

# An individual-based model for the Lenski experiment, and the deceleration of the relative fitness

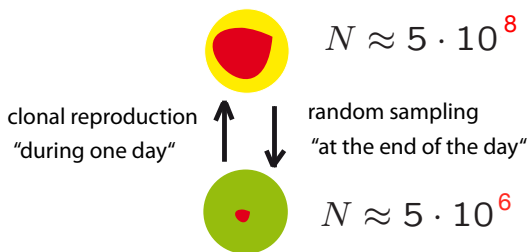
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Joint work with Noemi Kurt, Anton Wakolbinger and Linglong Yuan

27-01-2016



# The Lenski experiment (one day cycle)



# Relative fitness

## Measuring adaptation

- ▶ A population of size  $A_0$  of the unevolved strain and a population of size  $B_0$  of the evolved strain perform a direct competition.
- ▶ The respective population sizes at the end of the day are denoted by  $A_1$  and  $B_1$ .
- ▶ The (empirical) *relative fitness*  $F(B|A)$  of strain  $B$  with respect to strain  $A$  is

$$F(B|A) = \frac{\log(B_1/B_0)}{\log(A_1/A_0)}.$$

## Lenski, Travisano, PNAS, 1994

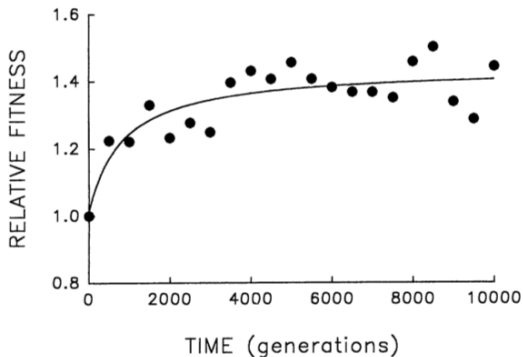
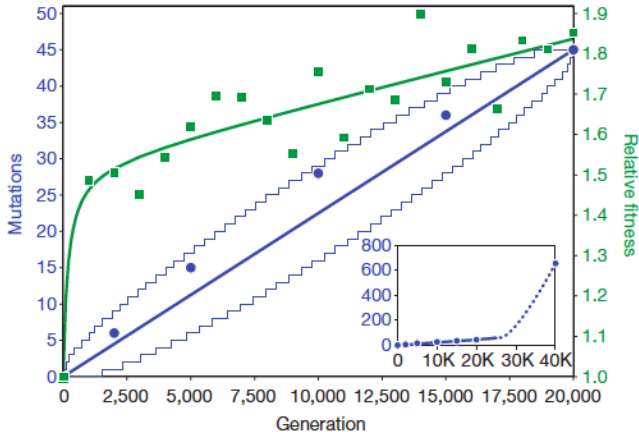


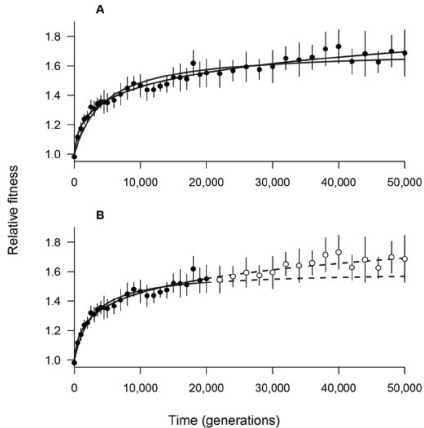
FIG. 4. Trajectory for mean fitness relative to the ancestor in one population of *E. coli* during 10,000 generations of experimental evolution. Each point is the mean of three assays. Curve is the best fit of a hyperbolic model.

# Barrick, Yu, Yoon, Jeong, Oh, Schneider, Lenski, Kim, Nature 2009

$$\omega(t) = 1 + at/(t + b)$$



# Wiser, Ribeck, Lenski, Science express 13-11-12



**Fig. 2. Comparison of hyperbolic and power-law models.** (A) Hyperbolic (red) and power-law (blue) models fit to the set of mean fitness values (black symbols) from all 12 populations. (B) Fit of hyperbolic (solid red) and power-law (solid blue) models to data from first 20,000 generations only (filled symbols), with model predictions (dashed lines) and later data (open symbols). Error bars are 95% confidence limits based on the replicate populations.

$$w(t) = (1 + ct)^{1/2g}$$

## Questions

- ▶ Which curve describes better the trajectory of the relative fitness?
- ▶ Why is the relative fitness decelerating?

