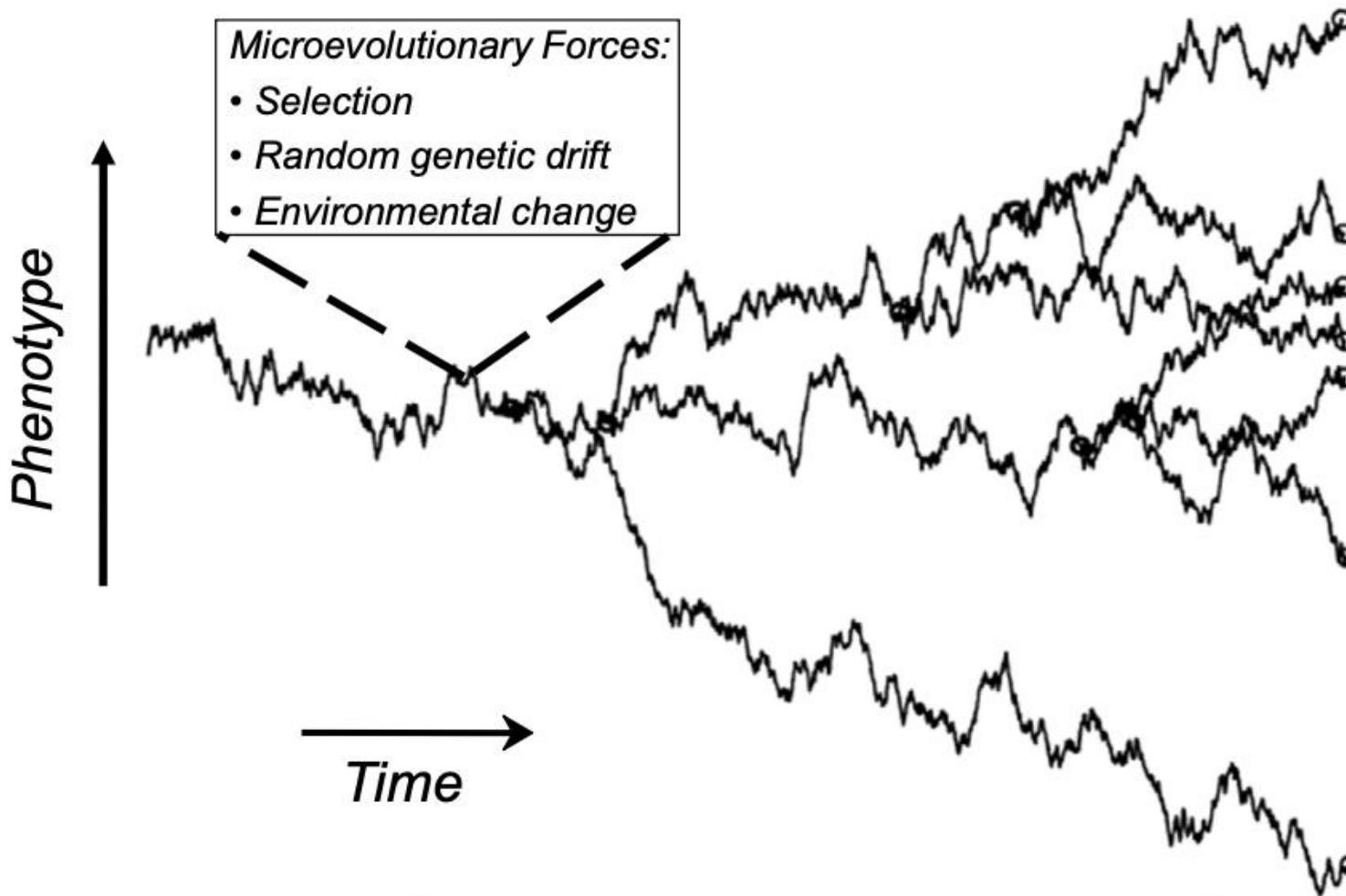


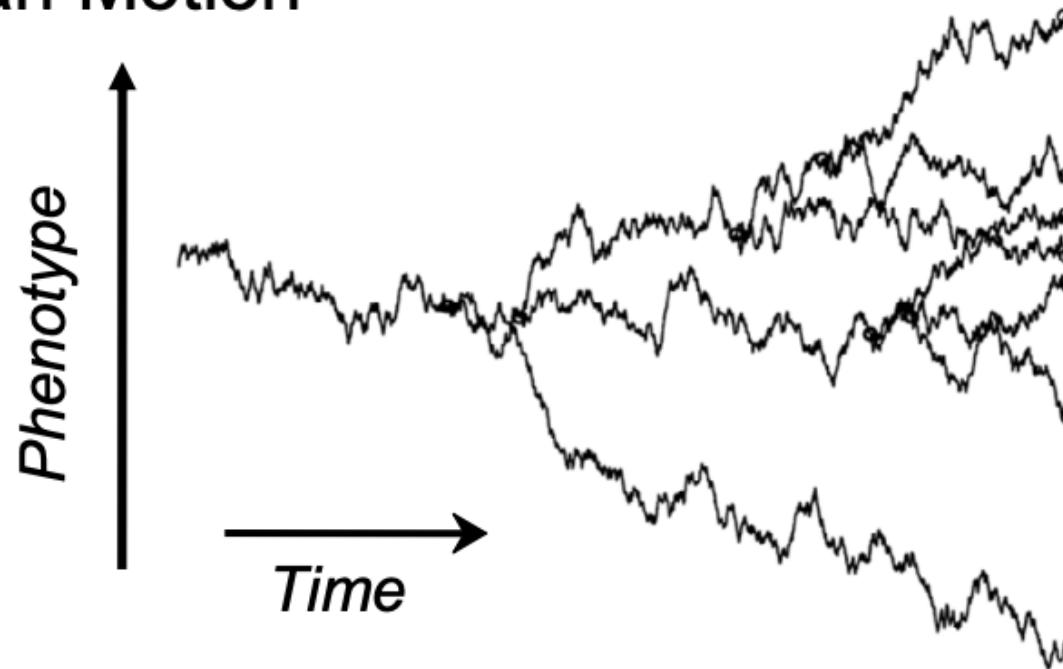
# Continuous Trait Correlations



**Comparative Data**

***Evolutionary Thinking***

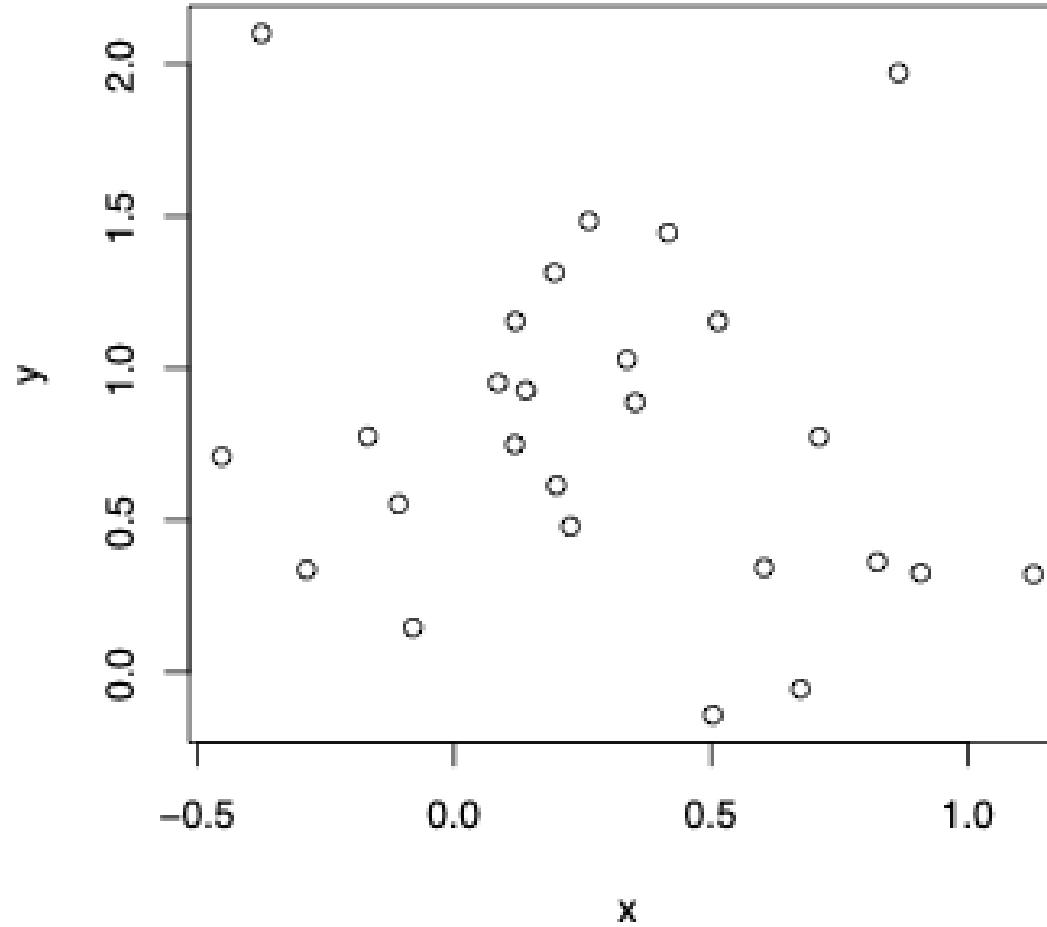
