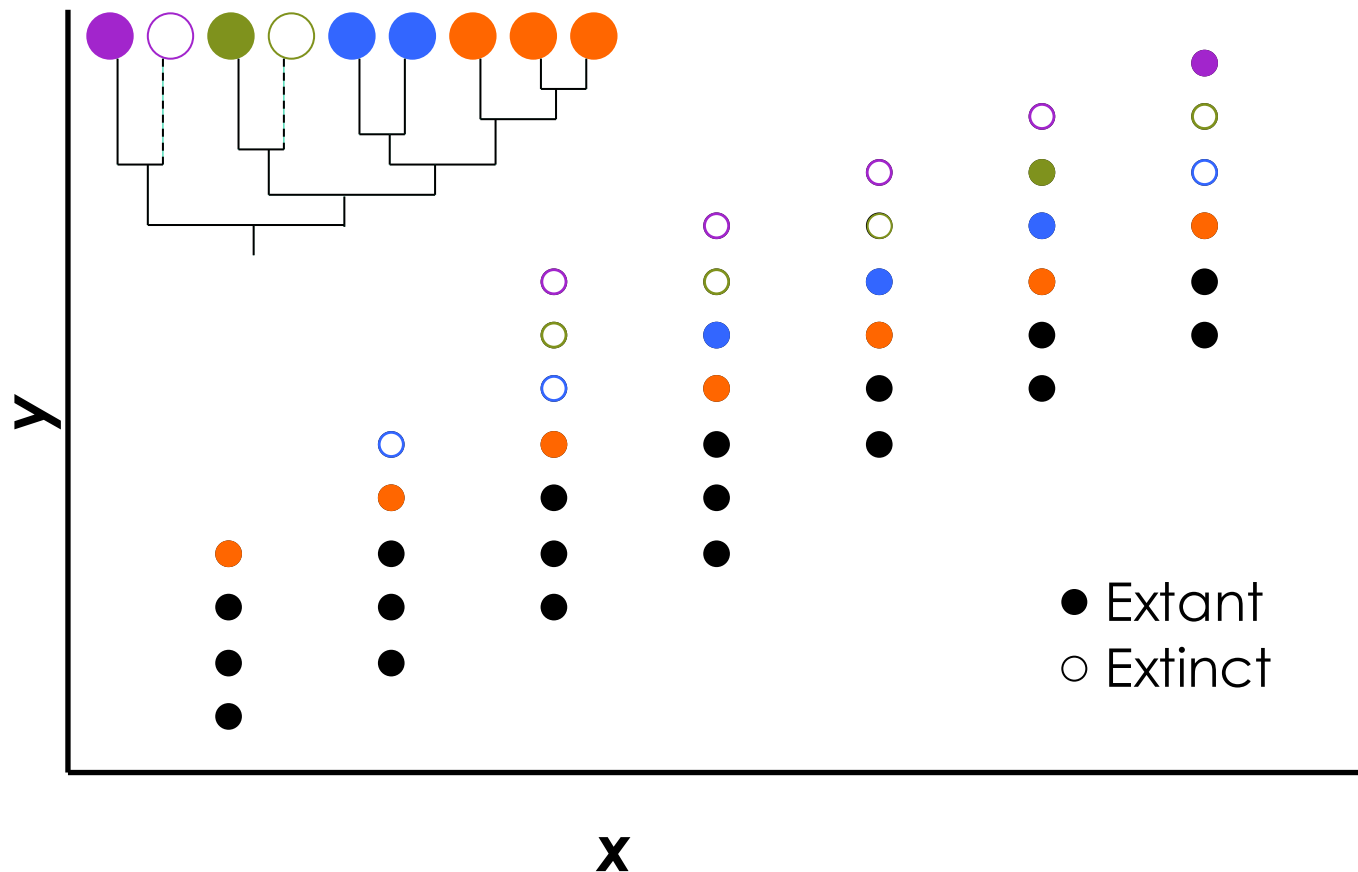
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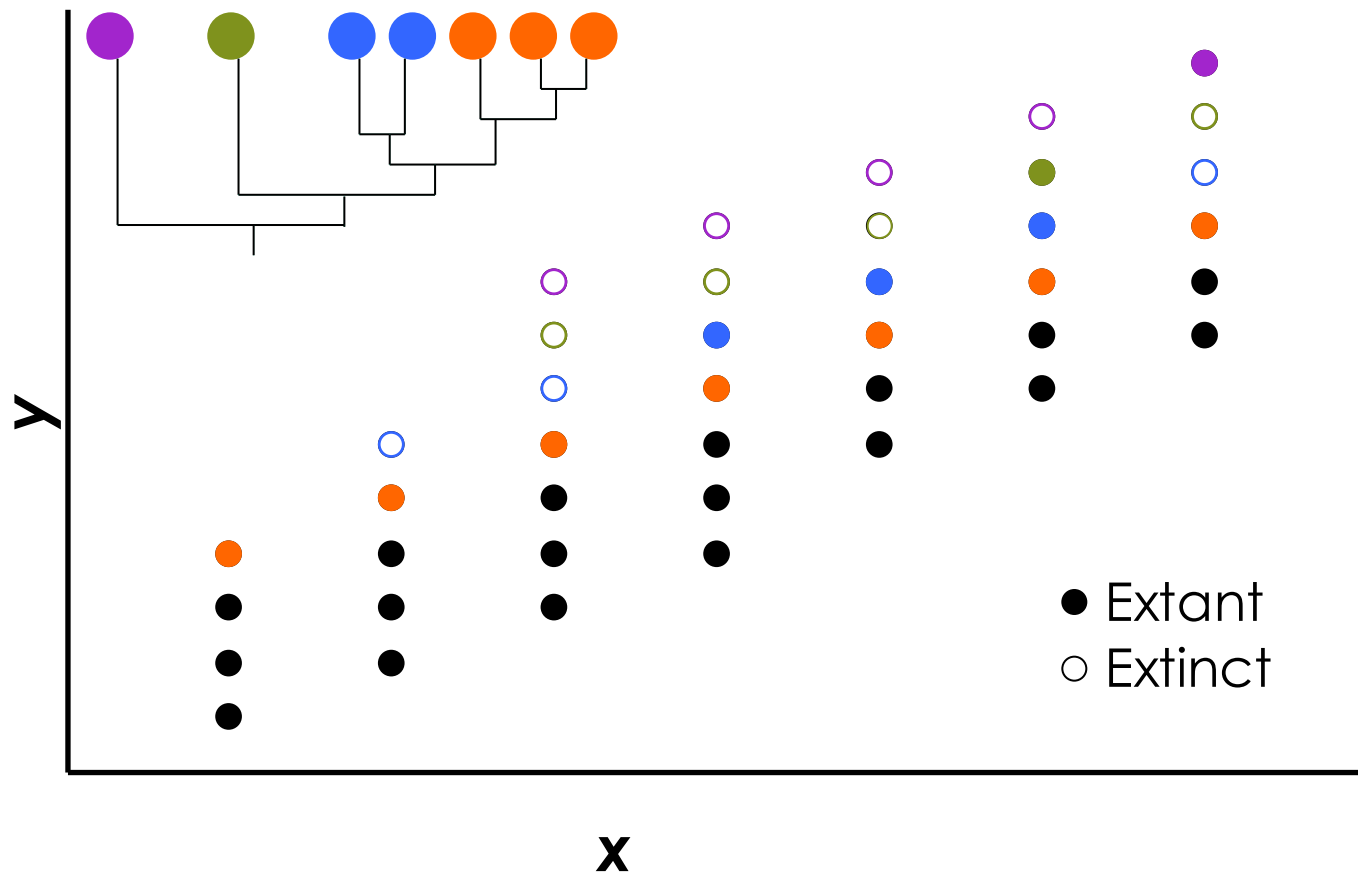
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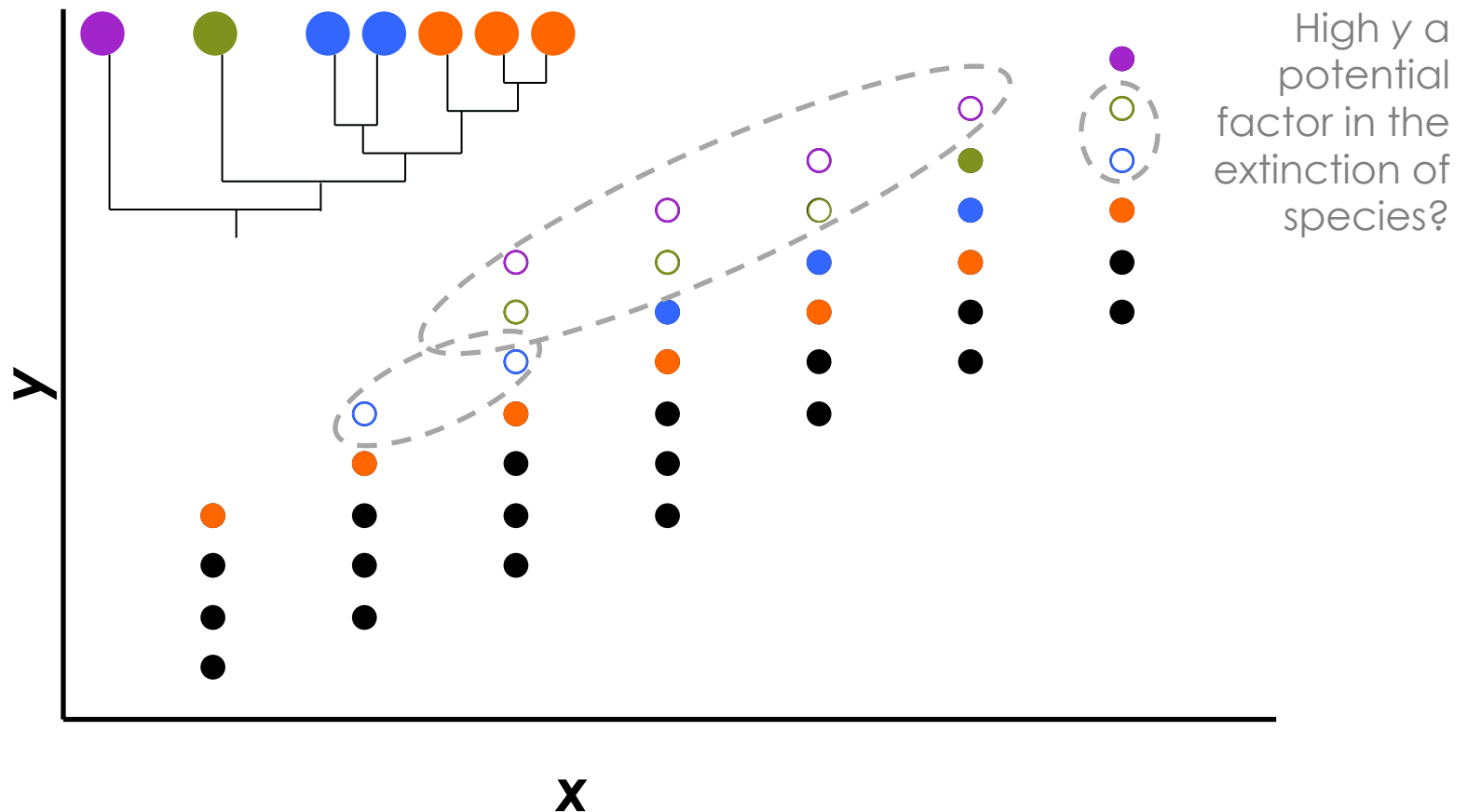
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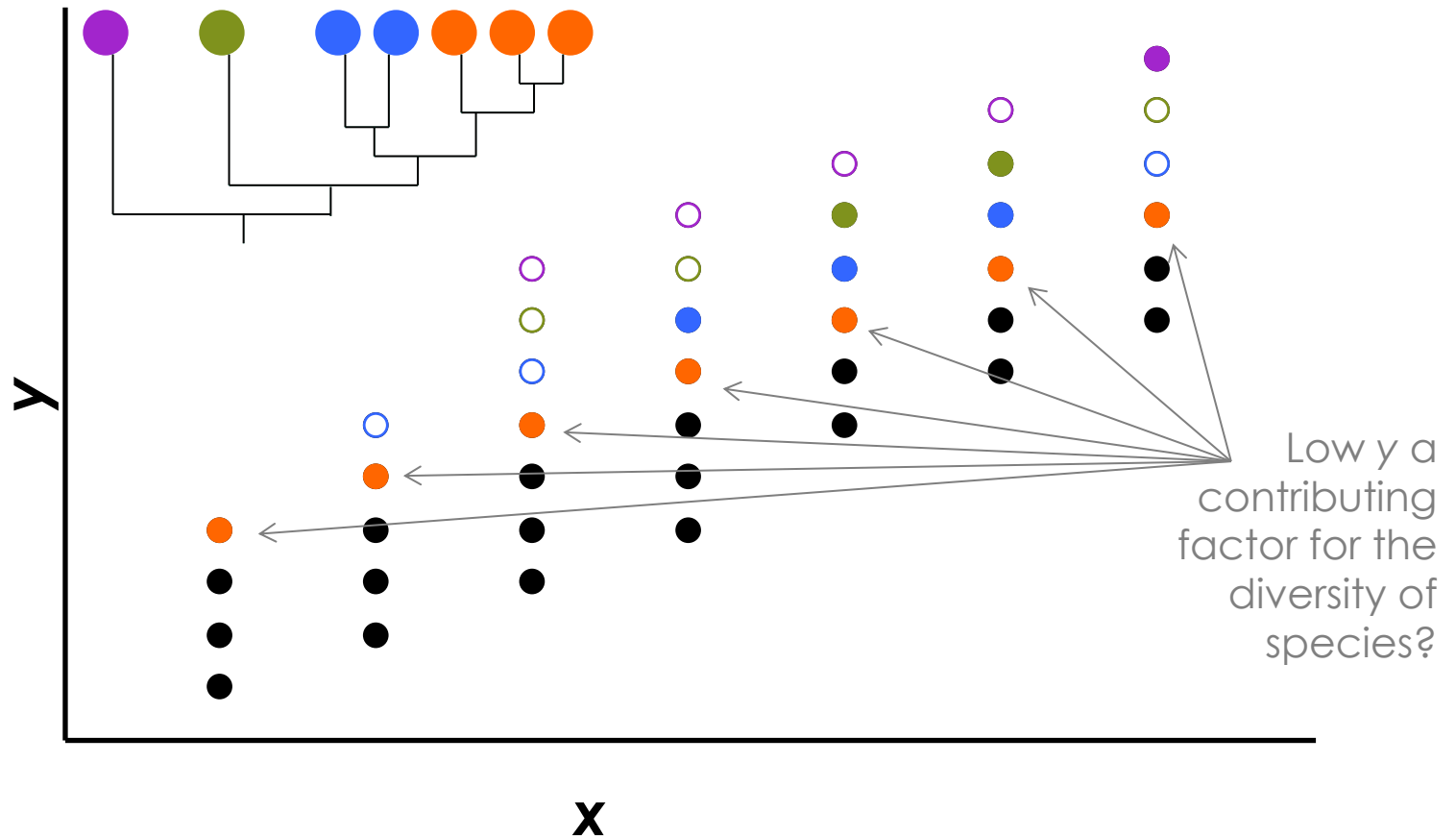
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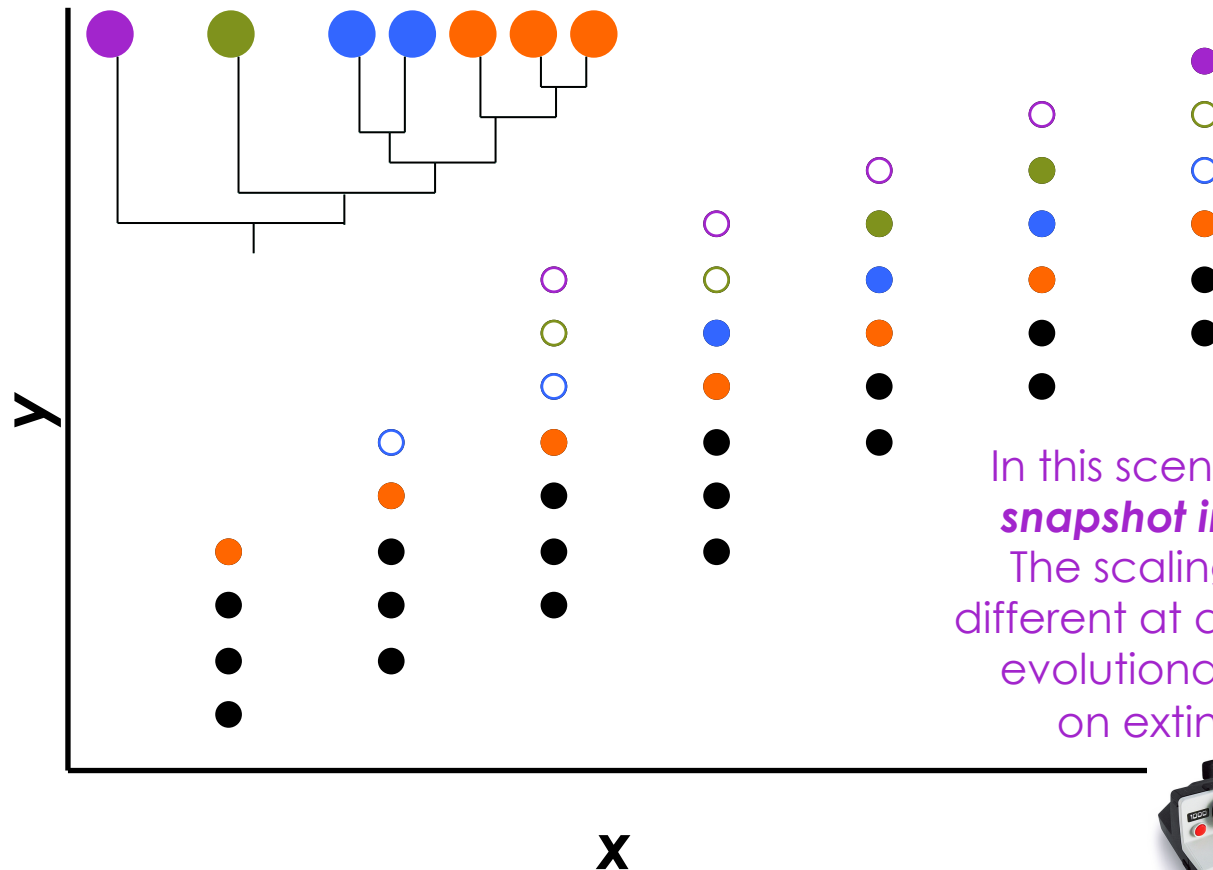
Is there a systematic phylogenetic structure in the dataset?





