

Discussion

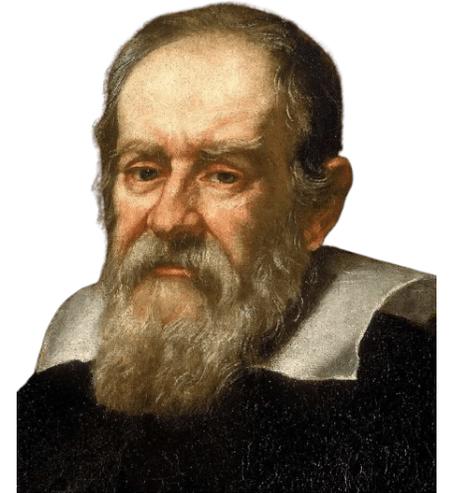
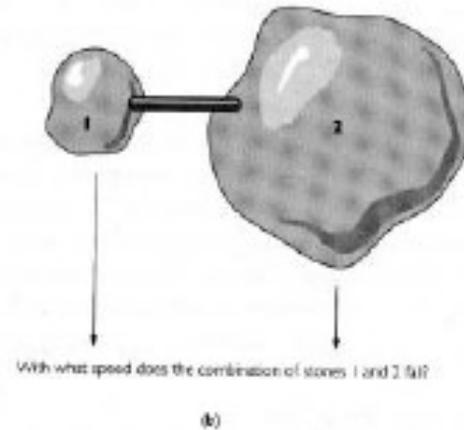
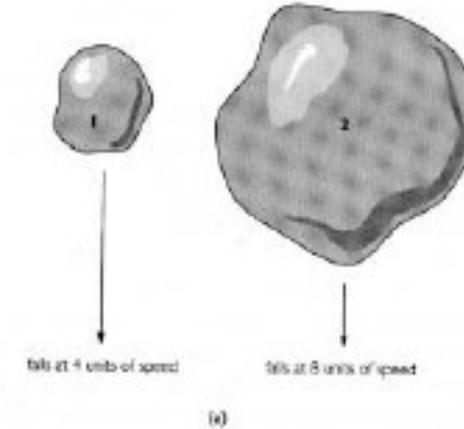
- Have you used theory or modelling already? How has it influenced your work?
- What (if any) is the difference between a model, a theory and a simulation?
- Do all models aim to answer the same type of question? If not, what are the different categories?
- What makes a good model versus a bad model?
- What is your aesthetic experience of modelling?

What does theory look like?

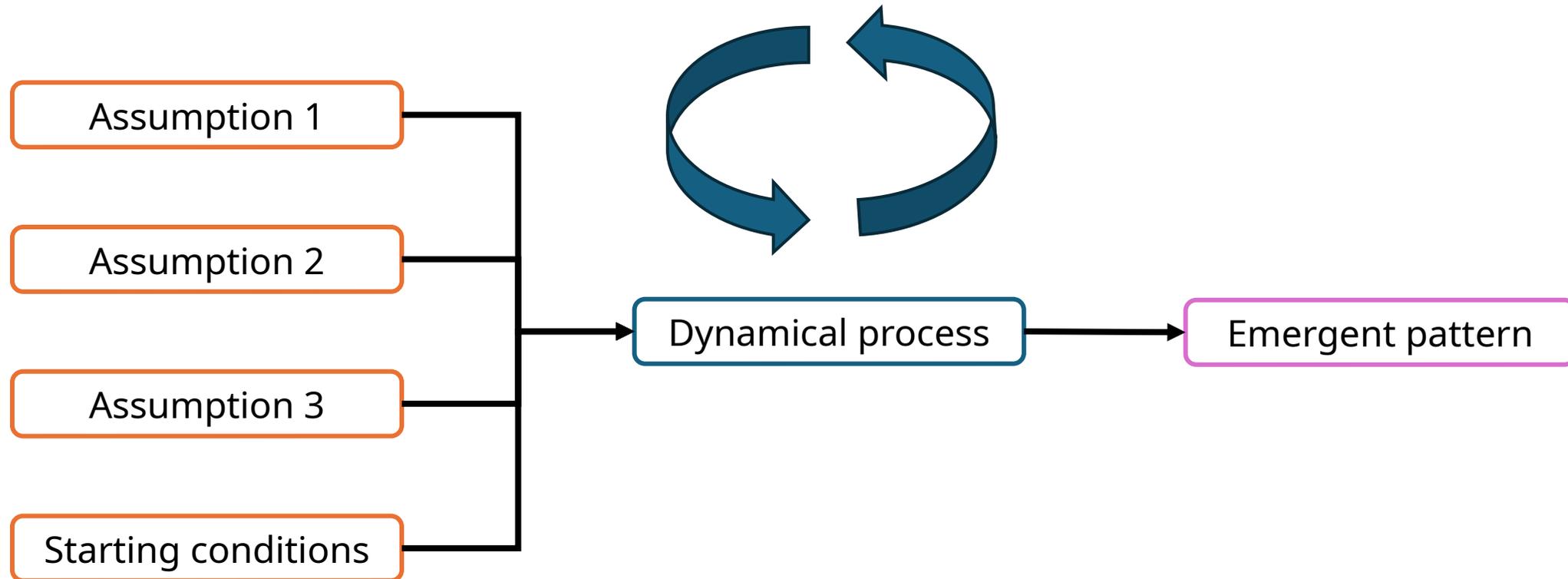
$$\frac{dx_i}{dt} = x_i \left(\sum_{j=1}^n \alpha_{ij} x_j \right)$$

'That's all Greek to me...'