## Brownian Motion



$$\text{Variance} = \sigma^2 t_a$$

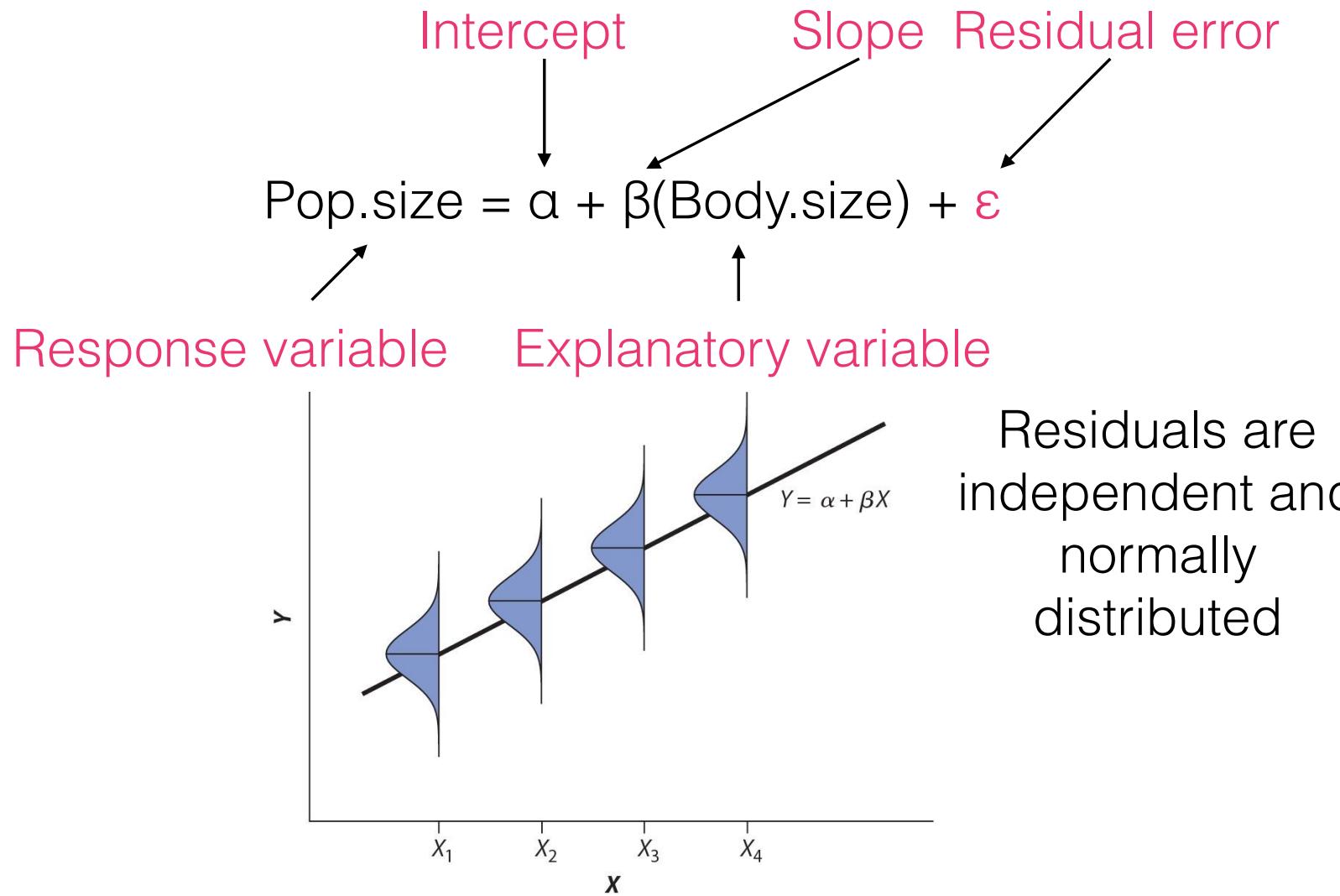
- Neutral evolution via random genetic drift
- Fluctuating directional selection
- Constant directional selection