Interpreting scaling: snapshots

Is there a systematic phylogenetic structure in the dataset?

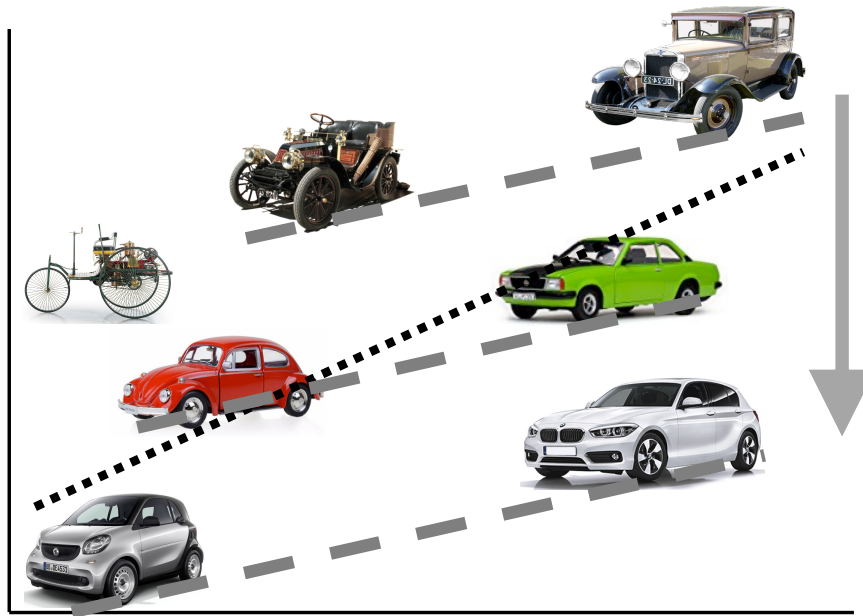


In this scenario, the scaling is a **snapshot in evolutionary time**. The scaling would have been different at different moments in evolutionary time (depending on extinction and radiation events)





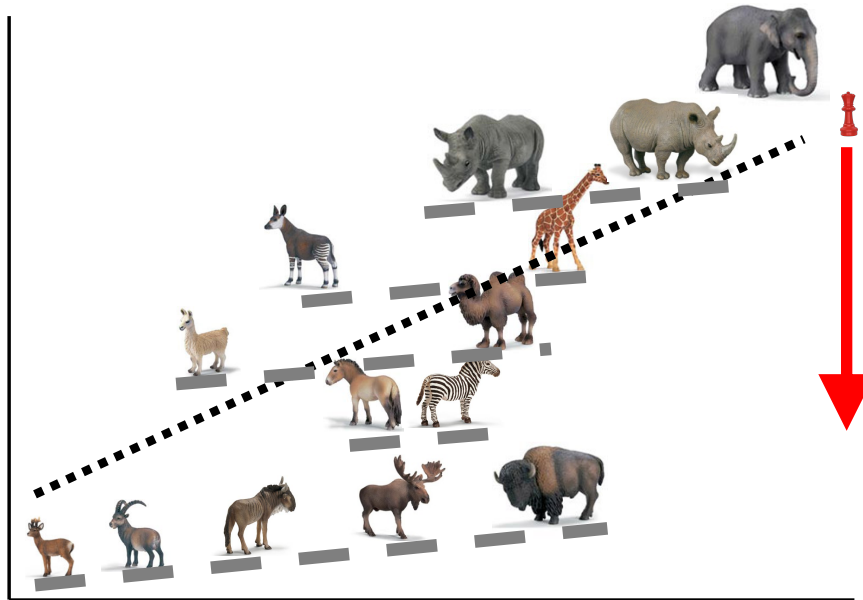
Energy per km



Mass

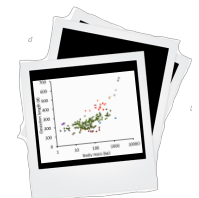
You would not consider the overall pattern a fixed law, but consider it with respect to technical progress.

Time per offspring



Mass

Why would you consider this a pattern due to fixed life history tradeoff laws, and not rather a **snapshot** in a process of optimization?





Assessing 'direction' / *Red Queen* / escalation / progress in life history

using the PanTheria dataset
(Jones et al. 2009)



Niche-specific assessment

Because niche space is less diverse at larger body sizes, large herbivores may be a particularly fruitful area of research for 'directed evolution'.



Herbivore
basic™



Herbivore
2.0™



Herbivore
professional™



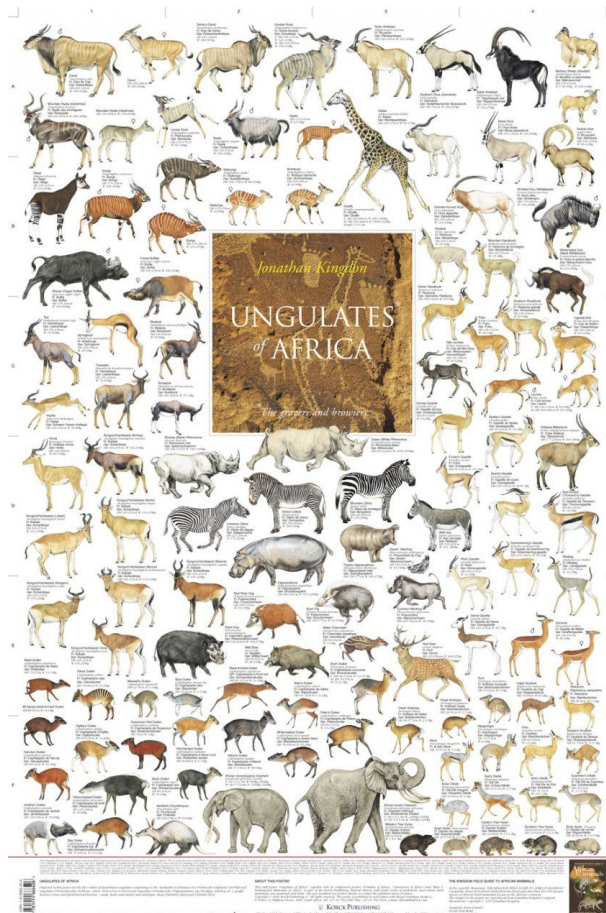
Herbivore
ultimate™





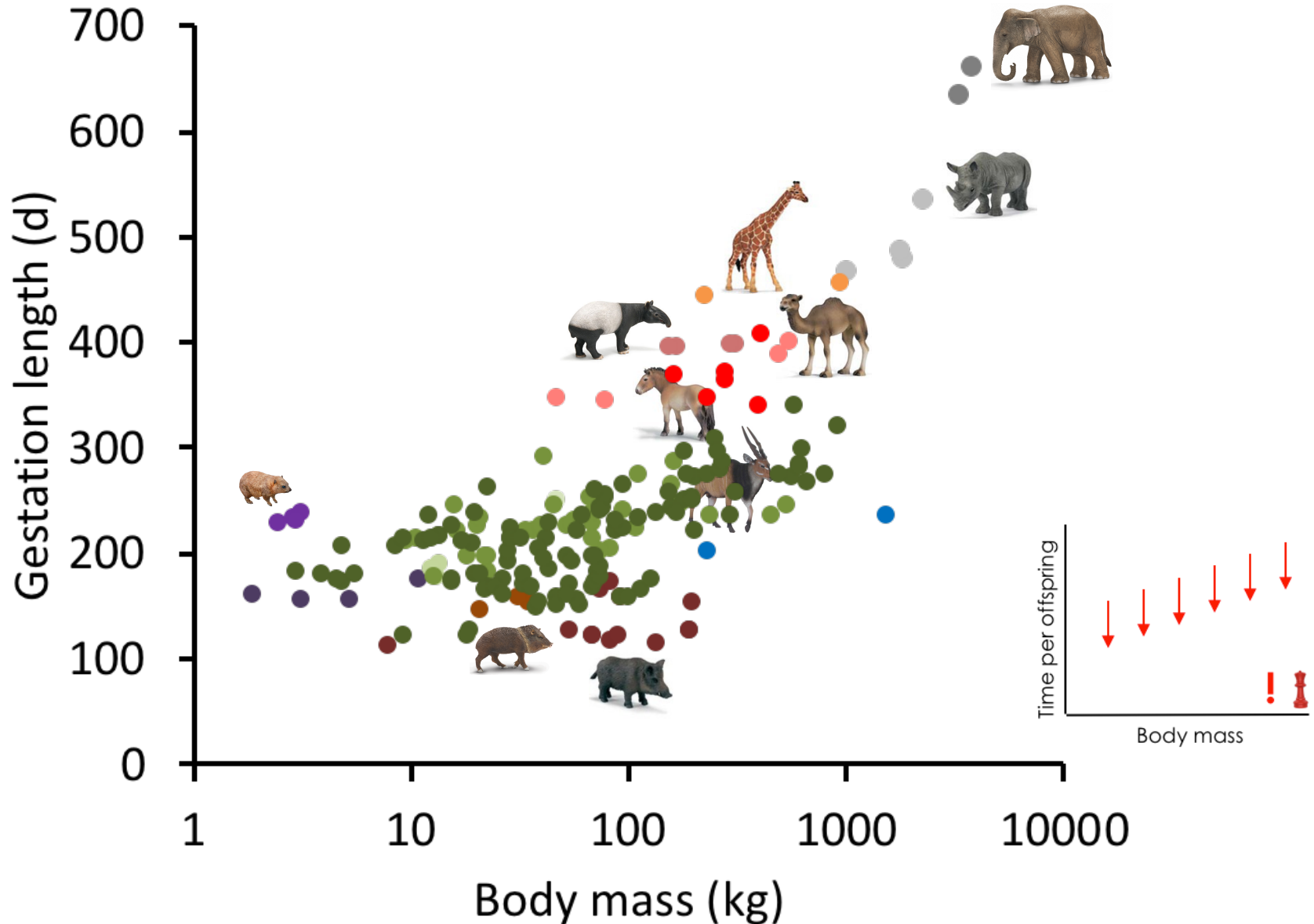
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A clear picture for gestation length





(Precocial) Mammal gestation period

For any mammal, achieving the same degree of neonatal development in a shorter gestation period – if not associated with higher costs – should be advantageous (higher fecundity due to shorter generation times).

Days of gestation period (to apparently similar level of precociality)

Cattle:	app. 280 days
Horse:	app. 340 days
Dromedary:	app. 390 days
Okapi:	app. 440 days



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The difference cannot be due to body size!