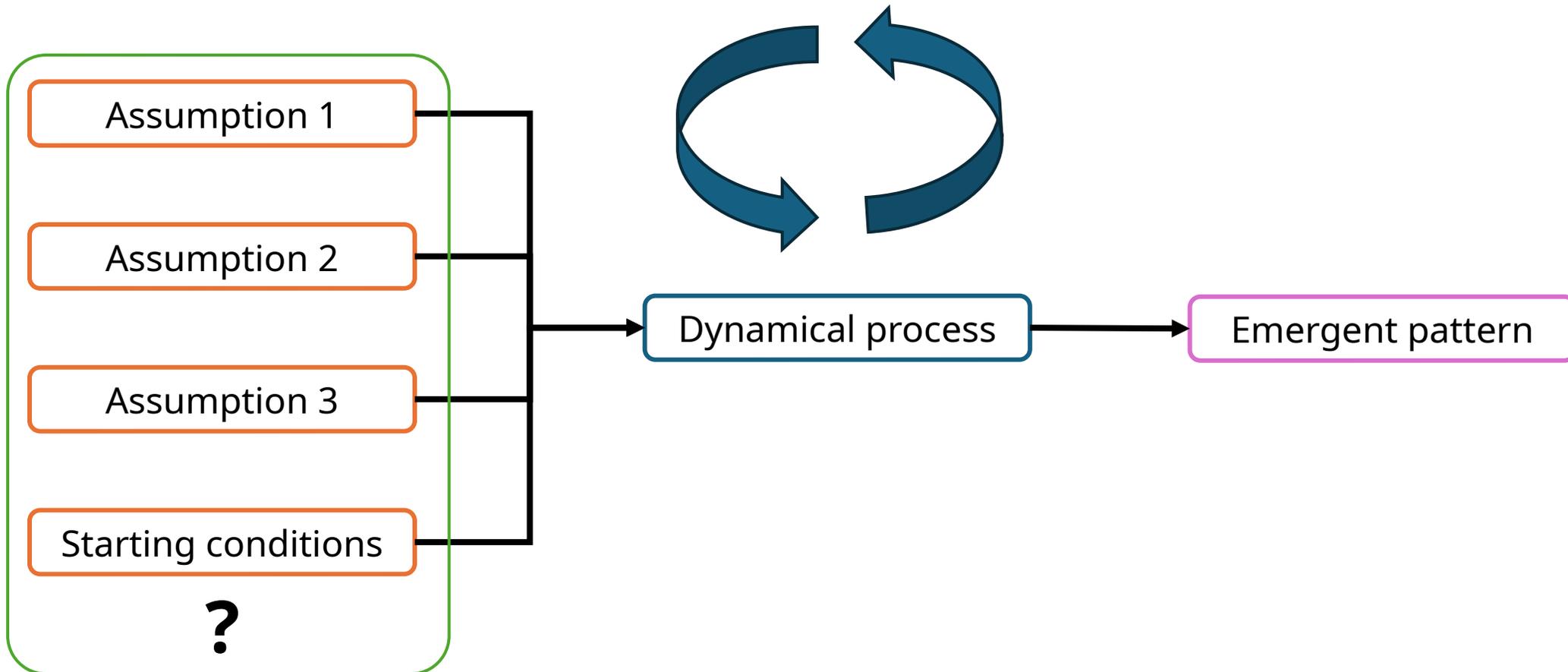
Galileo's gedankenexperiment



The 'emergence paradigm' for systemising theoretical approaches