Problem:  
How much  
of  $X$  explains  
 $Y$



# Assumptions of a linear model (linear regression)

1. Linear relationship
2. No Multicollinearity
3. Independence
4. Homoscedasticity
5. Residuals distribution



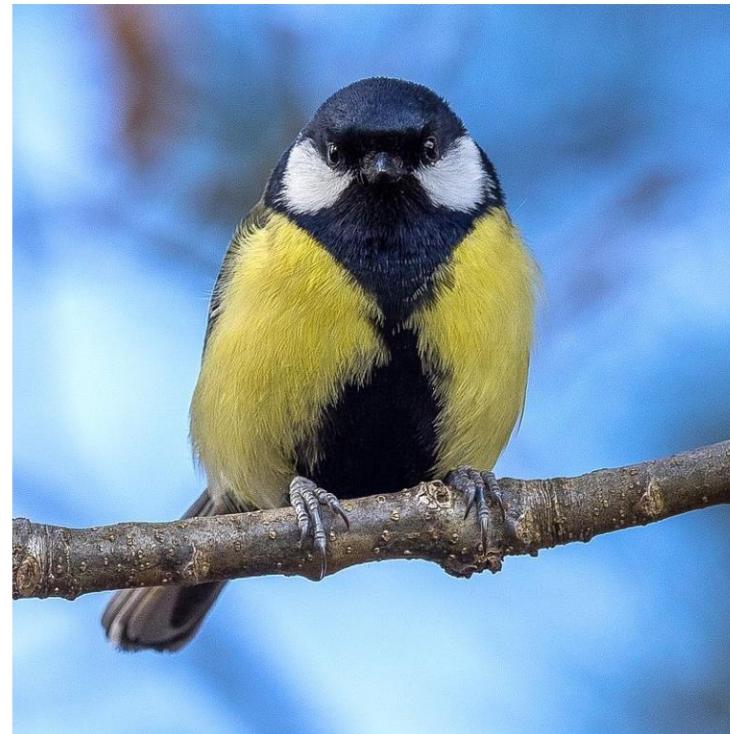
# *Do bird species with larger average body size have higher abundance?*

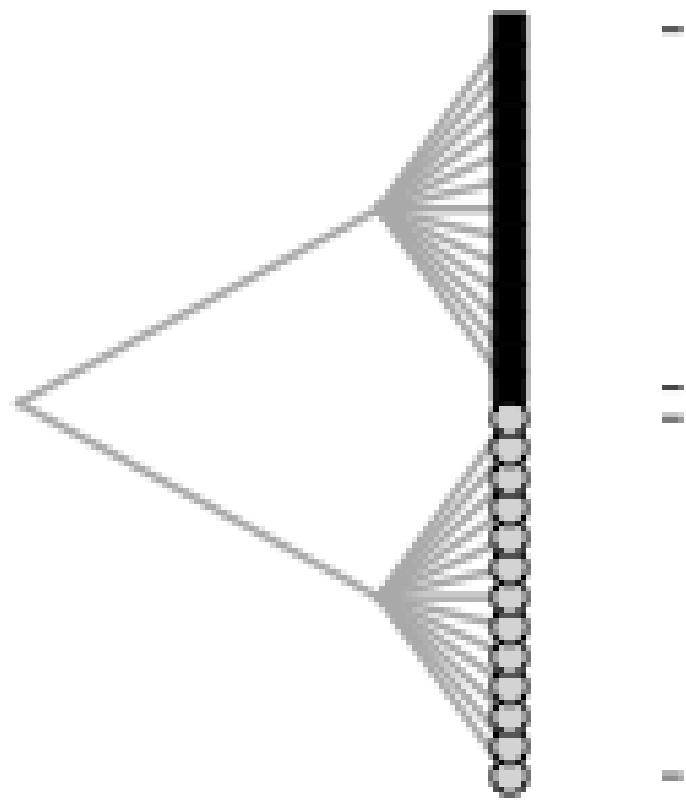
**Sean Nee, Andrew F. Read, Jeremy J. D. Greenwood\***  
**& Paul H. Harvey**

AFRC Unit of Ecology and Behaviour, Department of Zoology,  
University of Oxford, South Parks Road, Oxford OX1 3PS, UK