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nearly extinct in a
very limited
geographical range



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only in extreme,
resource-poor
habitats



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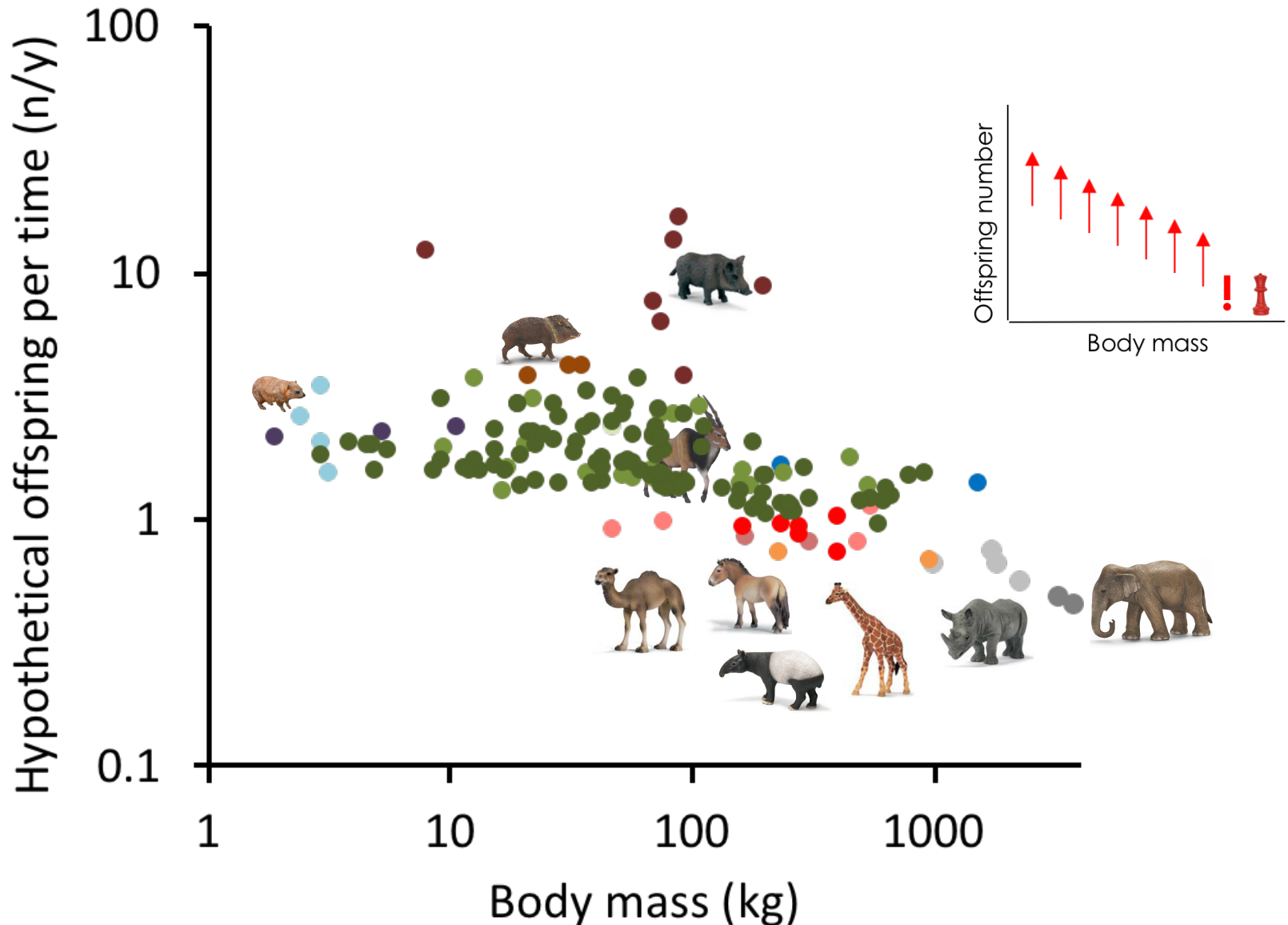
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rule the world !!

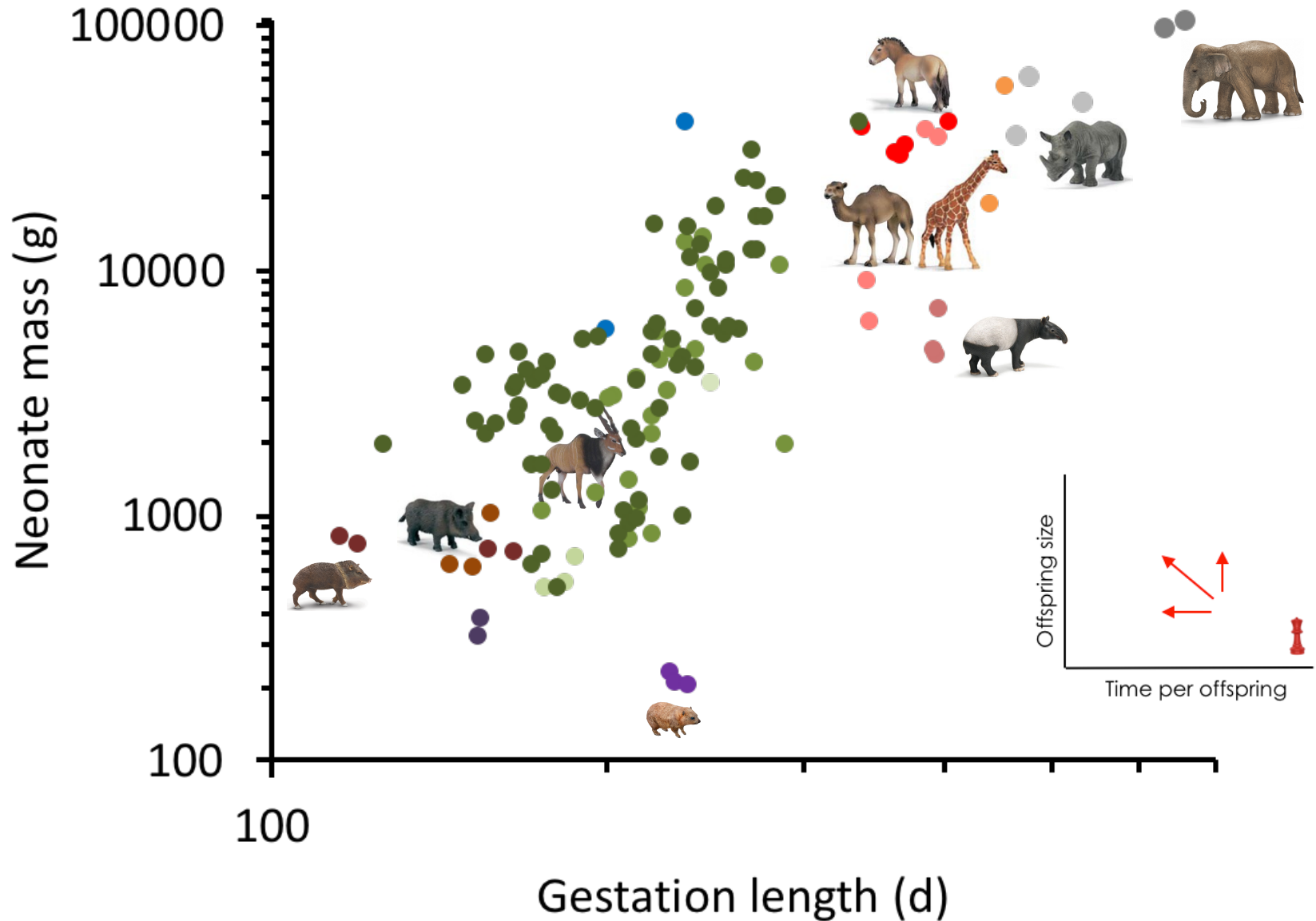


Clear effect for yearly offspring



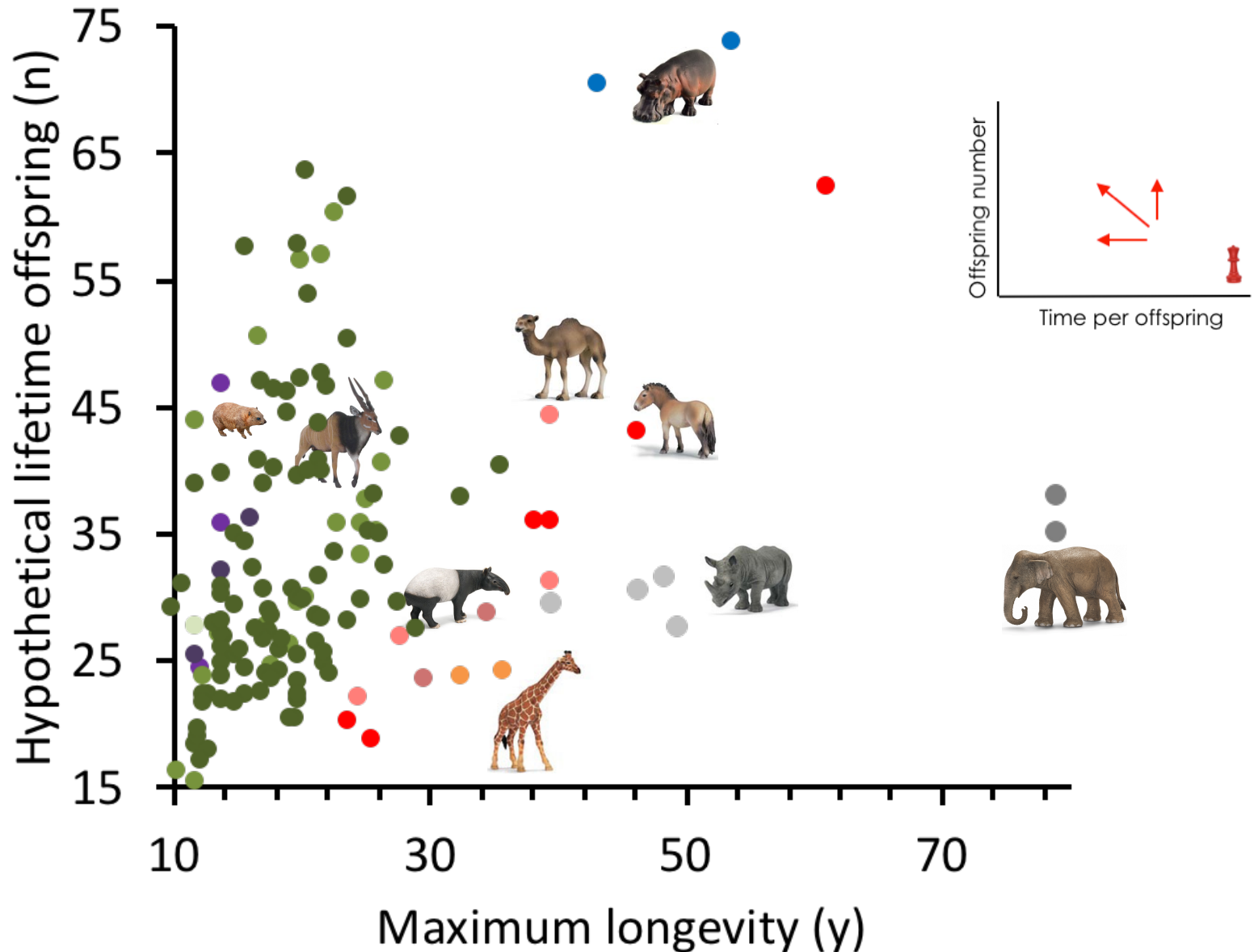


A clear picture for intrauterine growth





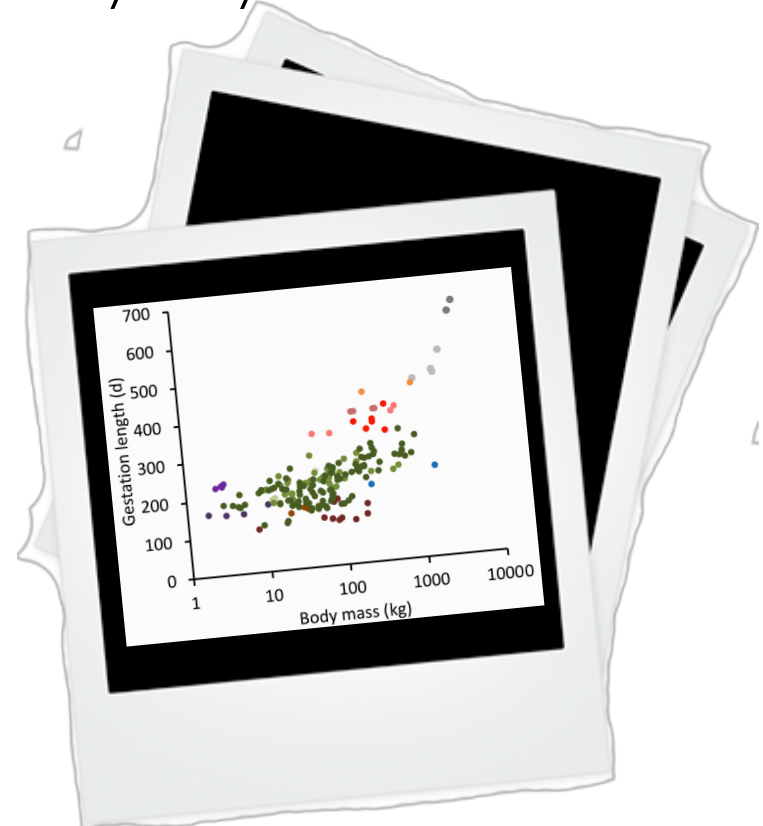
A clear picture for lifetime offspring production





Summary, Conclusions & Outlook

Rather than understanding tradeoffs along the fast-slow continuum as fixed physical laws, they can be considered as representing the efficiency of the organisms from which the data was taken – and that efficiency may evolve.

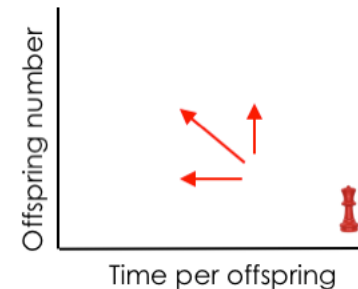




Summary, Conclusions & Outlook

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Within the boundaries of a specific niche, species possibly compete by demographic means: by evolving a faster reproduction.