## Possible explanations

- ▶ Clonal interference
- ▶ Epistasis
- ▶ The design of the experiment

# The daily cycle model<sup>1</sup>

## Information about the experiment

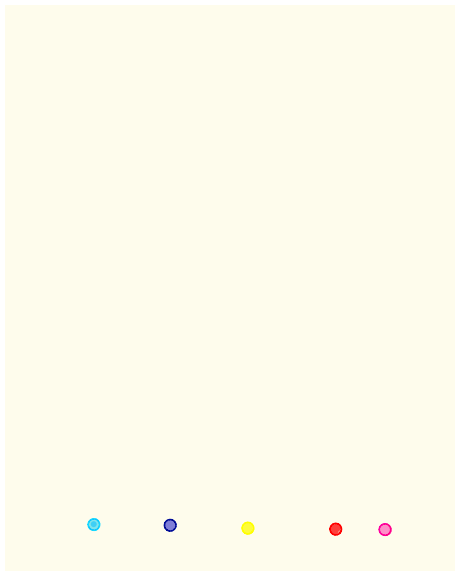
- ▶ At the beginning of each day there are  $N$  individuals.
- ▶ Within each day individuals split at constant rate.
- ▶ The reproduction process will stop when the glucose has been consumed. (This happens when there are around  $100N$  individuals).
- ▶  $N$  individuals out of the  $\sim 100N$  are uniformly sampled without replacement, to be starting individuals at the next day.

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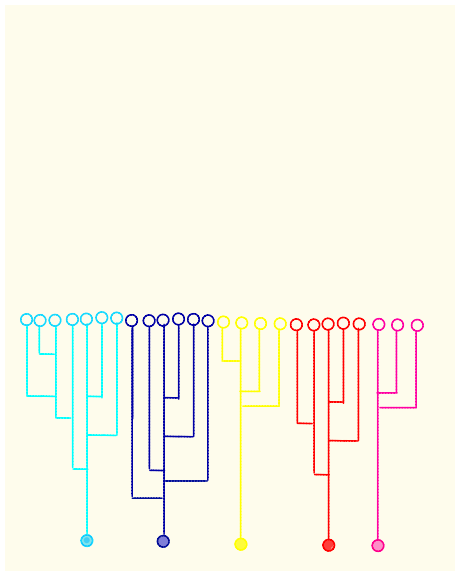
<sup>1</sup>An individual-based model for the Lenski experiment, and the deceleration of the relative fitness. Adrian Gonzalez Casanova, Noemi Kurt, Anton Wakolbinger and Linglong Yuan. (2015) arXiv 1505.0175



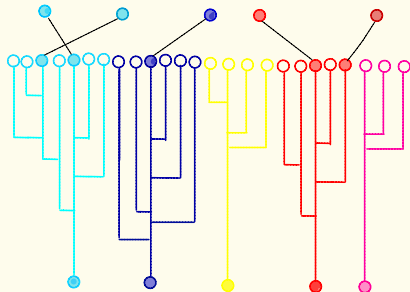
# Pruned Yule trees



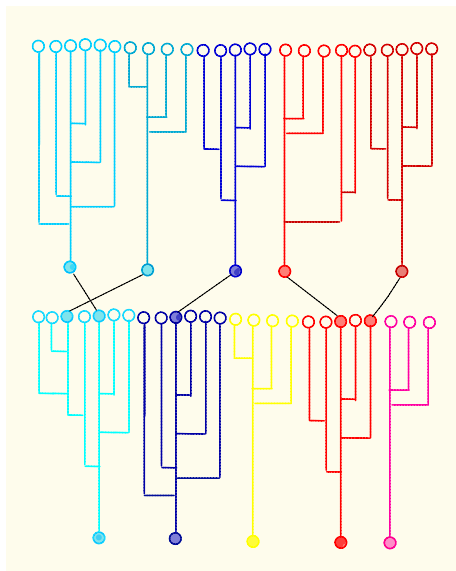
# Pruned Yule trees



# Pruned Yule trees



# Pruned Yule trees



# Inside a day

Let  $Y_{i,j}(t)$  be a pure birth process with rate  $r \in \mathbb{R}^+$ , for every  $i \in \mathbb{N}$  and  $j \in \{1, 2, \dots, N\}$ . (Yule Processes with parameter  $r$ ).