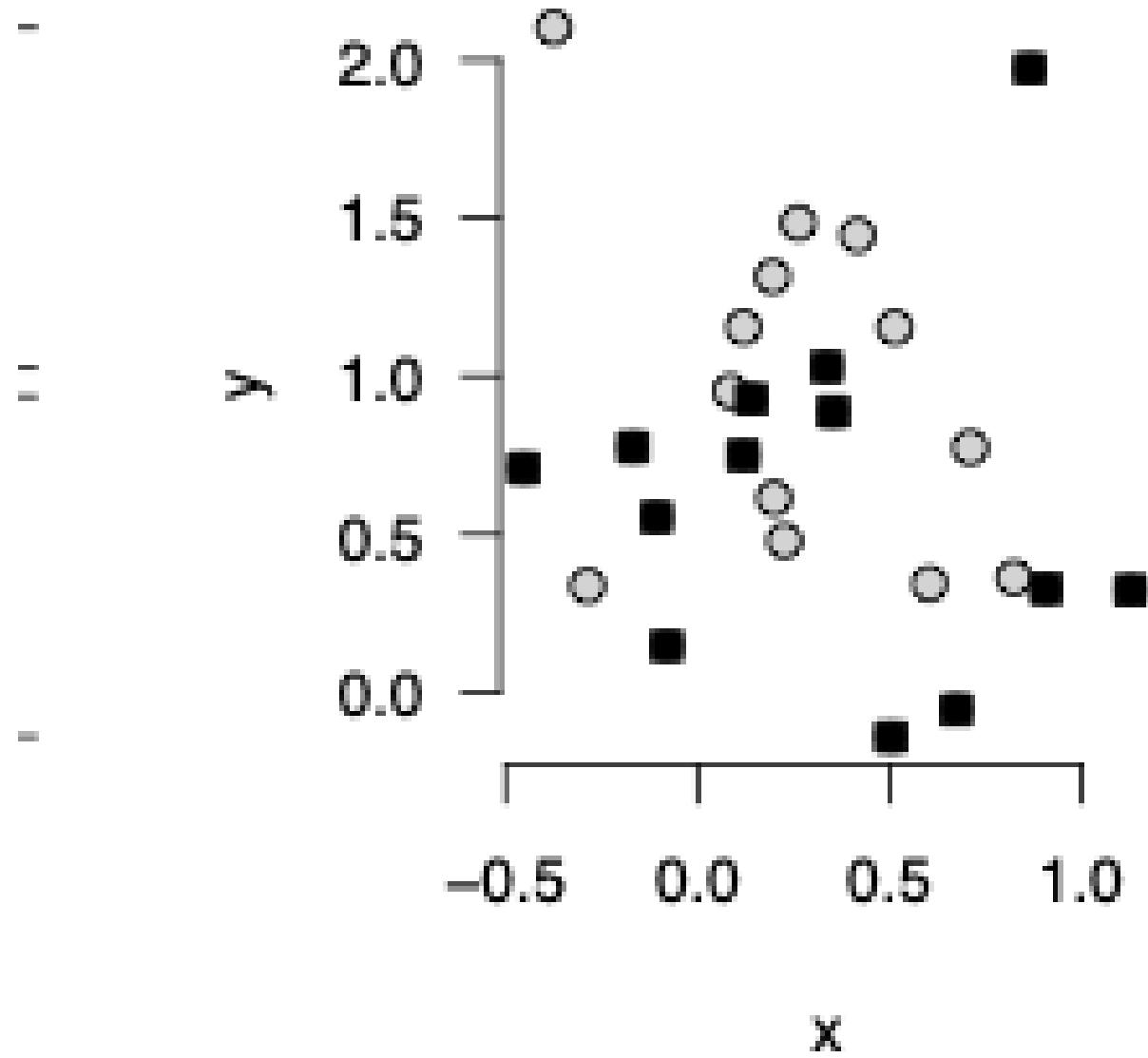
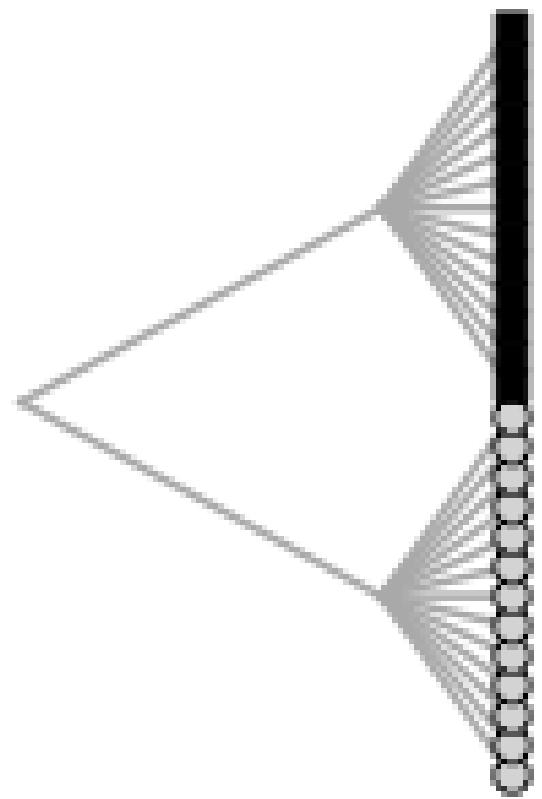
\* British Trust for Ornithology, Thetford, Norfolk IP24 2PU, UK

THE relationship between abundance and body size is the subject of considerable debate in ecology<sup>1–15</sup>. Several data sets spanning a large range of body sizes show linear negative relationships between abundance and weight<sup>1–6</sup> when these are measured on a logarithmic scale. But other studies of the abundances of species from single taxa, such as birds, which span a narrower range of body sizes reveal either little or no relationship, or a triangular relationship<sup>9–15</sup>. Errors in estimating abundance might obscure relationships that do exist over a narrow range of body sizes. We