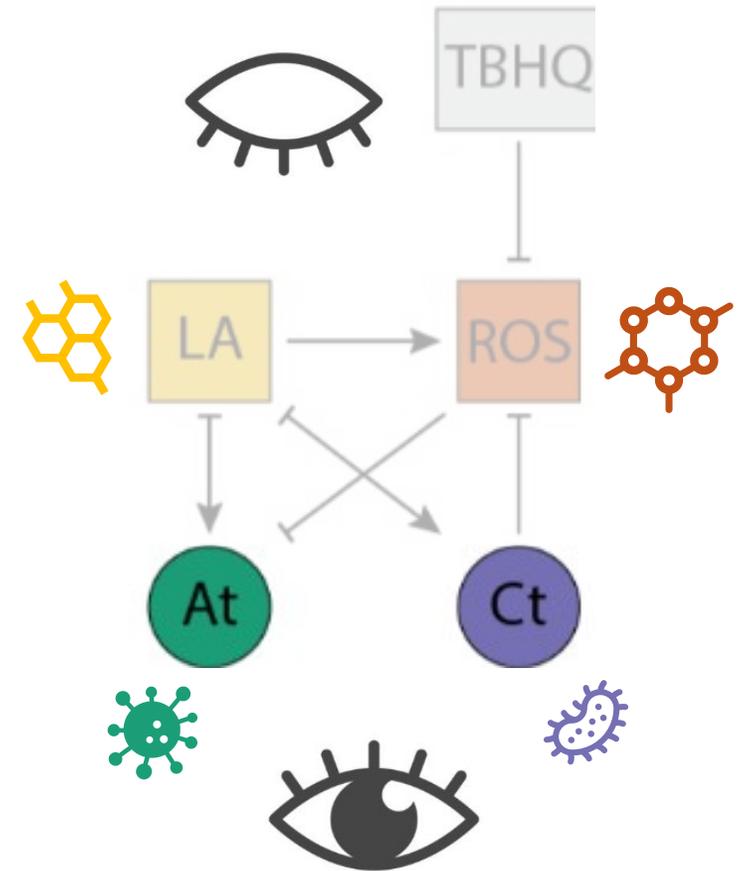
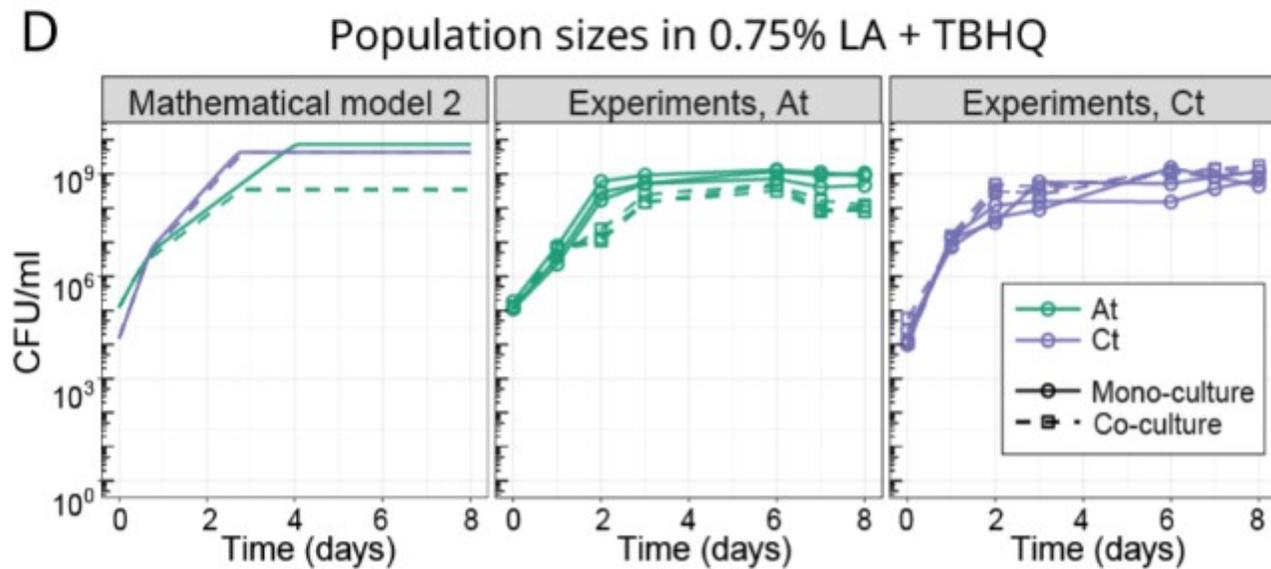
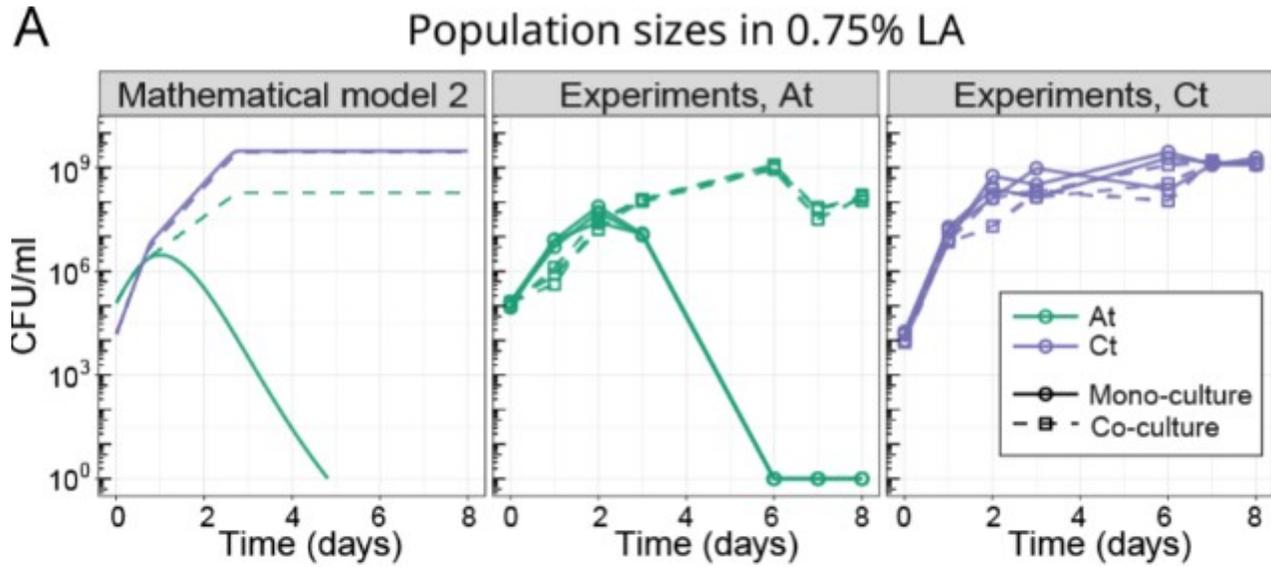