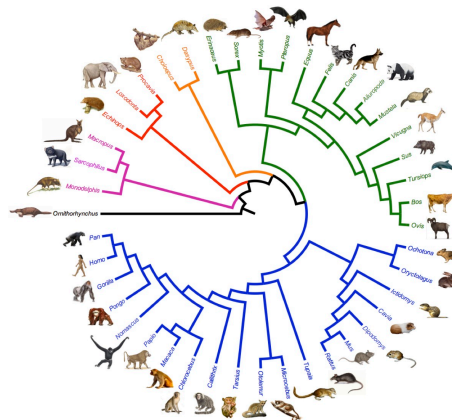


Summary, Conclusions & Outlook

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Within the boundaries of a specific niche, species possibly compete by demographic means: by evolving a faster reproduction.

Life history characteristics appear to be linked to taxonomic groups.





The interesting question ...

... what allowed the remaining extant species of the 'slower' taxa to survive?





Summary, Conclusions & Outlook

Rather than understanding tradeoffs along the fast-slow continuum as fixed physical laws, they can be considered as representing the efficiency of the organisms from which the data was taken – and that efficiency may evolve.

Within the boundaries of a specific niche, species possibly compete by demographic means: by evolving a faster reproduction.

Life history characteristics appear to be linked to taxonomic groups.

We would predict that during earth history, 'faster' species were not replaced by 'slower' species.

Application: large herbivore diversity through time

Historical Biology, 1994, Vol. 8, pp. 15–29
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MODELLING EQUID/RUMINANT COMPETITION IN THE FOSSIL RECORD

CHRISTINE M. JANIS¹, IAIN J. GORDON² and ANDREW W. ILLIUS³

¹*Department of Ecology and Evolutionary Biology, Brown University, Providence, Rhode Island 02912, USA*

²*Macaulay Land Use Research Institute, Craigiebuckler, Aberdeen AB9 2QY, UK*

³*School of Agriculture, University of Edinburgh, West Mains Road, Edinburgh EH9, 3JG, UK*

(Received November 2, 1993)

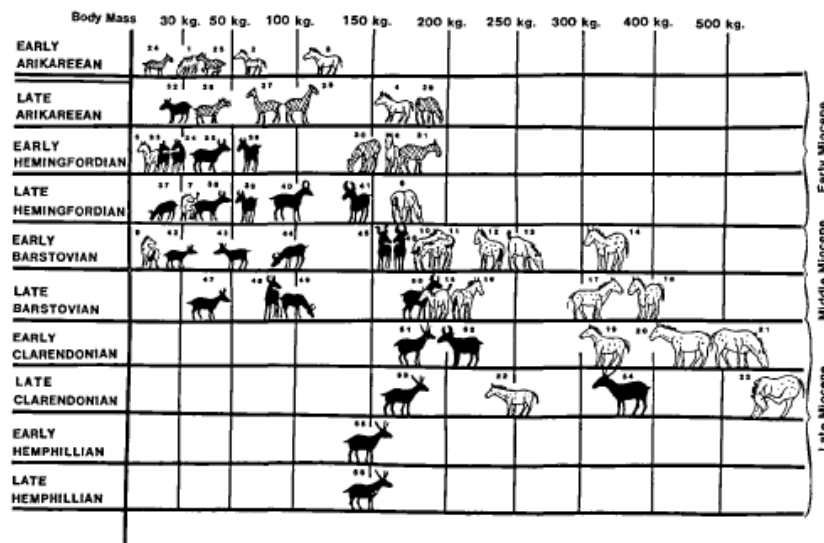


Figure 3 Body size distribution of browsing Miocene equids and ruminants. Key to ungulate taxa: A. Color of taxon: Striped horses=Mesohippines; white horses=Anchitherines; spotted horses=Hypohippines; black artiodactyls=Pecorans; cross-hatched artiodactyls=Tylopods.



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CHRISTINE M. JANIS¹, IAIN J. GORDON² and ANDREW W. ILLIUS³

Historical Biology, 1994, Vol. 8, pp. 15–29

Body Mass	30kg.	50kg.	100kg.	150kg.	200kg.	250kg.	300kg.	400kg.	500kg.	
EARLY ARIKAREEAN										EARLY MIOCENE
LATE ARIKAREEAN										
EARLY HEMINGFORDIAN										
LATE HEMINGFORDIAN										
EARLY BARSTOVIAN										MIDDLE MIOCENE
LATE BARSTOVIAN										
EARLY CLARENDONIAN										LATE MIOCENE
LATE CLARENDONIAN										
EARLY HEMPHILLIAN										
LATE HEMPHILLIAN										

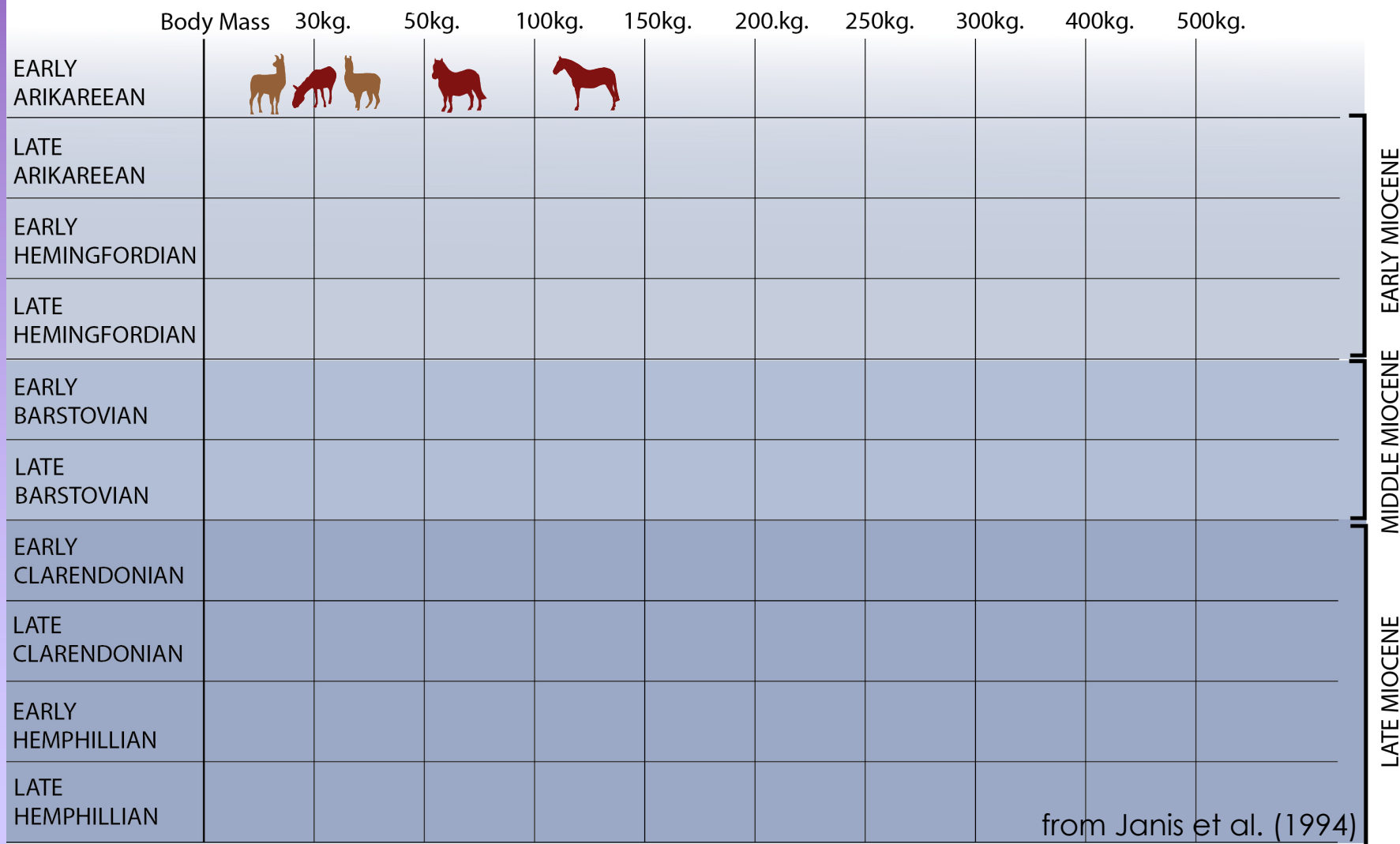
from Janis et al. (1994)



MODELLING EQUID/RUMINANT COMPETITION IN THE FOSSIL RECORD

CHRISTINE M. JANIS¹, IAIN J. GORDON² and ANDREW W. ILLIUS³

Historical Biology, 1994, Vol. 8, pp. 15–29

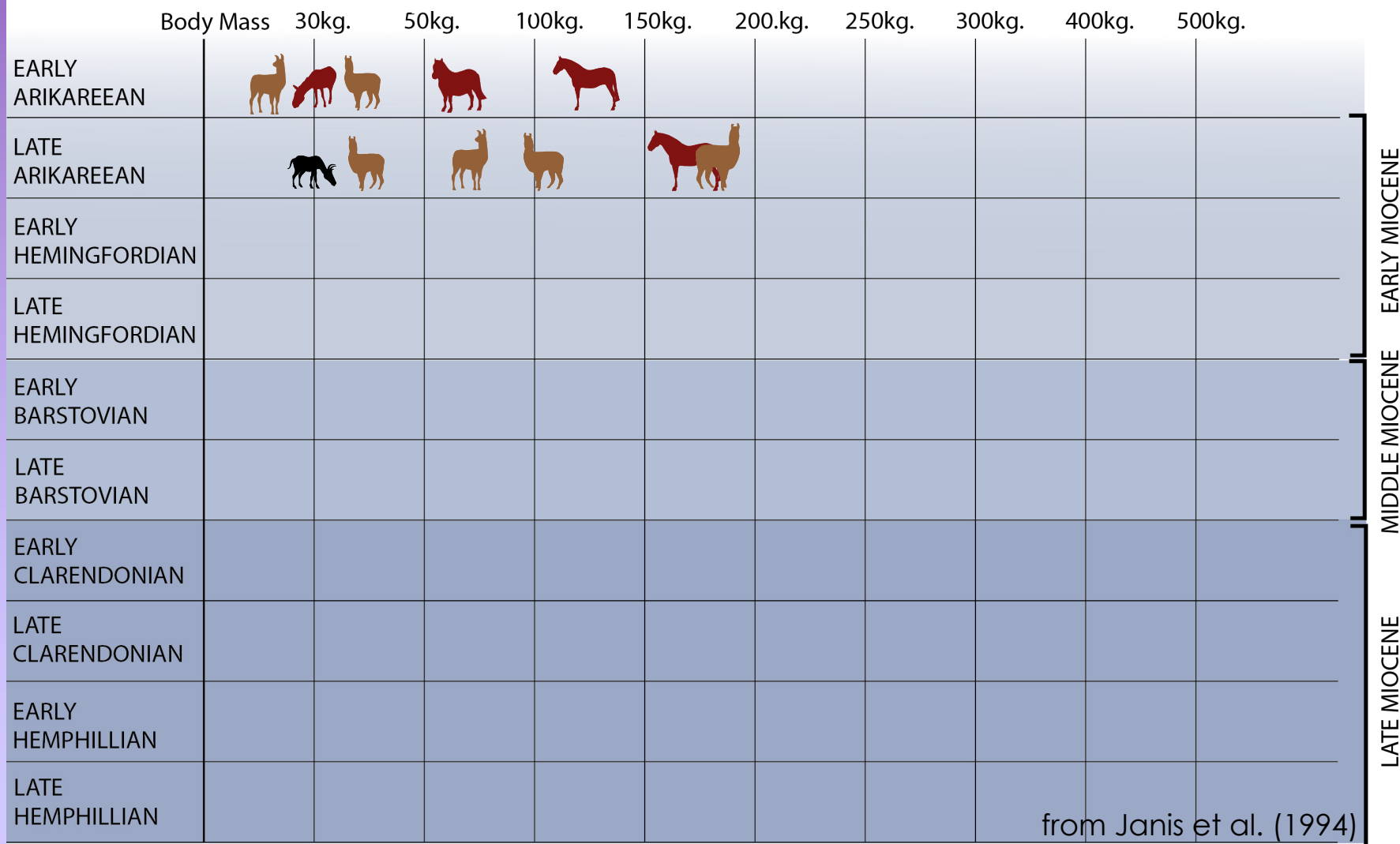




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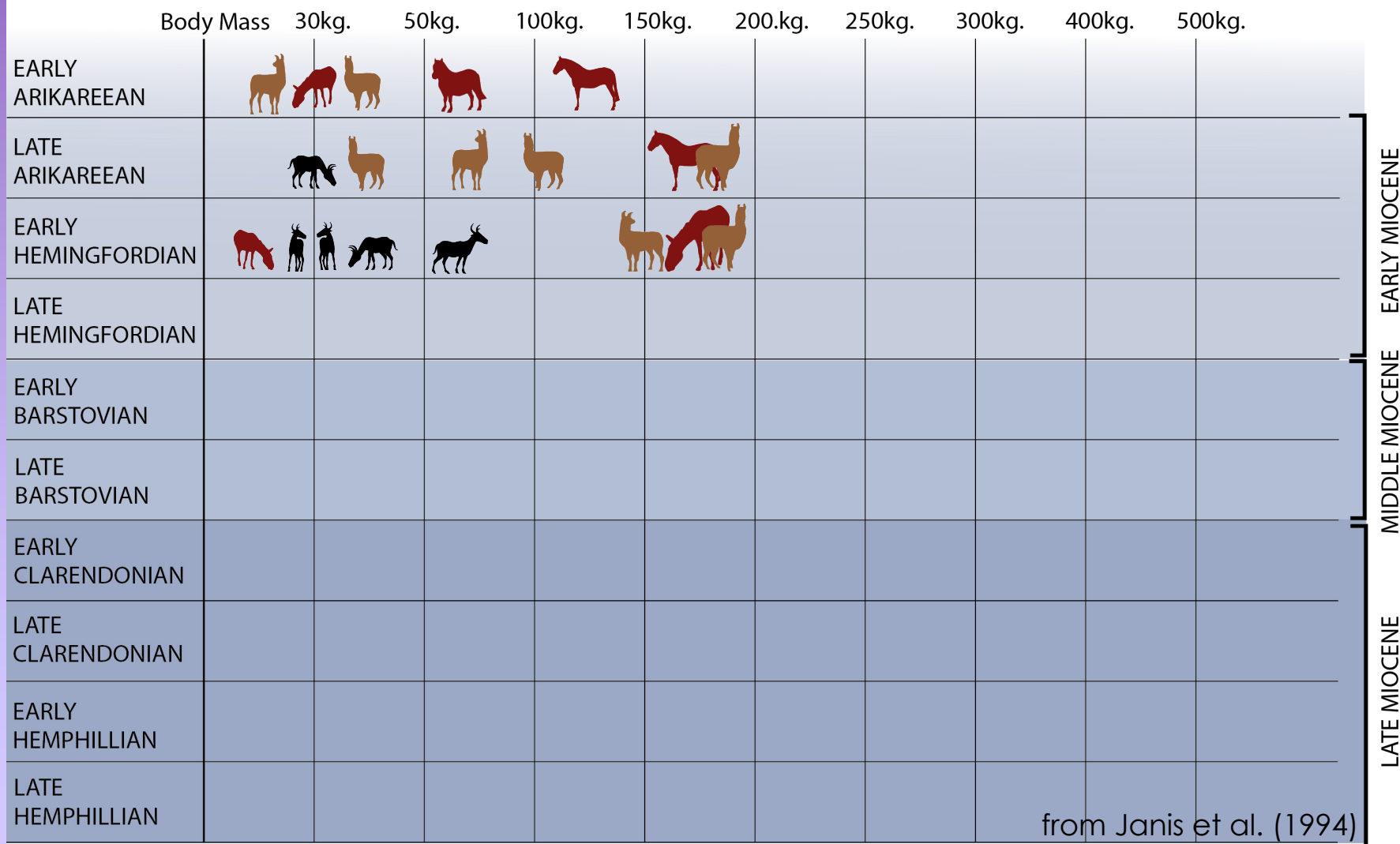