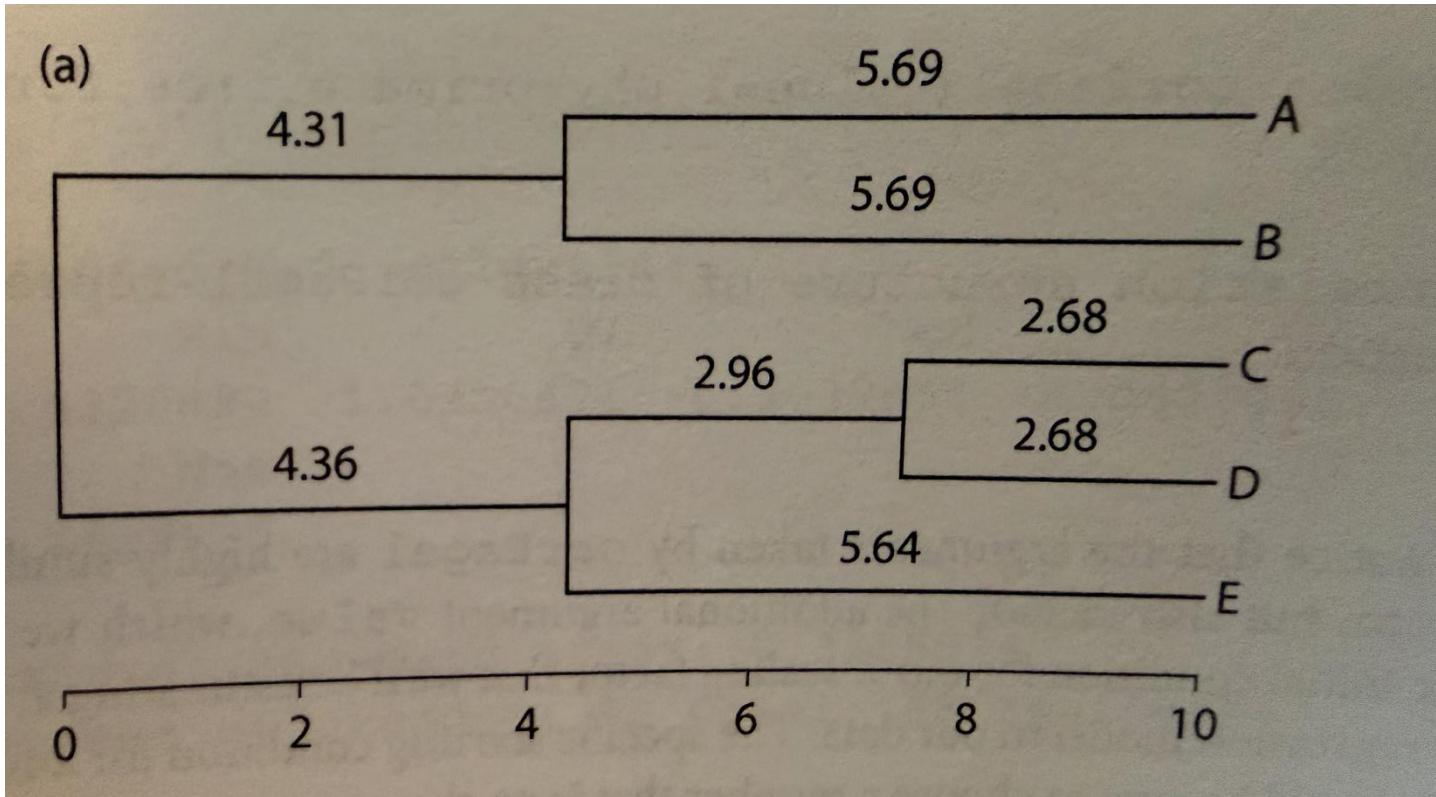


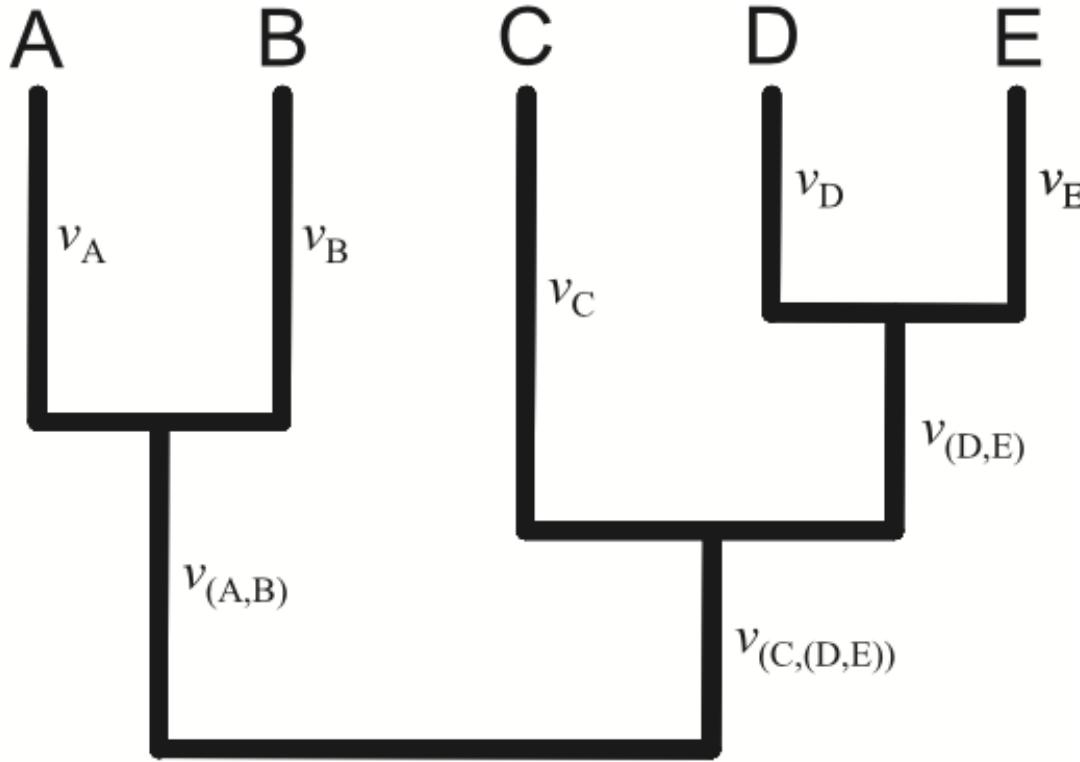


- - - - -



(a)





**(b)**

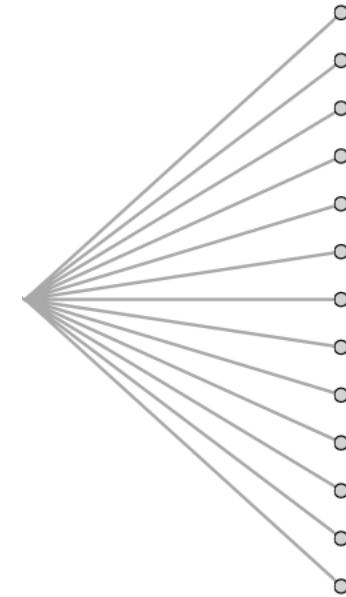
$$\mathbf{C} = \begin{bmatrix} & \mathbf{A} & \mathbf{B} & \mathbf{C} & \mathbf{D} & \mathbf{E} \\ \mathbf{A} & v_{(A,B)} + v_A & v_{(A,B)} & 0.00 & 0.00 & 0.00 \\ \mathbf{B} & v_{(A,B)} & v_{(A,B)} + v_B & 0.00 & 0.00 & 0.00 \\ \mathbf{C} & 0.00 & 0.00 & v_{(C,(D,E))} + v_C & v_{(C,(D,E))} & v_{(C,(D,E))} \\ \mathbf{D} & 0.00 & 0.00 & v_{(C,(D,E))} & v_{(C,(D,E))} + v_{(D,E)} + v_D & v_{(C,(D,E))} + v_{(D,E)} \\ \mathbf{E} & 0.00 & 0.00 & v_{(C,(D,E))} & v_{(C,(D,E))} + v_{(D,E)} & v_{(C,(D,E))} + v_{(D,E)} + v_E \end{bmatrix}$$

$$Y = \beta X + \varepsilon$$

$$\varepsilon \sim N(0, \sigma^2 I)$$

Strong, rapid selection.  
Any force that erases history.