Identifying underlying mechanisms based on emergent patterns

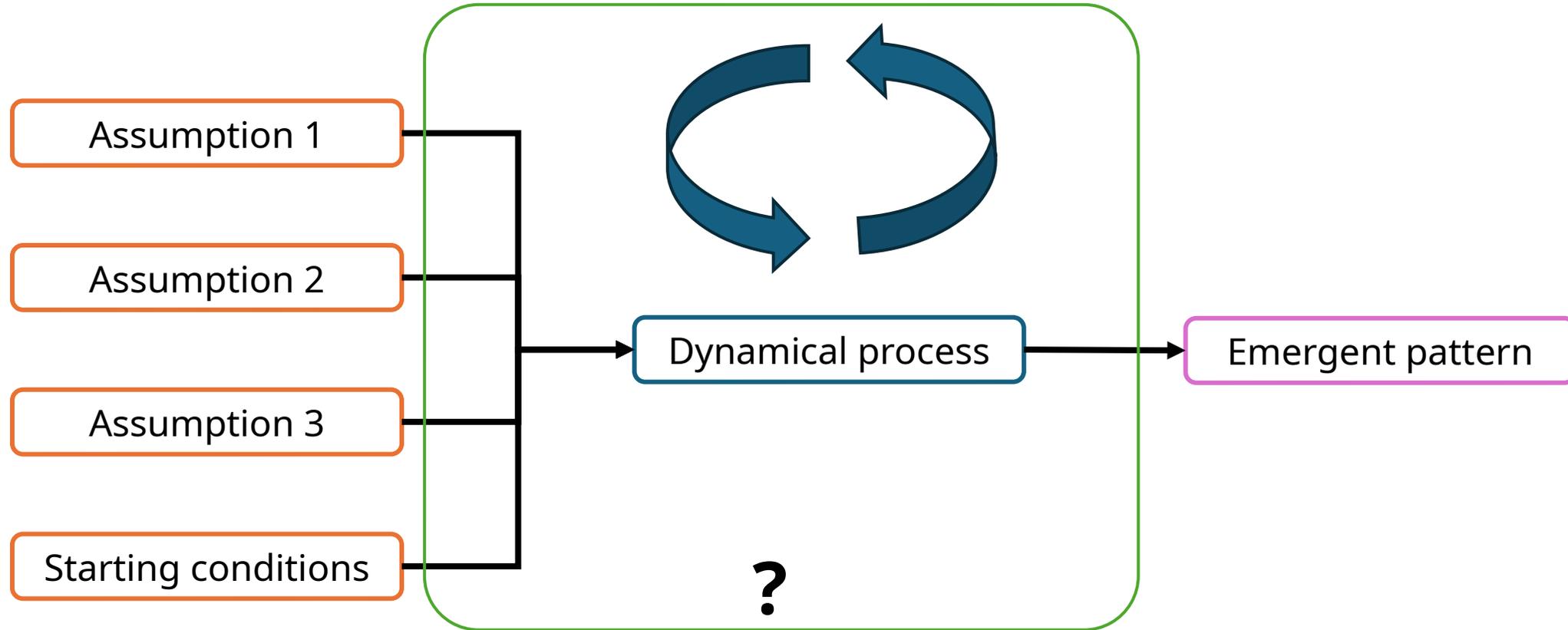


Identifying hidden mechanisms

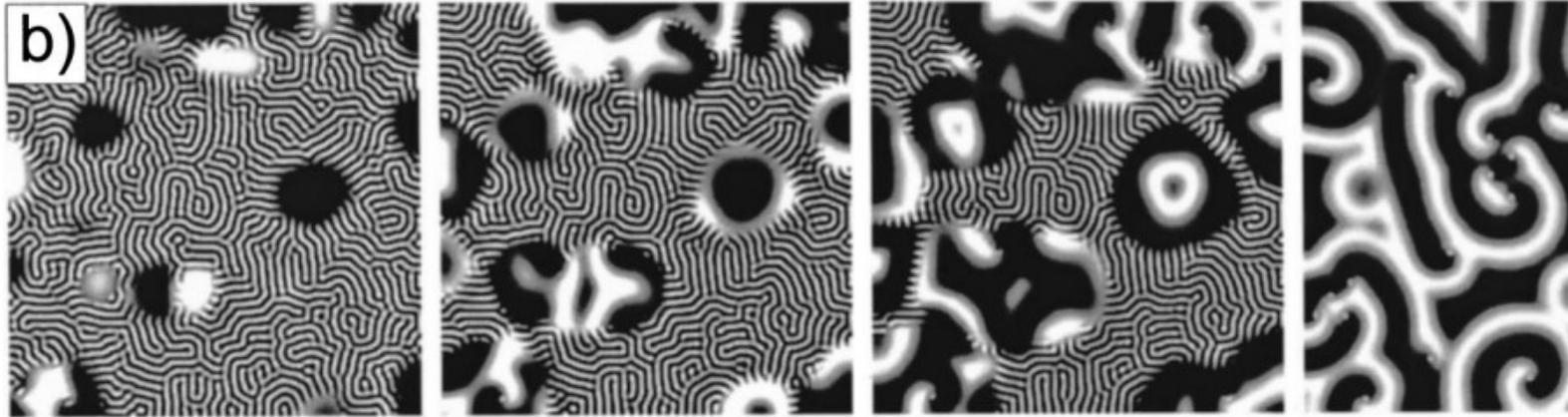
Di Martino et al. 2024, PLOS Bio.



Understanding emergence

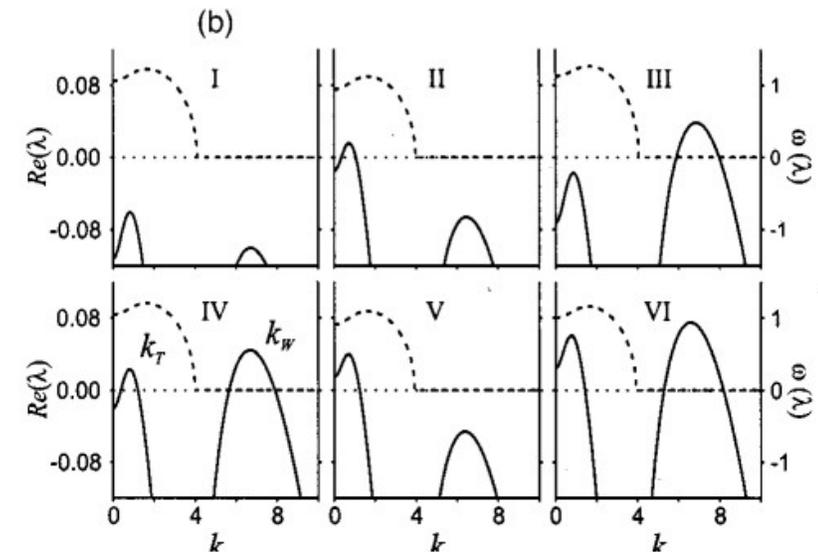
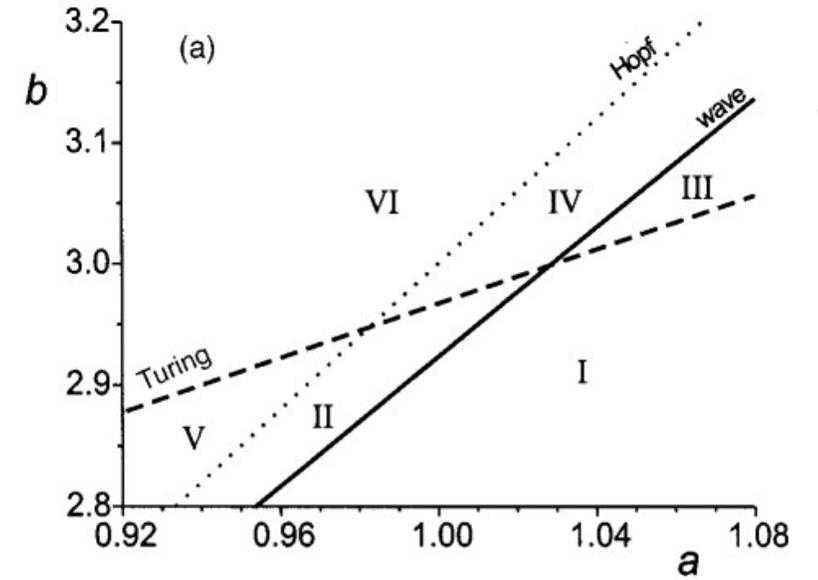