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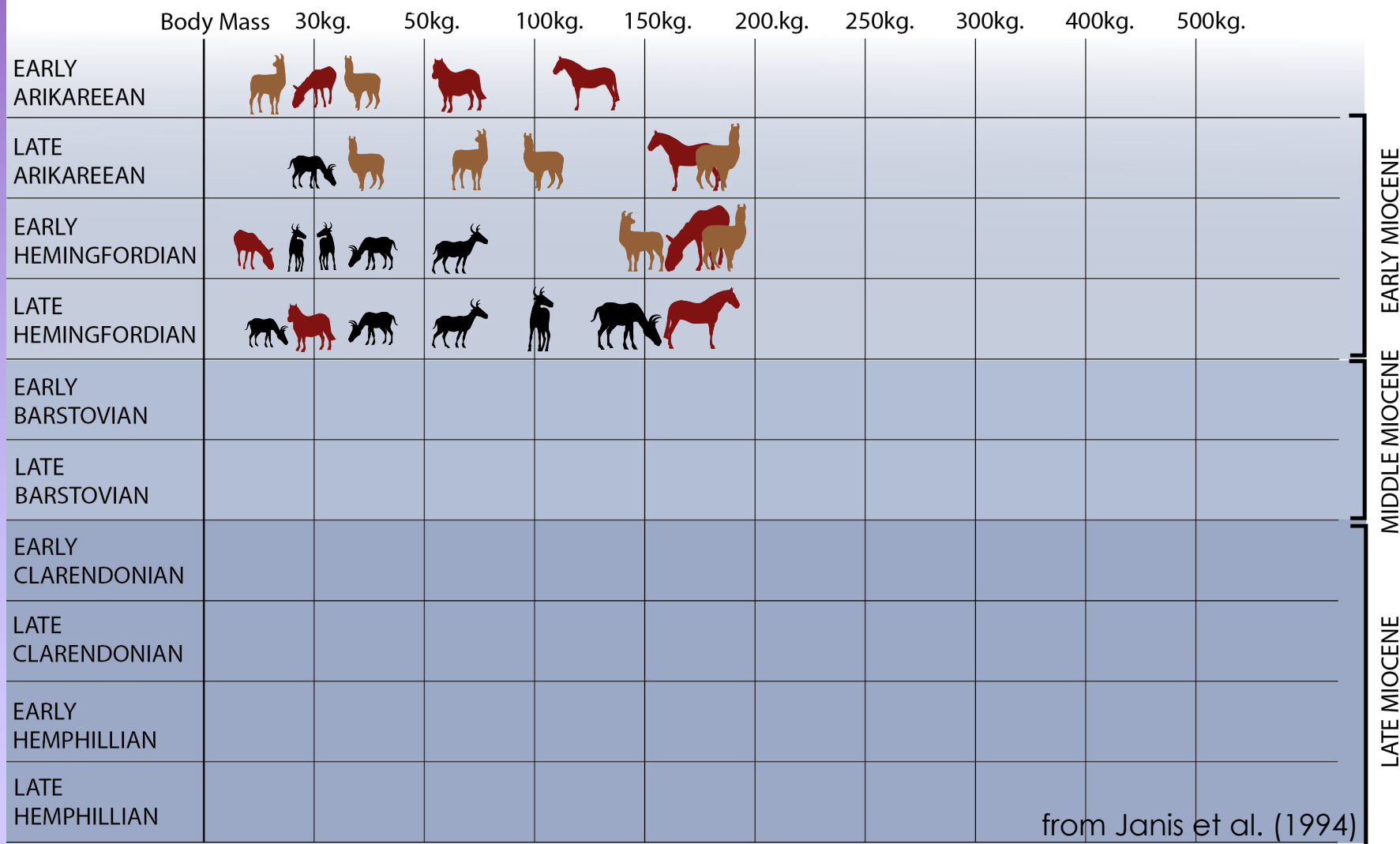




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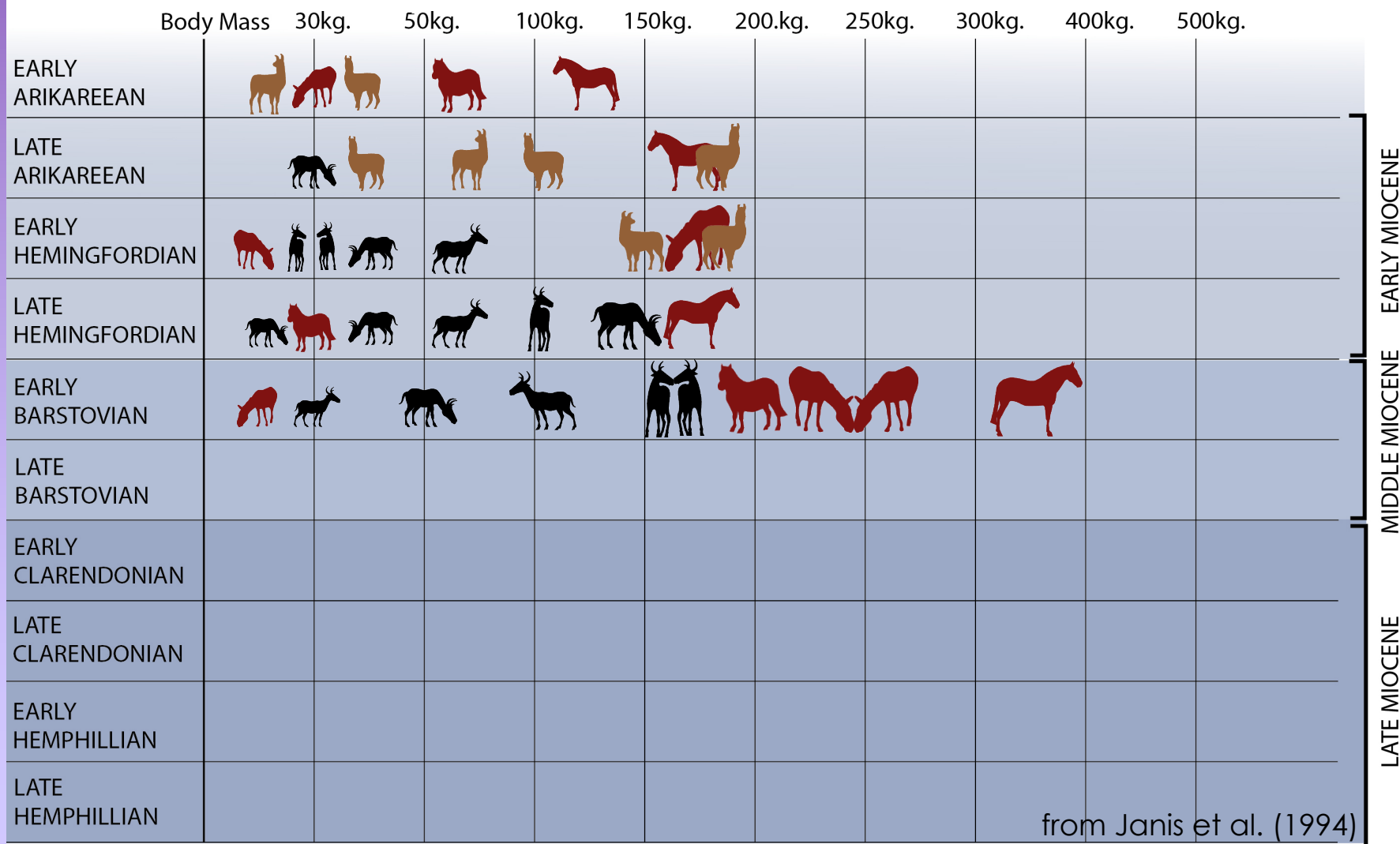




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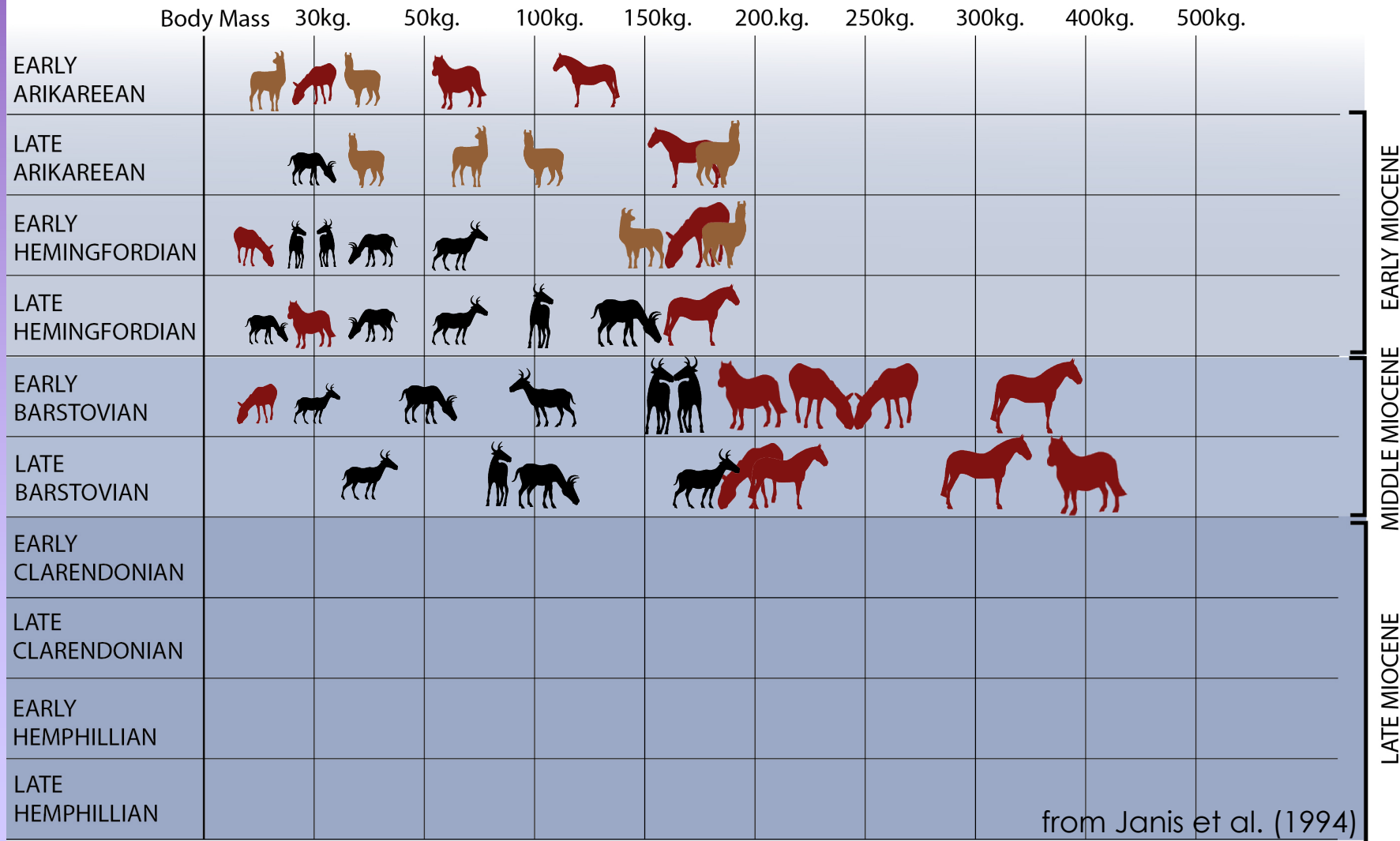