	A	B	C	D	E	F	G	H
A	$\sigma^2$	0	0	0	0	0	0	0
B	0	$\sigma^2$	0	0	0	0	0	0
C	0	0	$\sigma^2$	0	0	0	0	0
D	0	0	0	$\sigma^2$	0	0	0	0
E	0	0	0	0	$\sigma^2$	0	0	0
F	0	0	0	0	0	$\sigma^2$	0	0
G	0	0	0	0	0	0	$\sigma^2$	0
H	0	0	0	0	0	0	0	$\sigma^2$



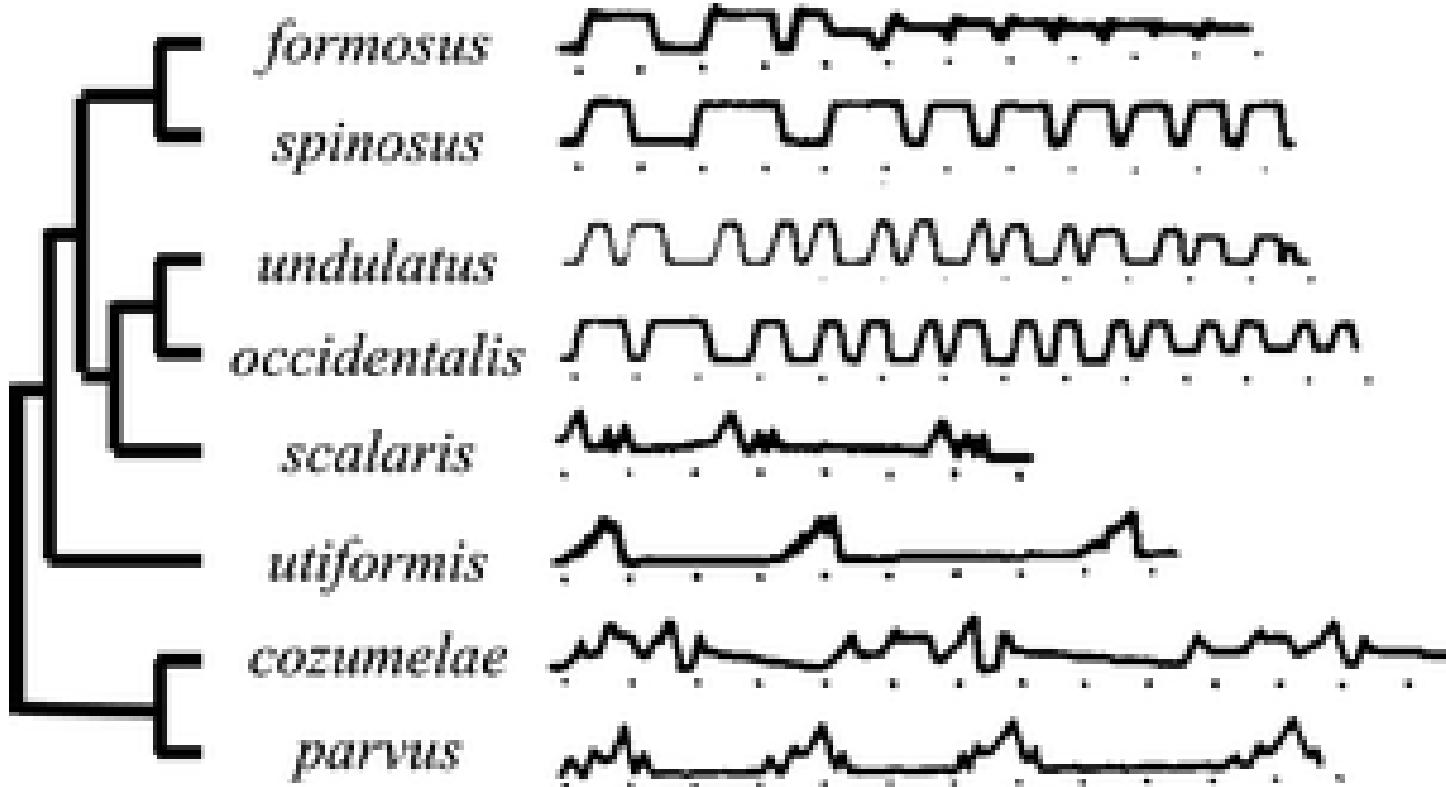
# Phylogenetic linear model assumptions

# Phylogenetic linear model assumptions

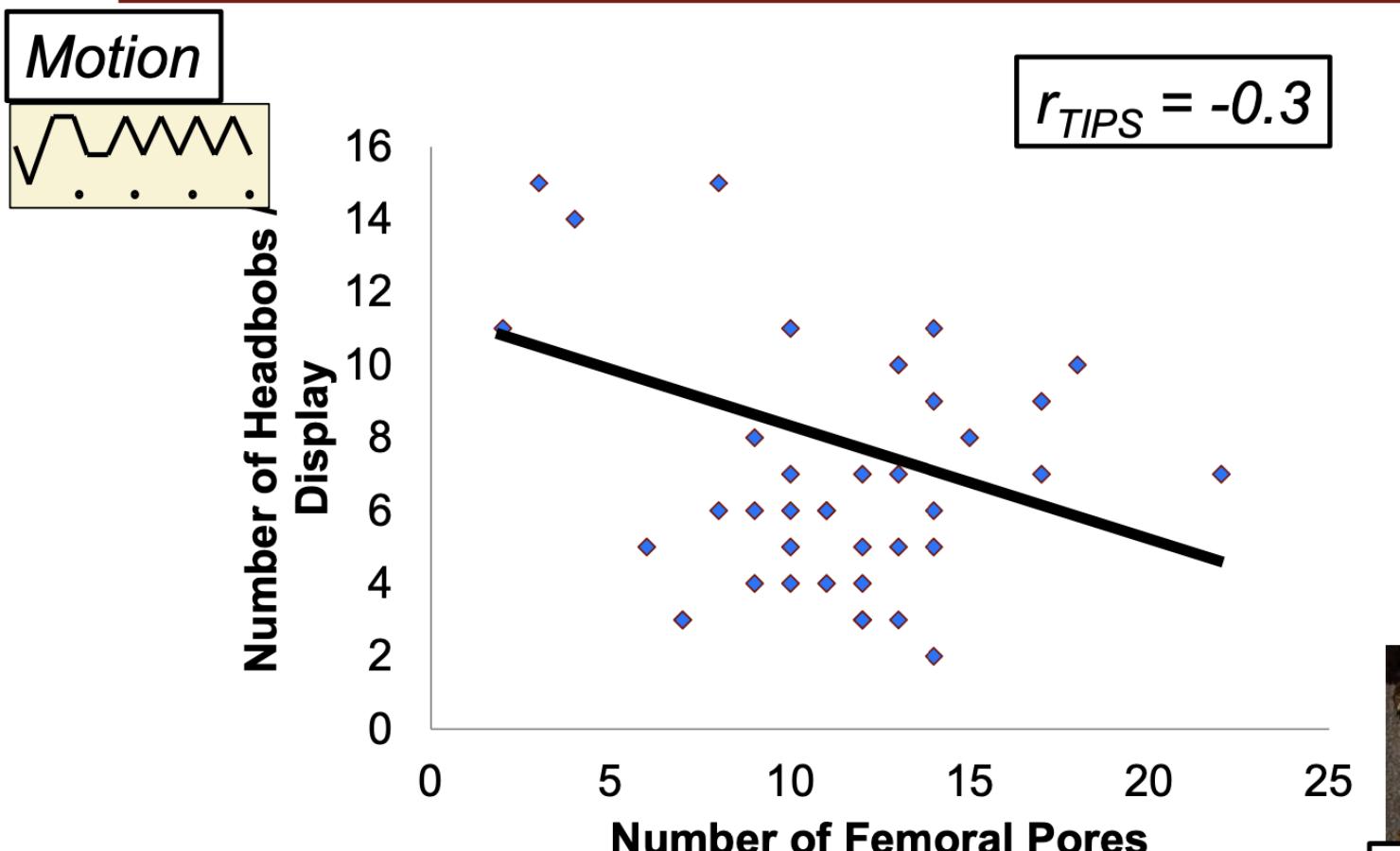