Understanding emergence: Turing patterns



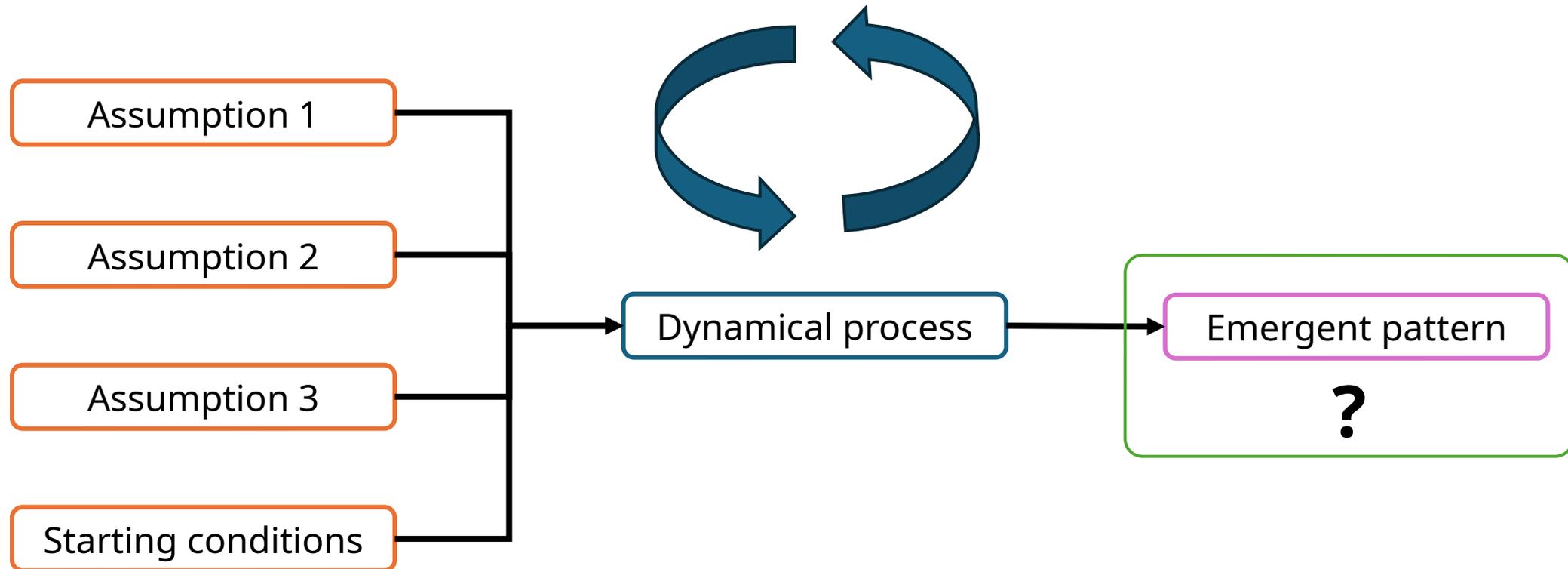
$$\frac{\partial z}{\partial \tau} = D_z \nabla^2 z + x - z, \quad (14)$$

$$\frac{\partial r}{\partial \tau} = D_r \nabla^2 r + \frac{1}{\epsilon_1} (cx - dr), \quad (15)$$