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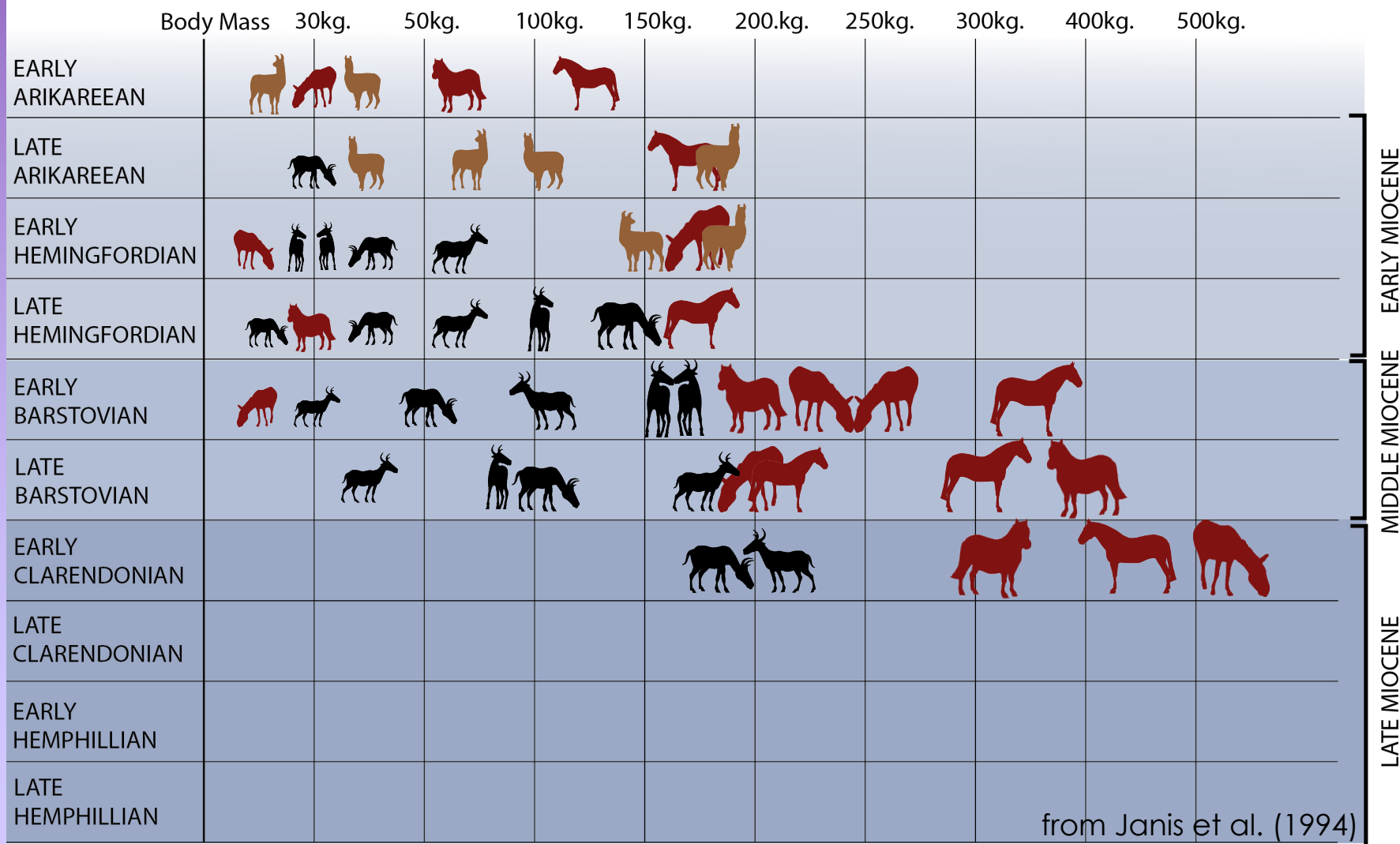




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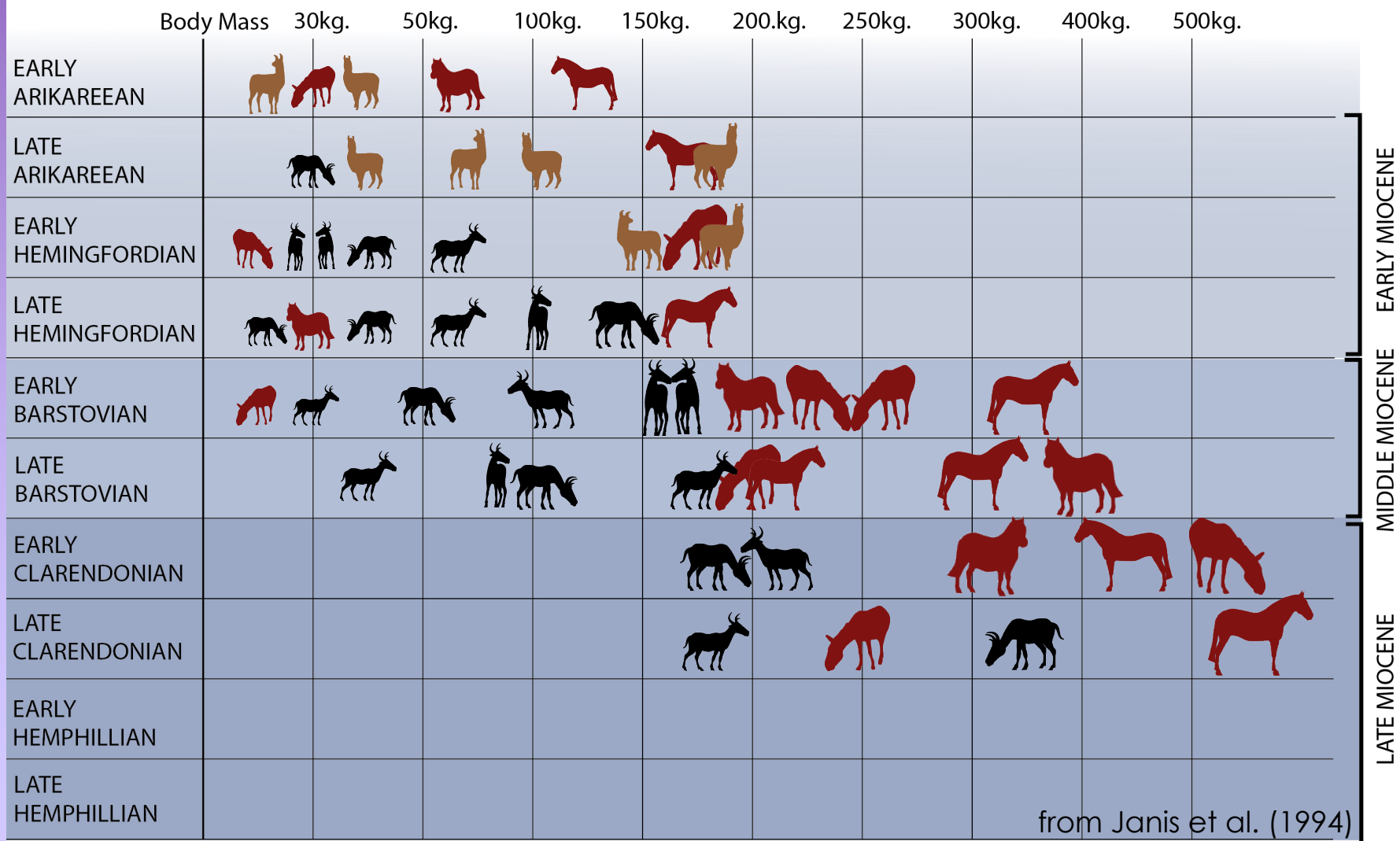




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Summary, Conclusions & Outlook

Rather than understanding tradeoffs along the fast-slow continuum as fixed physical laws, they can be considered as representing the efficiency of the organisms from which the data was taken – and that efficiency may evolve.

Within the boundaries of a specific niche, species possibly compete by demographic means: by evolving a faster reproduction.

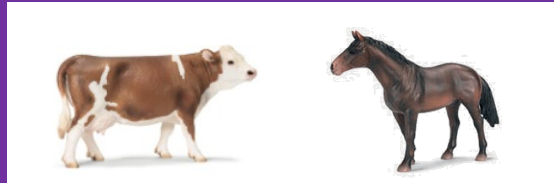
Life history characteristics appear to be linked to taxonomic groups.

We would predict that during geological history, 'faster' species were not replaced by 'slower' species.

The physiological means by which species differ in their life history are not well explored.



By what means do cattle achieve faster intrauterine growth than horses?





A priori conditions and their consequences

By which means can organisms become more efficient?



A priori conditions and their consequences

By which means can organisms become more efficient?

adapting to
optimal
resource use



A priori conditions and their consequences

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resource use

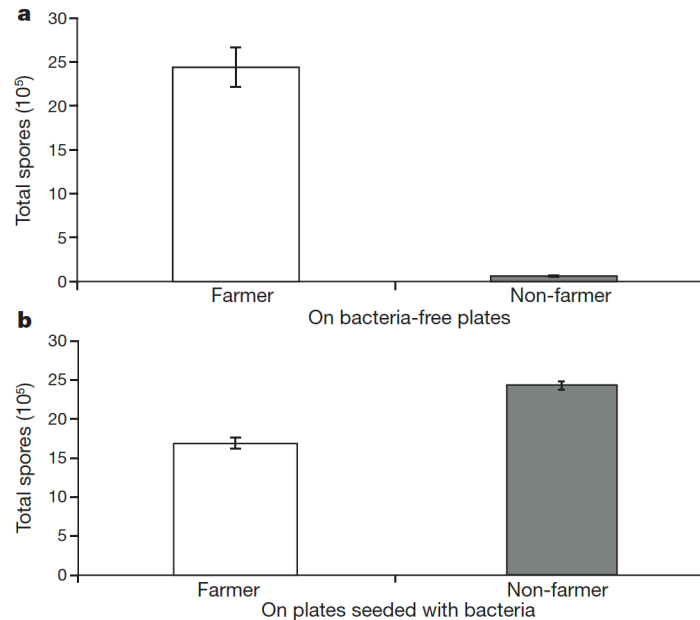
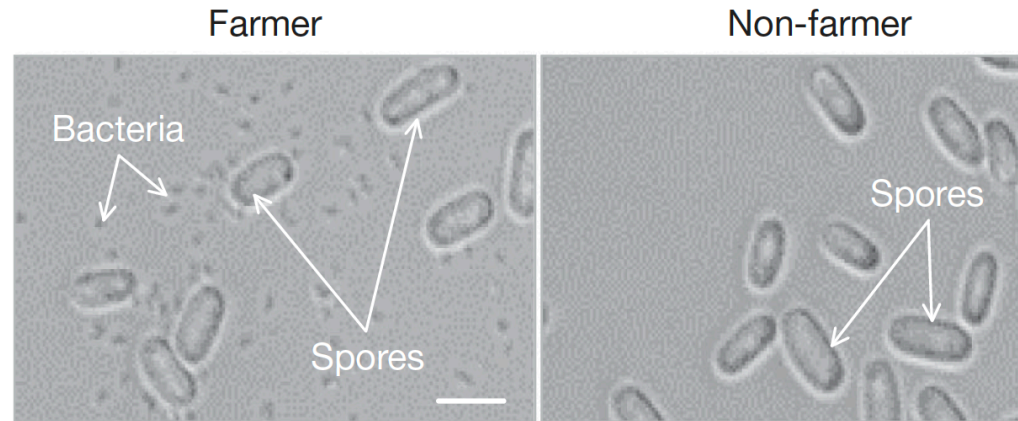
controlling
resources



Primitive agriculture in a social amoeba

Debra A. Brock¹, Tracy E. Douglas¹, David C. Queller¹ & Joan E. Strassmann¹

20 JANUARY 2011 | VOL 469 | NATURE | 393





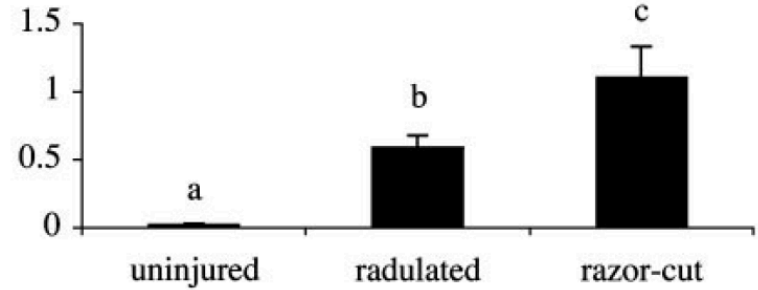
Fungal farming in a snail

Brian R. Silliman*[†] and Steven Y. Newell[‡]

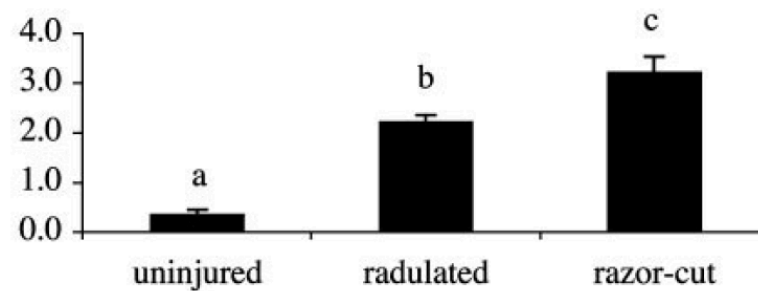
PNAS | December 23, 2003 | vol. 100 | no. 26 | 15643–15648



Fungal Biomass
(erg./ cm² leaf)



Snail growth (mm)



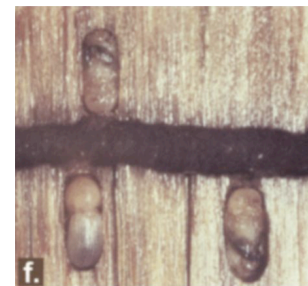
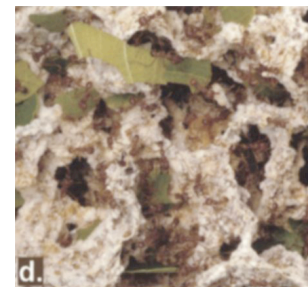


THE EVOLUTION OF AGRICULTURE IN INSECTS

Ulrich G. Mueller,^{1,2} Nicole M. Gerardo,^{1,2,3}

Duur K. Aanen,⁴ Diana L. Six,⁵ and Ted R. Schultz⁶

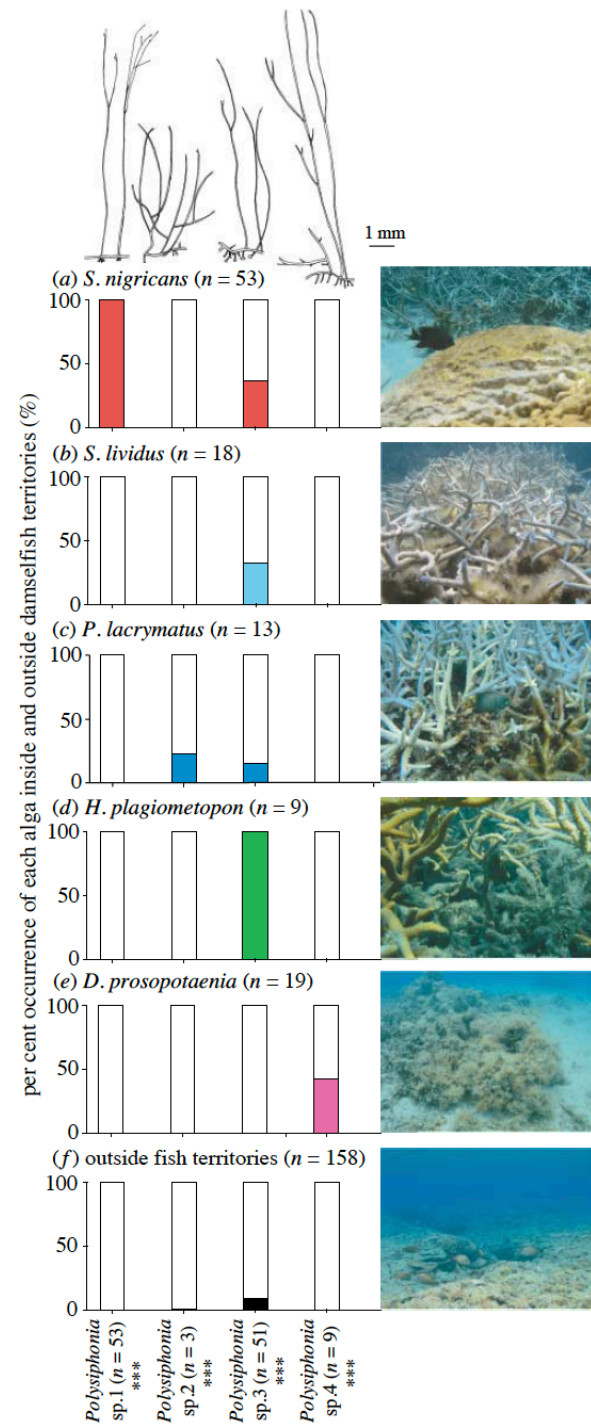
Annu. Rev. Ecol. Evol. Syst. 2005. 36:563–95





A novel obligate cultivation mutualism between damselfish and *Polysiphonia* algae

Hiroki Hata^{†,*} and Makoto Kato





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resource use

controlling
a broad set of resources



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(niche specificity)

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(niche generalism)
“super-niche” emergence



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controlling
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“super-niche” emergence

Probabilistic directionality IV: from use towards control



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“super-niche” emergence

Probabilistic directionality IV: from use towards control

“In the course of the process described by Darwinian evolution, which includes a not-so-sharply-defined relationship between specific niches and more-or-less specific adaptations to them, including arms races and directionality of reproductive efficiency, evolution might (but does not have to), helped by its probabilistic directionality of increasing variability and complexity, evolve an organism whose adaptations are of such a general application that its existence changes the basic, probabilistic rules of evolution.”