Maximum Likelihood estimates

$$\hat{\beta} = (\mathbf{X}' \mathbf{C}^{-1} \mathbf{X})^{-1} \mathbf{X}' \mathbf{C}^{-1} \mathbf{y}. \quad \sigma_{\varepsilon}^2 = (1/n)(\mathbf{y} - \mathbf{X}\hat{\beta})' \mathbf{C}_{\lambda}^{-1} (\mathbf{y} - \mathbf{X}\hat{\beta}).$$

# Lizard head bobbing

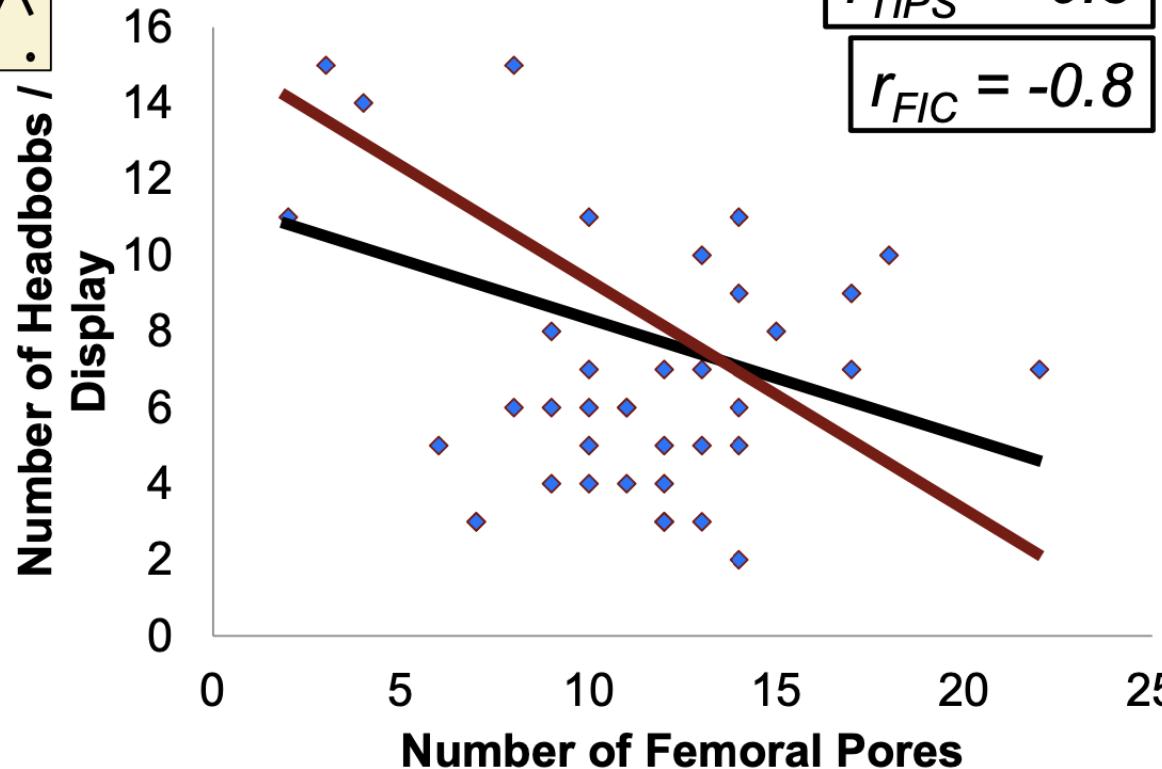
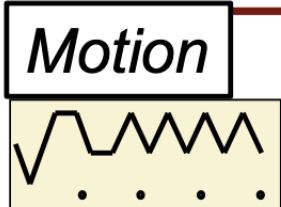