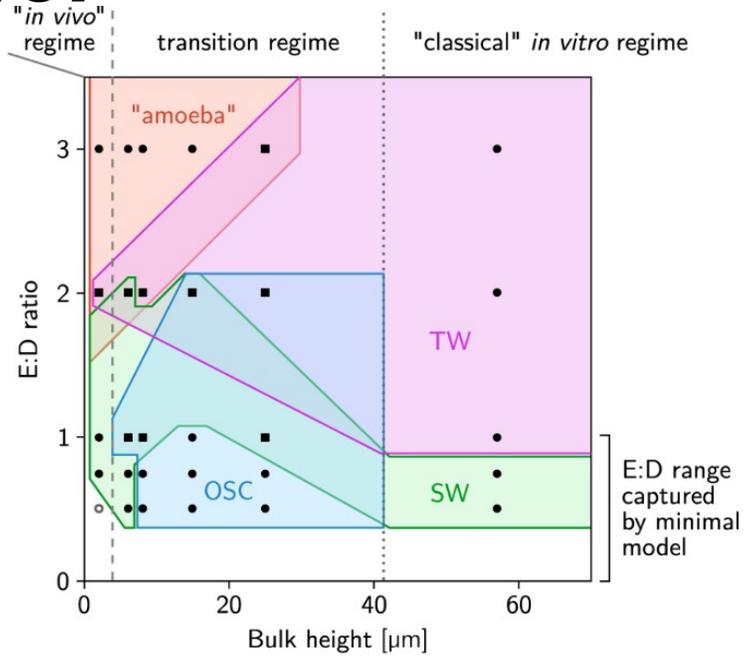


Yang et al. 2002, *J. Chem. Phys*

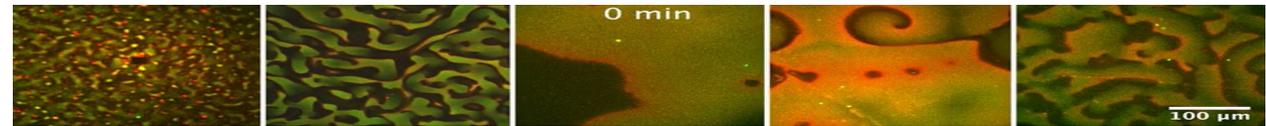
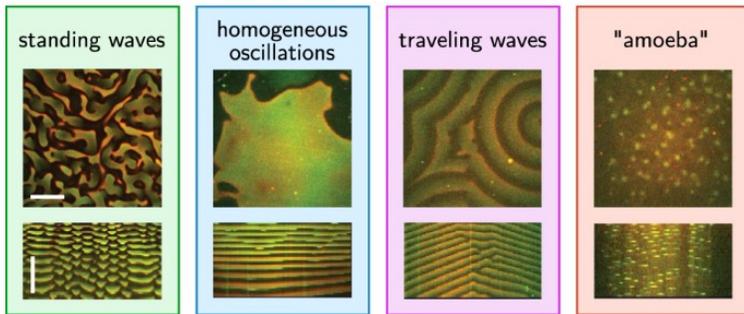
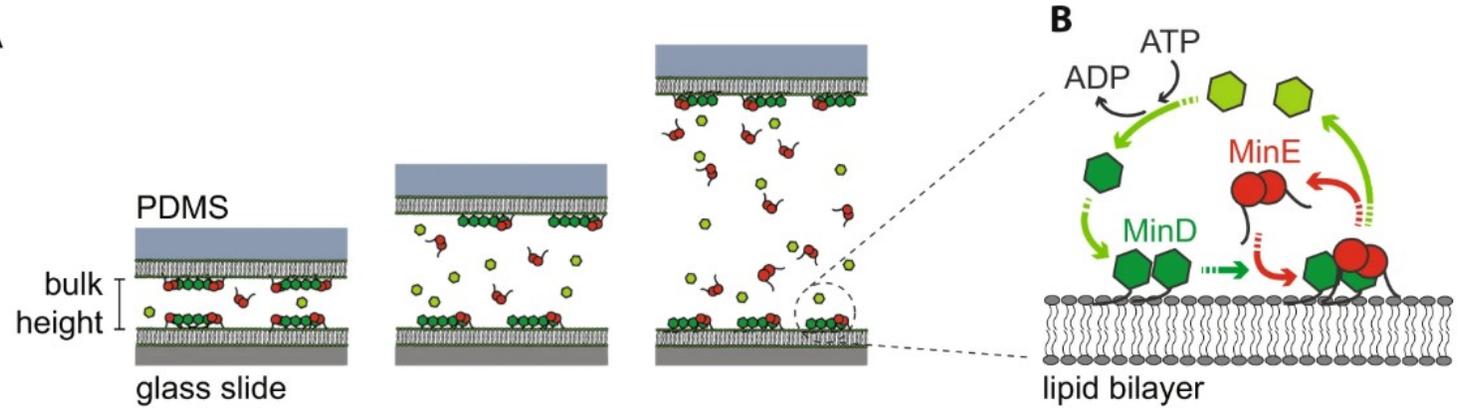
Identifying new patterns to search for