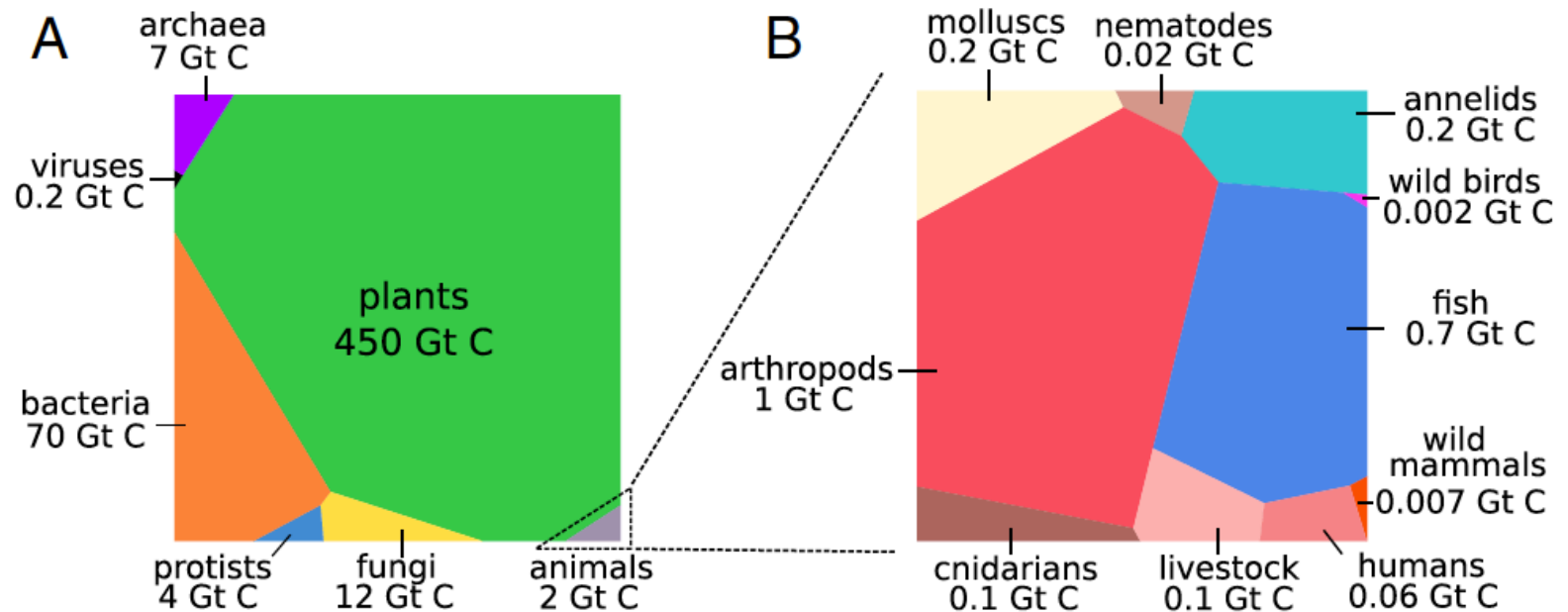


The state of the planet

The biomass distribution on Earth

Yinon M. Bar-On^a, Rob Phillips^{b,c}, and Ron Milo^{a,1}

www.pnas.org/cgi/doi/10.1073/pnas.1711842115



estimate for reptiles similar magnitude as wild birds (but more assumptions; estimate for amphibians not possible)



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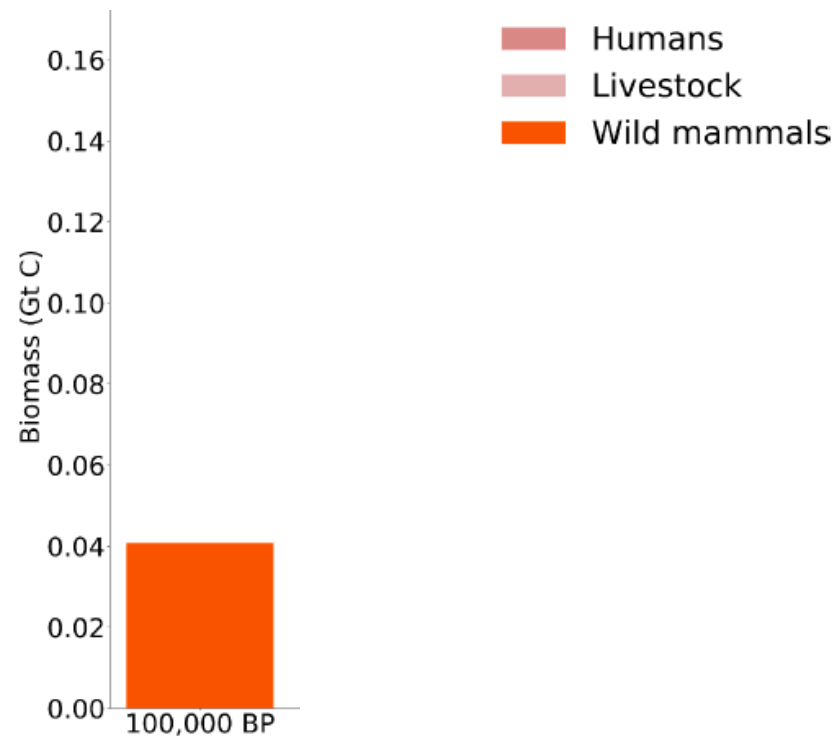


Fig. S5. The impact of human civilization on the biomass of mammals.



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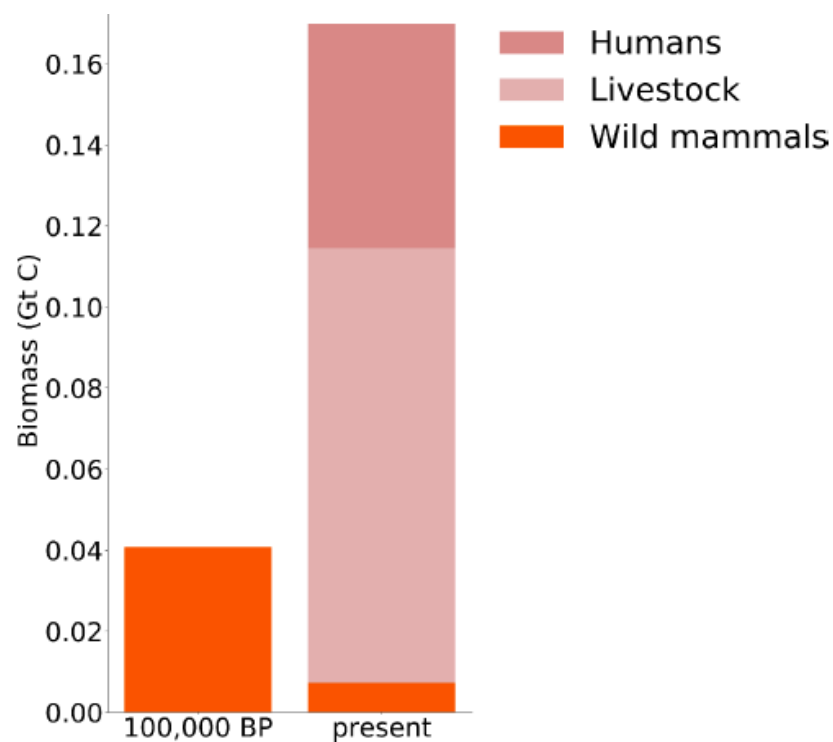


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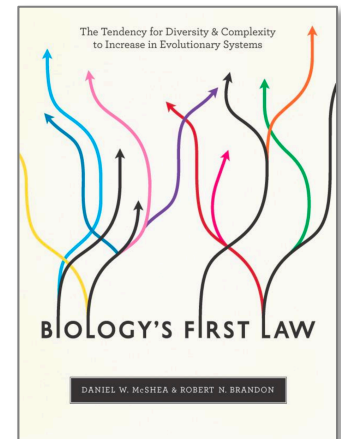
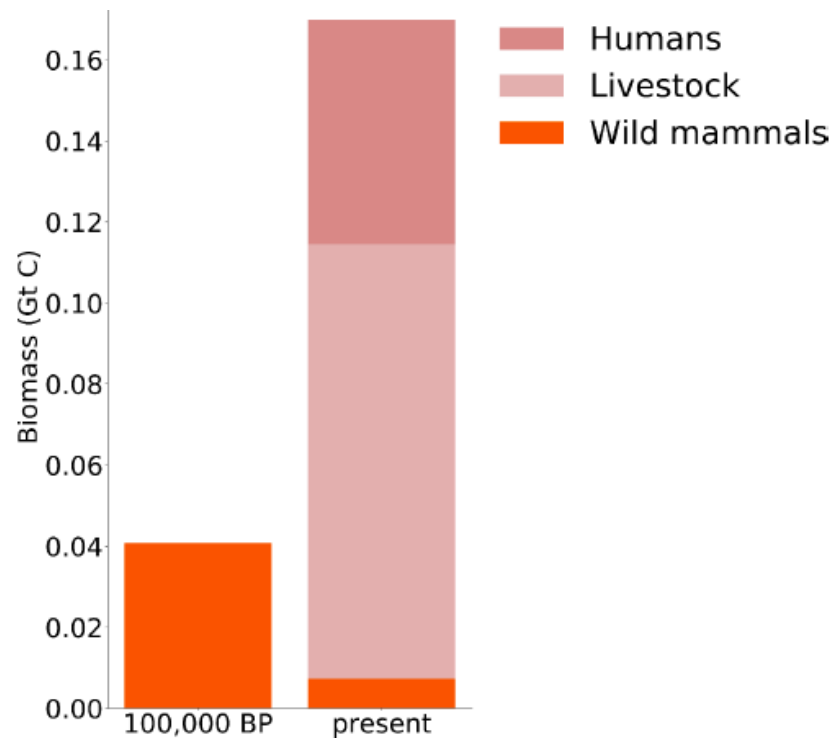


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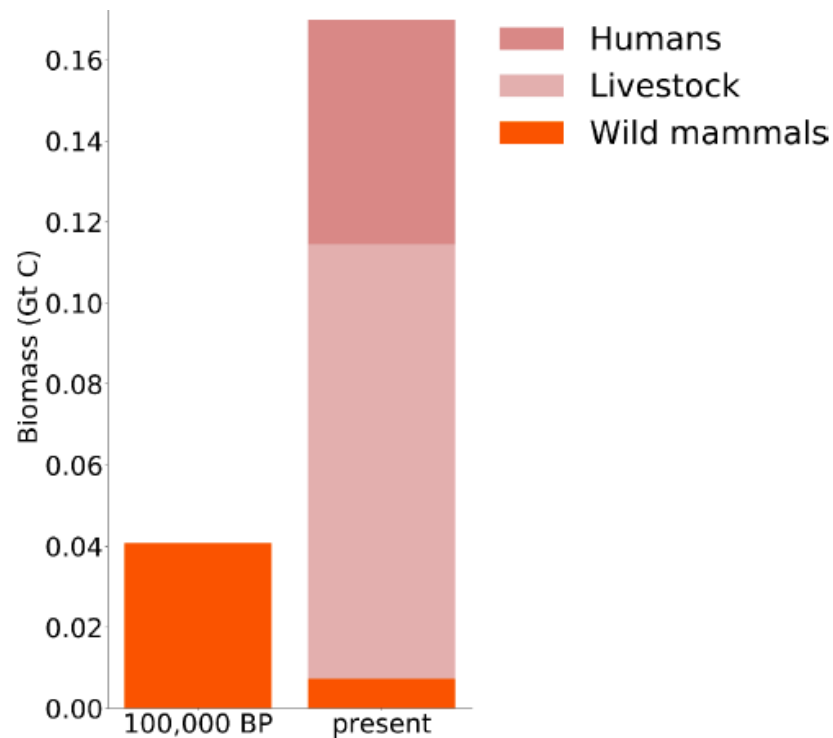