# Motion & chemical signals



**Chemicals**

# Motion & chemical signals

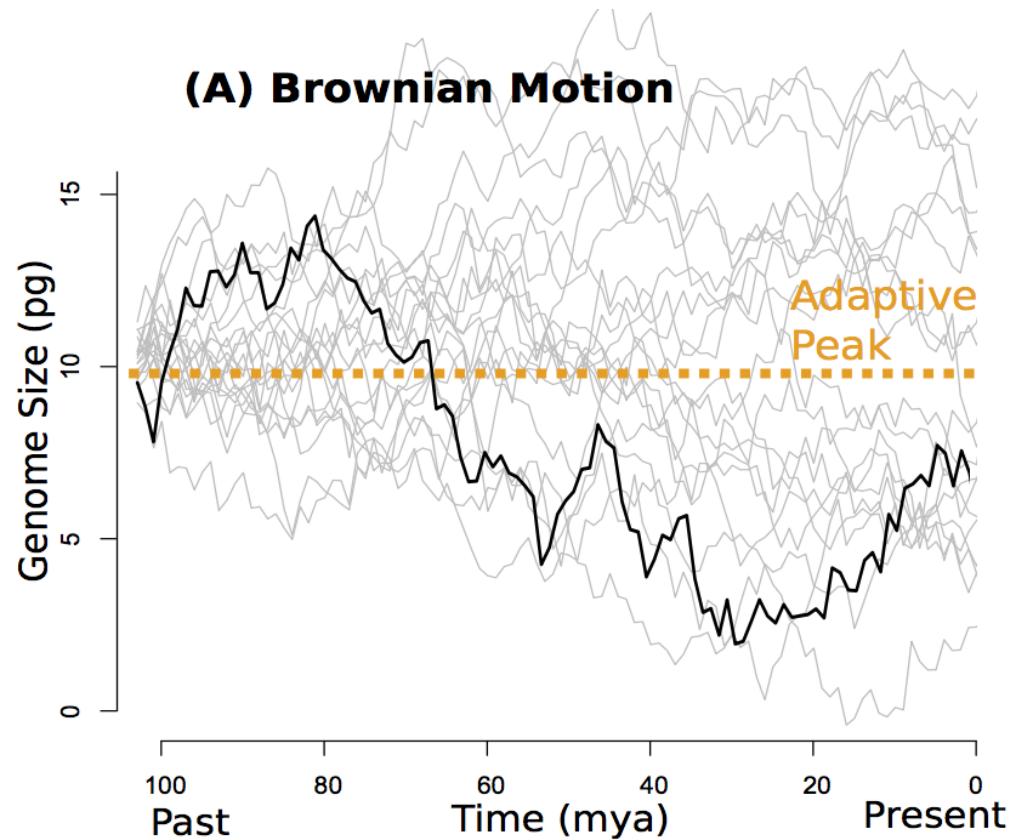


*Chemicals*

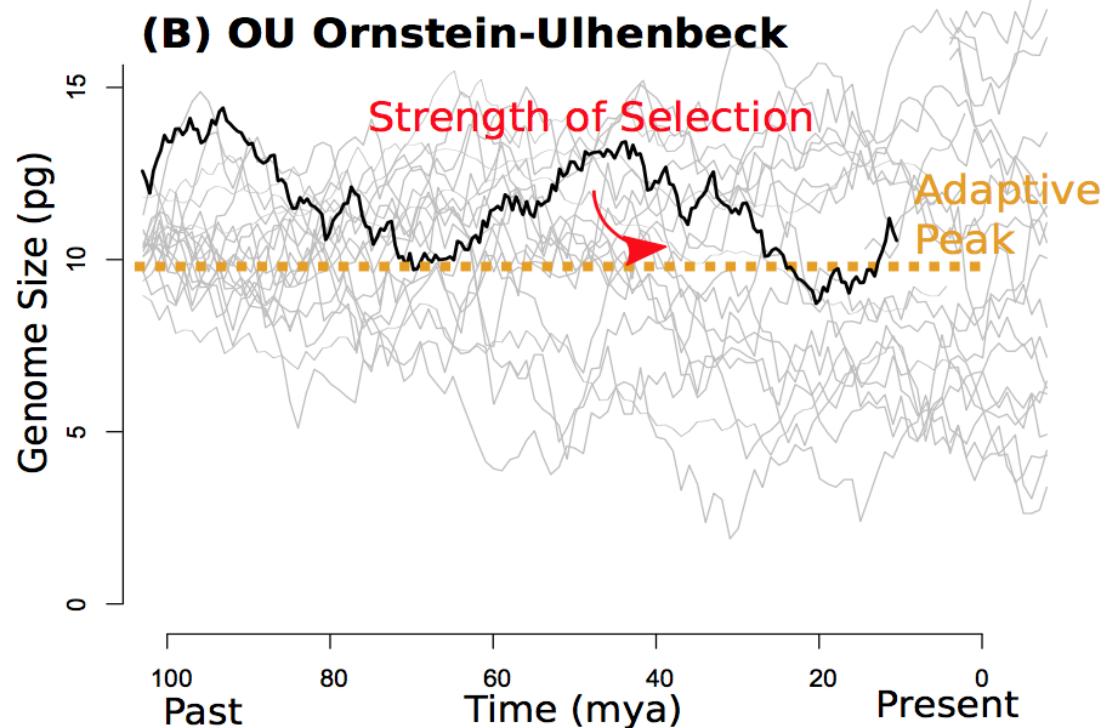
*Trade-off between motion and chemicals  
Phylogeny clarifies relationship between traits.*

BM may not be a good model of :

1. Most forms of selection
2. Phenotypic plasticity



$$\text{BM: } \text{var}(\varepsilon)_{ij} = \sigma^2 t_a$$



$$\text{OU: } \text{var}(\varepsilon)_{ij} = \gamma \exp[-\alpha t_{ij}]$$