Identifying new patterns to search for

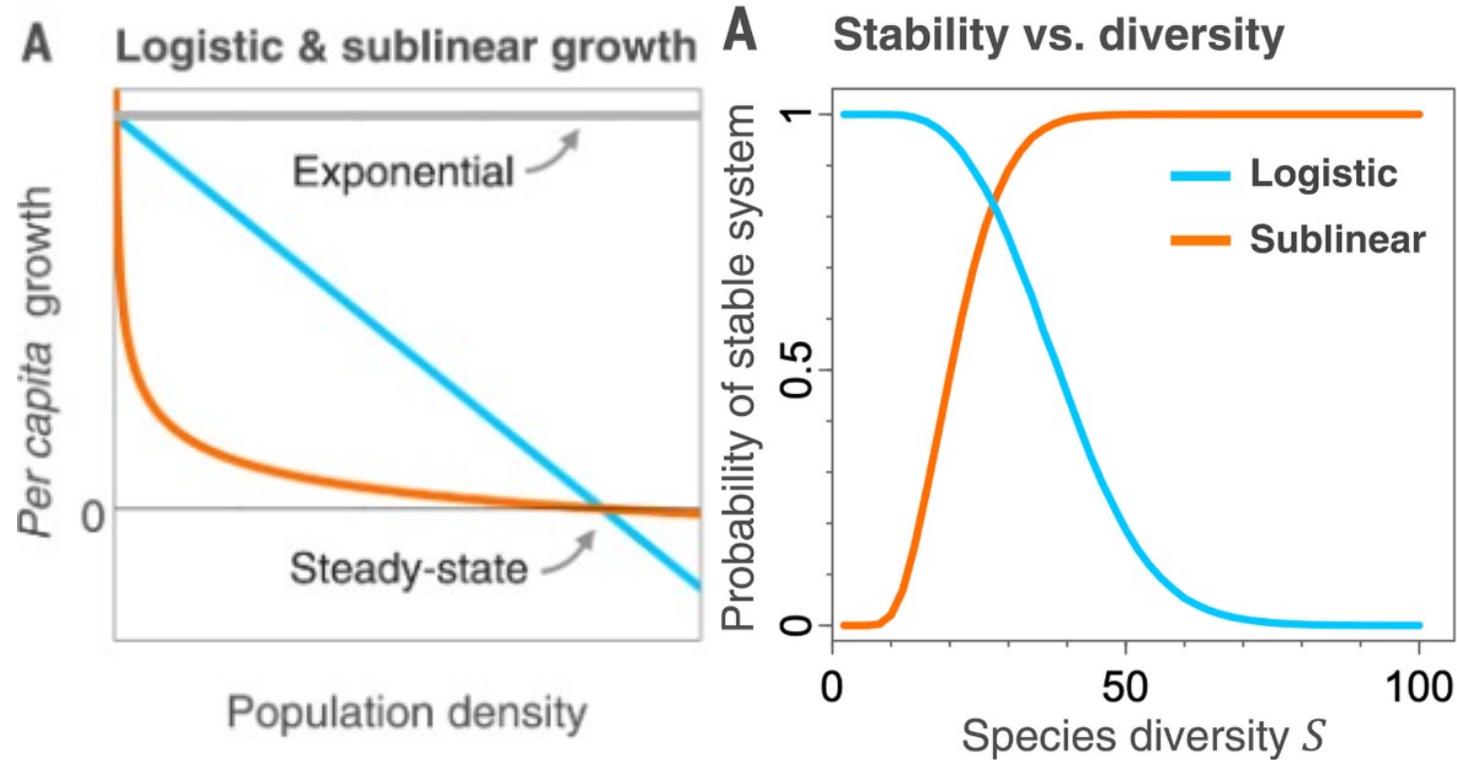


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Brauns et al. 2021, *Nat. Comms.*

The sins of theory: Faulty assumptions



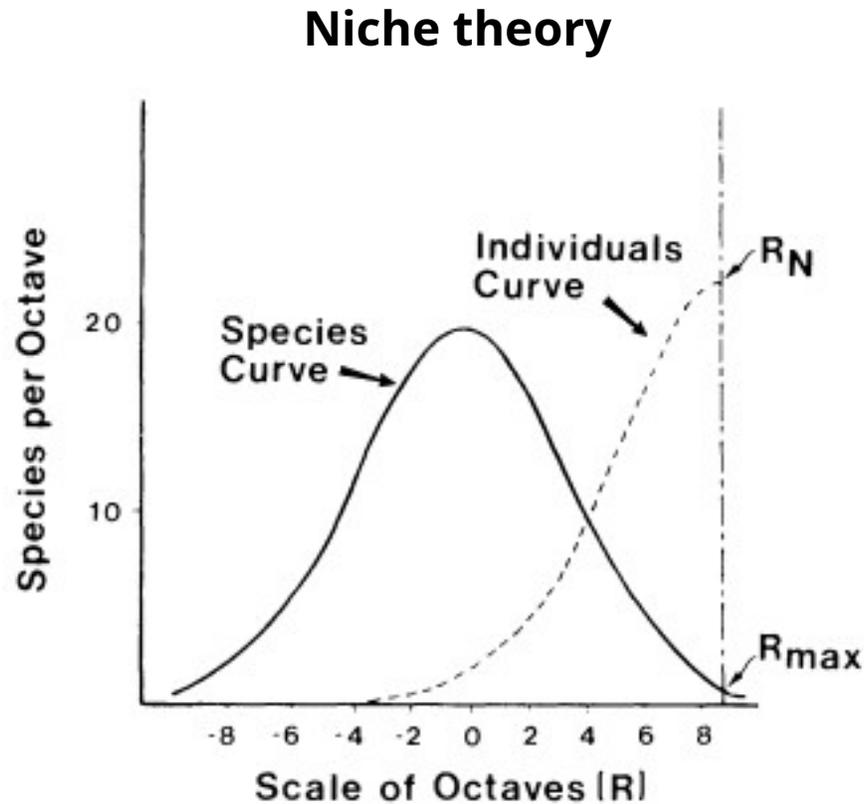
$$P_i(B_i) = \begin{cases} (r_i - z_i)B_i(1 - B_i/K) & (\text{logistic}) \\ r_i B_0^{1-k} B_i^k - z_i B_i & (\text{sublinear}) \end{cases}$$

Hatton et al. 2024, *Science*
(current references: 121)

"Here we clarify why SG [Sublinear-Growth] models do not solve the paradox of species coexistence. This is because, in the SG model, coexistence emerges from an unrealistic property, in which population per capita growth rates tend to infinity at low abundance, preventing species from ever going extinct due to competitive exclusion."

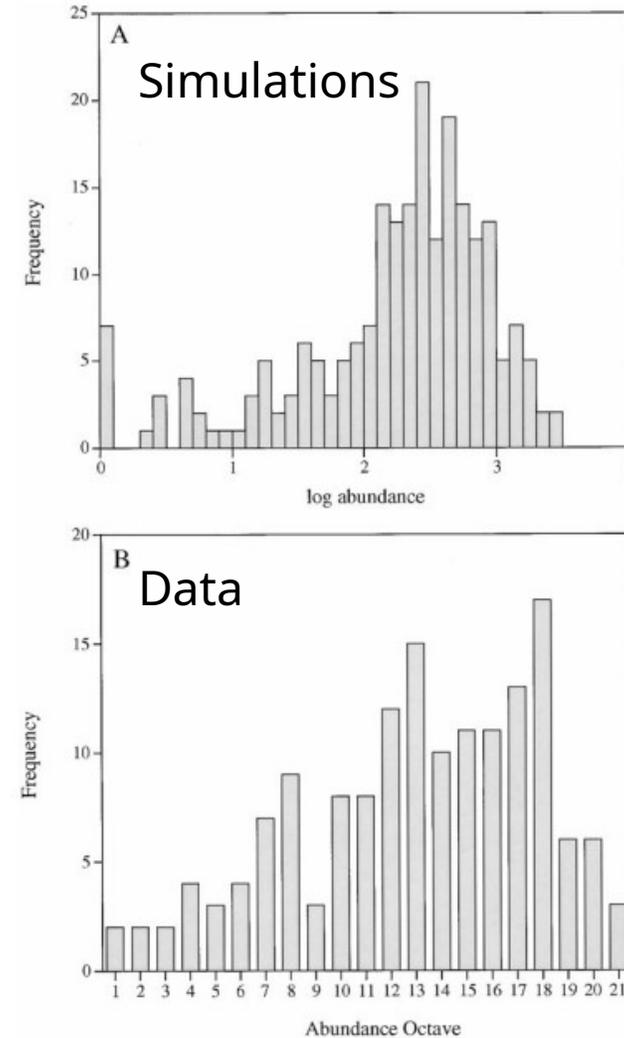
Aguadé-Gorgorió et al. 2024, *Oikos*
(current references: 7)