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“super-niche” emergence

Probabilistic directionality IV: from use towards control



A priori conditions and their consequences

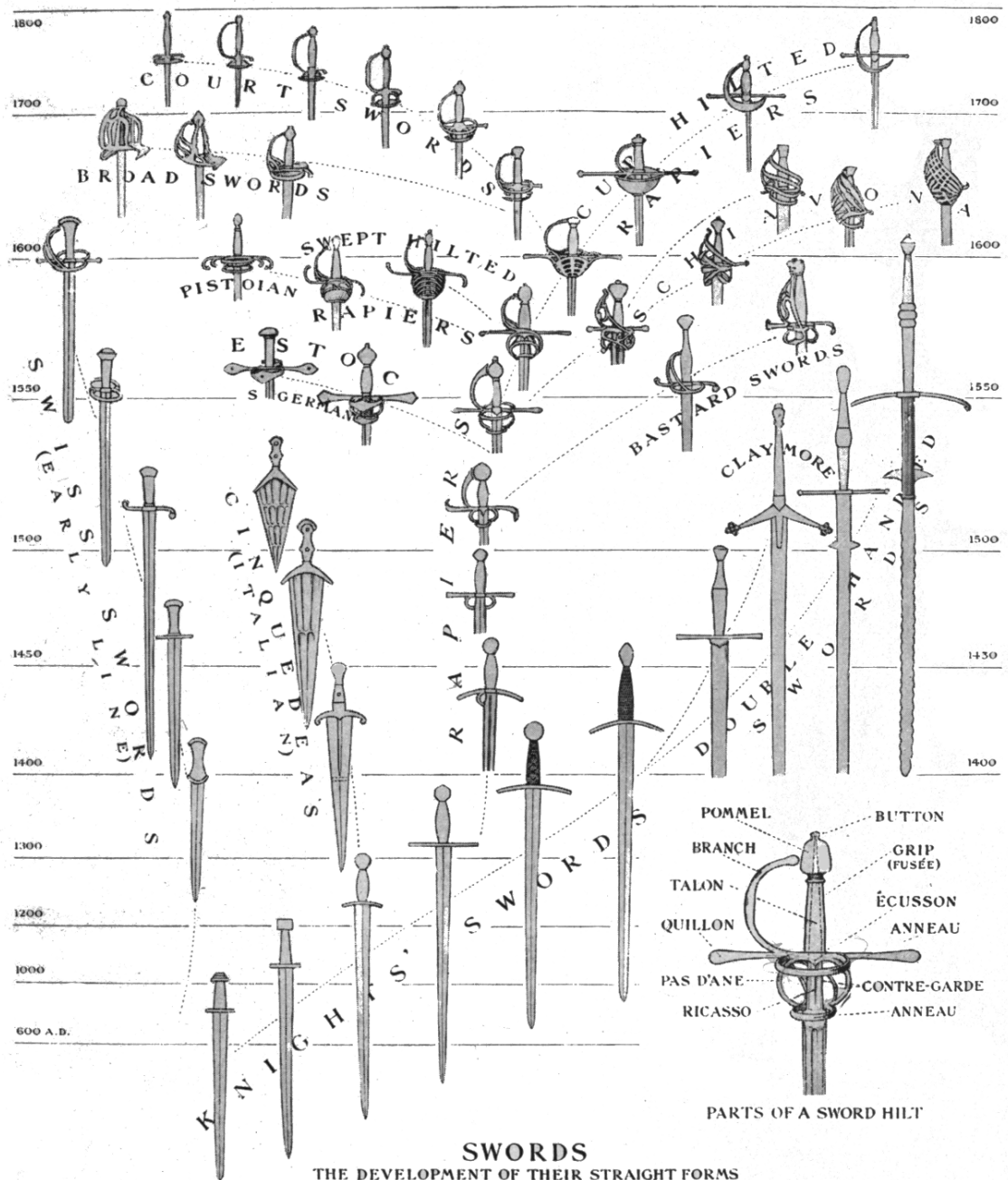
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no species diversity but diversity of
sub-niches, culture, tools ...
intraspecific competition = history



SWORDS

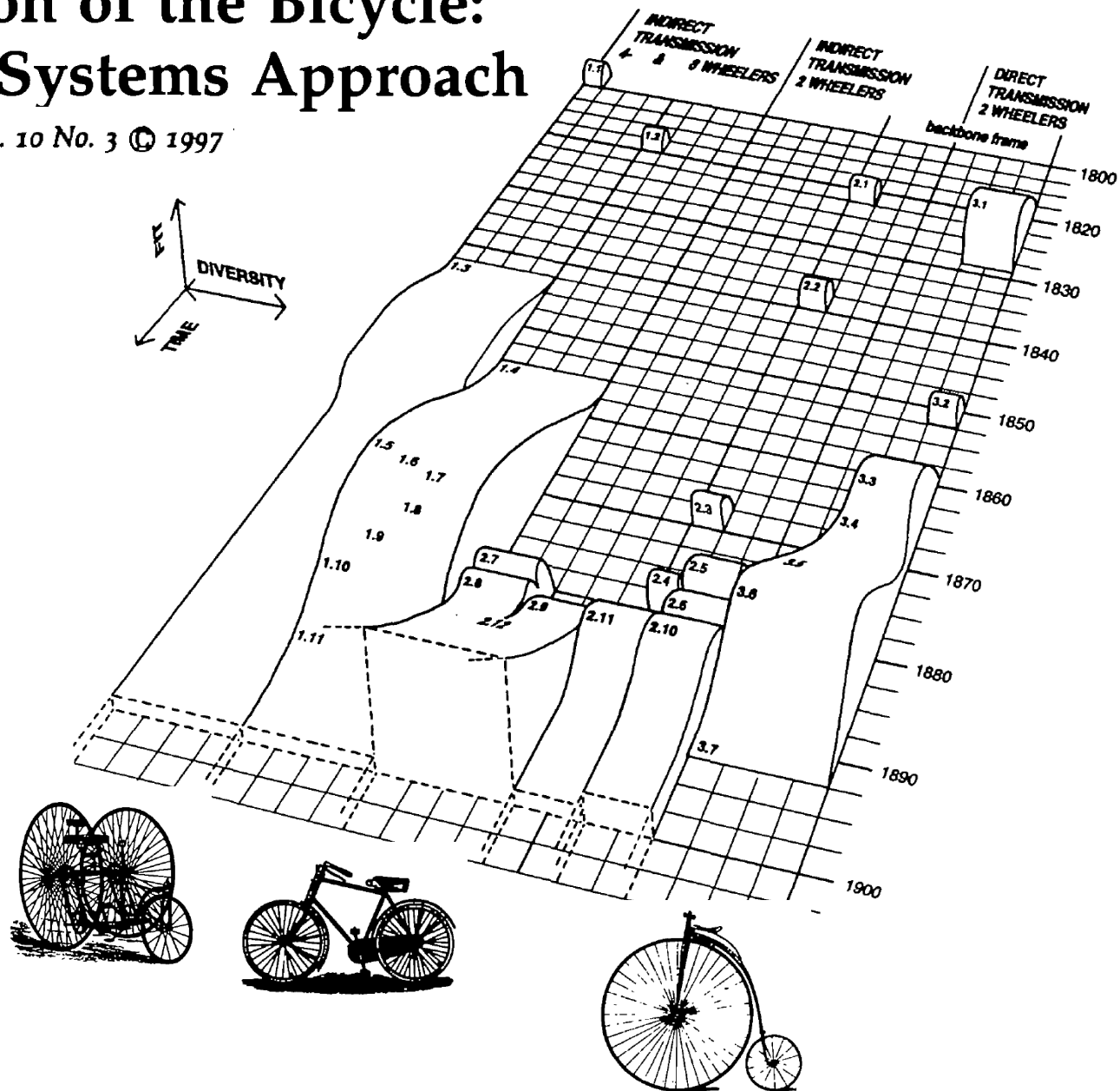
THE DEVELOPMENT OF THEIR STRAIGHT FORMS DURING THE CENTURIES

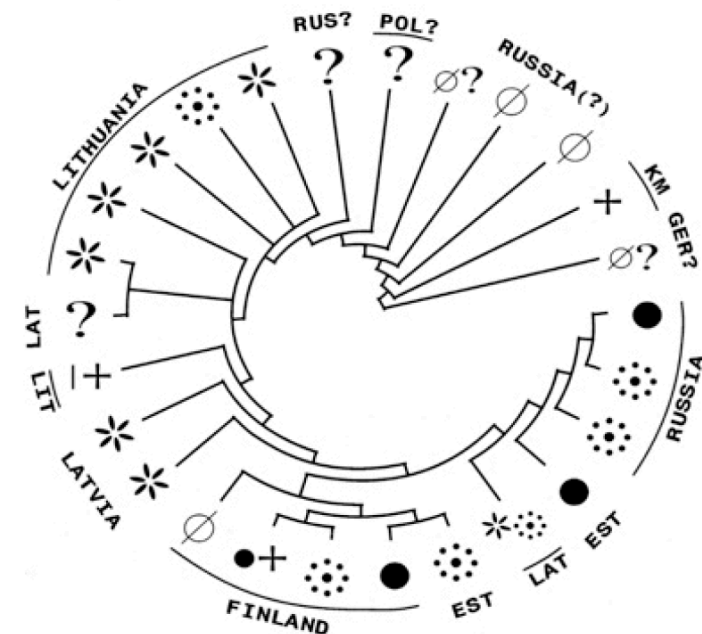


O. A. van Nierop, A. C. M. Blankendaal, and C. J. Overbeeke

The Evolution of the Bicycle: A Dynamic Systems Approach

Journal of Design History Vol. 10 No. 3 © 1997

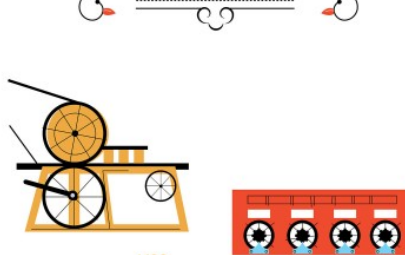




THE EVOLUTION OF COMPUTERS

PC NINJA TRAVELS THROUGH TIME, revealing the history of how computers became our sidekicks. From sliding pebbles on a simple machine to swiping your fingers across a touchscreen, technology has transformed radically!

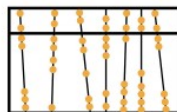
HISTORIC



- 1439** Johannes Gutenberg invents the printing press.
- 1642** Blaise Pascal invents the **Pascaline** (Pascals' Calculator), a mechanical adding machine.

2400 BC

The **abacus**, the first known calculator, invented in Babylonia. Decks of beads and rods used to calculate addition, subtraction, division and multiplication, along with being able to extract square and cube roots.



1617

Jean Napier creates the **Napier's Bones**, a manually operated calculating device consisting of a base board with a row and rods, unrelated with multiplication tables, which reduce multiplication to addition and division to subtraction.

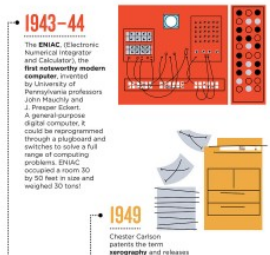
1822

Charles Babbage conceives designs for the **Difference Machine**, a steam-powered calculating machine that is considered one of the first programmable computers, though the technology was too advanced to be built in his time.

KEY

- PC HISTORY
- MAC HISTORY
- INTERNET HISTORY
- OTHER DEVICES HISTORY

MODERN



- 1943-44** The **ENIAC** (Electronic Numerical Integrator and Calculator), the first **reprogrammable** machine computer, invented by University of Pennsylvania professors John Mauchly and J. Presper Eckert. A general-purpose digital computer, it could be reprogrammed through a keyboard and switch to solve a full range of computing problems. ENIAC occupied a room 30 by 50 feet in size and weighed 30 tons!
- 1949** Chester Carlson patents the term **xerography** and releases the **Xerox** machine. Model A photocopiers, which required 30 steps to make one copy!
- 1952** The first mass-produced computer made and sold by the IBM Company.
- 1962** J.C.B. Licklider conceives of an **inter-galactic network** of computers, later known as the **Internet**.

1971

First e-mails sent by Ray Tomlinson on the ARPANet, the precursor to the Internet, which were able to reach users on different hosts. He used the "W" sign to separate a person from a machine, which became a standard part of the e-mail address.



1975

The IBM 5100 becomes the first commercially available portable computer.



1976

The first **Apple** computer invented by Steve Jobs and Steve Wozniak.

1946

Mauchly and Eckert receive funding from the Census Bureau to build the **UNIVAC** (Universal Automatic Computer), the first commercial computer for business and government applications.



1952

The first computer game, "Tic-Tac-Toe," designed and created by A.S. Douglas.



1971

IBM introduces the first **floppy disk** that is 8 1/2 inches wide with a capacity of 797 KB, and was made only a year later into a hard disk drive.



1964

Computer systems that use **tokens**, **windows**, and a **mouse** designed by Douglas Engelbart, which will lay the foundation for modern computing.



1975

Bill Gates and Paul Allen founded **Microsoft**.

MICROSOFT

1977

The **Apple II**, the first personal computer with **color graphics** is demonstrated.



1979

Word processing becomes a reality as **Microsoft International** releases **WordStar**.



1985

Bill Gates and Microsoft release the **Windows** operating system, forever changing desktop computing.



1989

The **"World Wide Web"** invented by Tim Berners-Lee, who developed HyperText Markup Language (HTML) to design webpages.



1990

Release of the original **Microsoft Office 1.0**, featuring Word 1.0, Excel 2.0, and PowerPoint 2.0.



1995

Amazon launches, and online shopping becomes popular.



2003

Electronic devices that could send e-mails, access the web, and receive phone calls become commercial.



2010

Apple releases the **iPad**, a tablet computer with a touchscreen that changes the way users view news, e-mails, devices, and in 80 days.



2001

Apple unveils the **Mac OS X** operating system, which provides protected memory architecture and pre-emptive multi-tasking among other benefits. Microsoft rolls out **Windows XP**, which has a significantly redesigned GUI.



2013

Microsoft releases **Windows 8**, a completely redesigned operating system that is tailored toward touchscreens, tablets, and cloud functionality.



1999

The term **Wi-Fi** becomes part of computing language and users begin connecting to the Internet without wires.

1995

Microsoft releases **Windows 95**, which introduces the web browser **Internet Explorer**.



2007

Apple debuts the **iPhone**, both a phone and handheld computer that runs "apps."





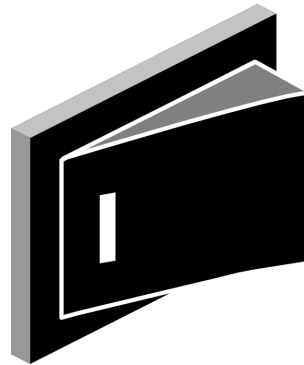
Directionality in Evolution:
beware of the *natural fallacy*



A priori conditions and their consequences

By which means can organisms become more efficient?

adapting to
optimal
resource use



controlling
a broad set of resources

Probabilistic directionality IV: from use towards control

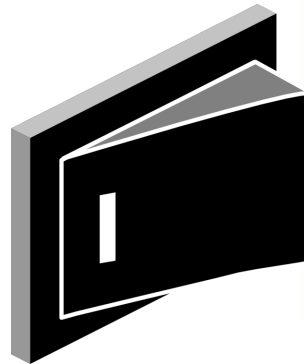
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