The sins of theory: Indistinguishable explanations



Bell 2000, *Am. Nat.*

Neutral theory



Sugihara 1980, *Am. Nat.*

The sins of theory: Obfuscation by mathematics

$$\kappa_i > \rho_i \left(\mathbf{P}^{\{-i\}} \right)^{-1} \boldsymbol{\kappa}^{\{-i\}}, \quad (44)$$

which is a direct generalization of the two-species condition 41 taking account of the indirect effects through the inverse matrix $\left(\mathbf{P}^{\{-i\}} \right)^{-1}$. Defining $\bar{\rho}^{\{-i\}}$ to be the average of the rows of $\mathbf{P}^{\{-i\}}$ and $\bar{\kappa}^{\{-i\}}$ to be the average the resident κ 's (i.e. the average of the elements of $\boldsymbol{\kappa}^{\{-i\}}$), the invader growth rate can now be written in terms of average fitness differences and stabilizing terms as follows

$$\frac{r_i}{b_i} = \kappa_i - \bar{\kappa}^{\{-i\}} + \left(\bar{\rho}^{\{-i\}} - \rho_i \right) \left(\mathbf{P}^{\{-i\}} \right)^{-1} \boldsymbol{\kappa}^{\{-i\}}. \quad (45)$$

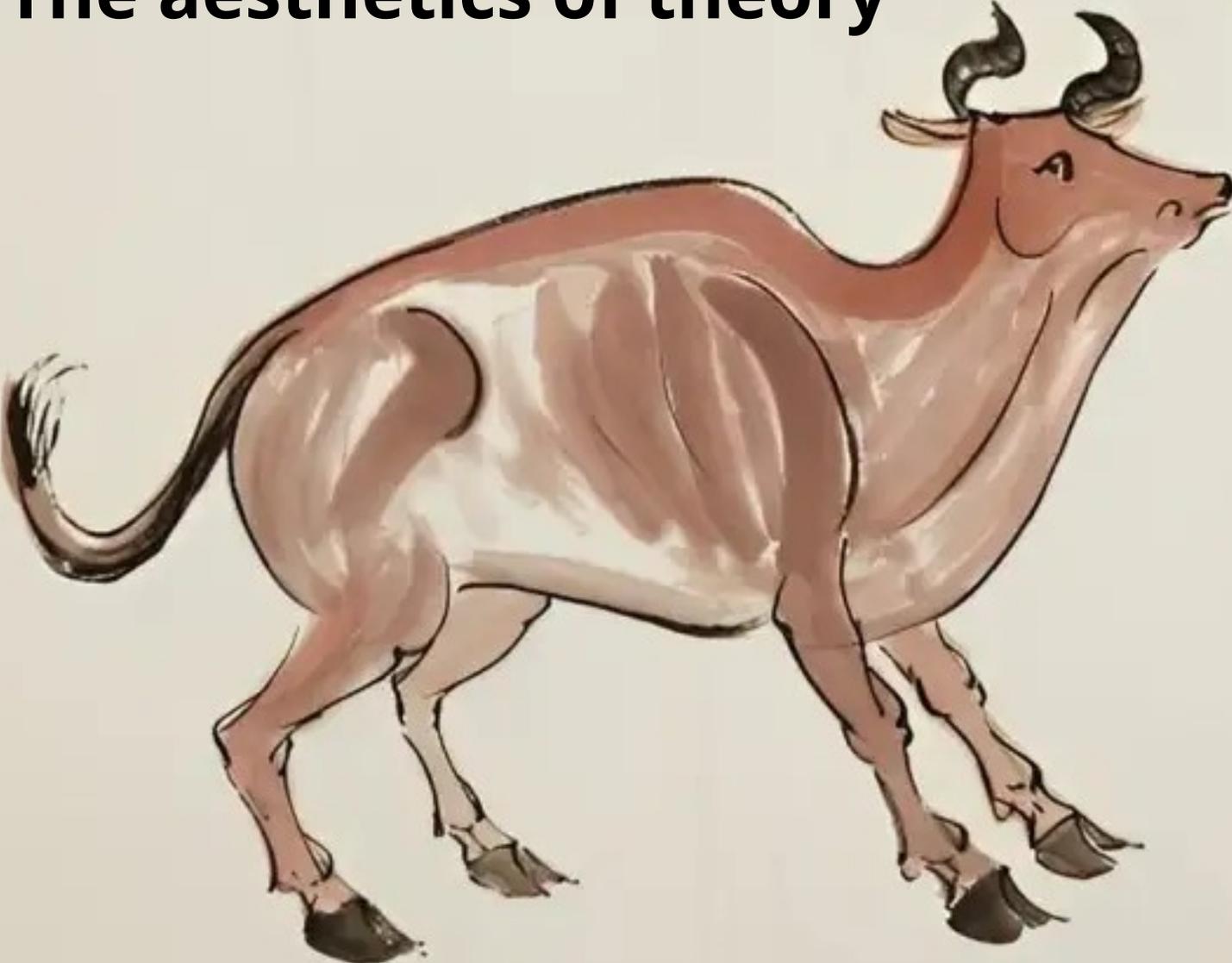
In Chesson (2000b), a general approximate invasion formula for diffuse competition is given as

$$\frac{r_i}{b_i} \approx \kappa_i - \bar{\kappa}^{\{-i\}} + \frac{D(1 - \rho)}{n - 1}, \quad (46)$$

where ρ is a common overlap measure for all pairs of species, and D is a constant depending on the circumstances, as derived in Chesson (1994). Expression 45 now allows the exact result for diffuse competition to be stated as

$$\frac{r_i}{b_i} = \kappa_i - \bar{\kappa}^{\{-i\}} + \frac{\bar{\kappa}^{\{-i\}}(1 - \rho)}{1 + (n - 2)\rho}. \quad (47)$$

The aesthetics of theory



Reading list/references

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