The total population size at time  $t$  of day  $i$  is

$$\sum_{j=1}^N Y_{i,j}(t)$$

## Stopping rule

Each day, the reproduction stops at time  $\sigma$  , where

$$\sigma = \inf\{t : E[\sum_{j=1}^N Y_{i,j}(t)] = \gamma N\}.$$

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$$\sigma = \frac{\ln(\gamma)}{r}.$$

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$$\sigma = \frac{\ln(\gamma)}{r}.$$

The total population size at the end of the day is

$$\sum_{j=1}^N Y_{i,j}(\sigma) \sim \gamma N.$$



## Sampling rule

To go from one day to the next, we sample uniformly at random  $N$  individuals (out of  $\sim \gamma N$ ), and we say that each sampled individual is a root of a Yule tree in the next day.

## Introducing selective advantage

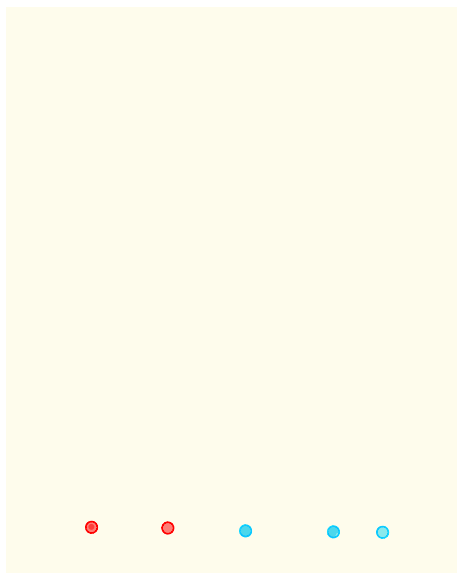
- ▶ Assume that some individuals reproduce at rate  $r + \varrho_N$  (mutants), while other reproduce at rate  $r$  (basis population).

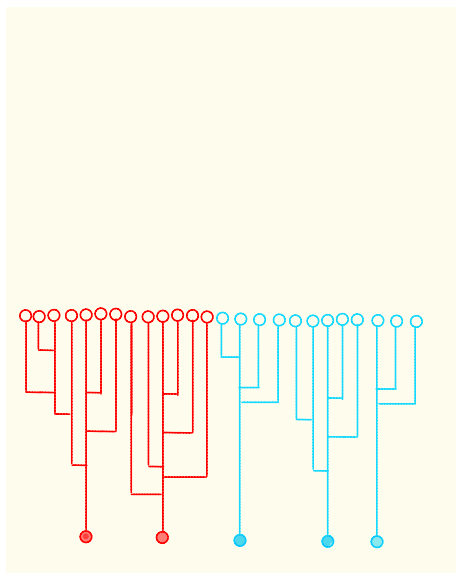
## Introducing selective advantage

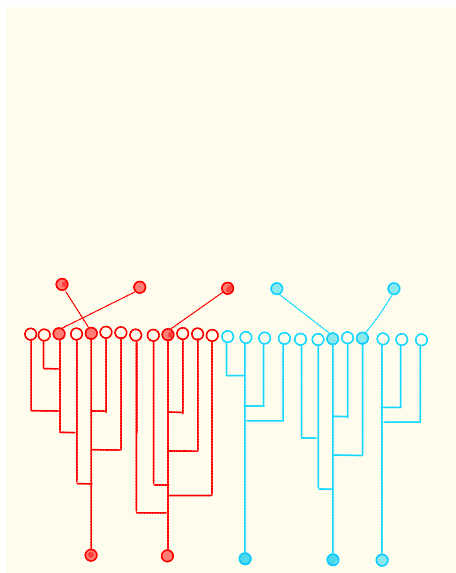
- ▶ Assume that some individuals reproduce at rate  $r + \varrho_N$  (mutants), while other reproduce at rate  $r$  (basis population).
- ▶ Let  $M_i(t)$  be the number of mutants at time  $t$  of day  $i$ .
- ▶ We are interested in the process

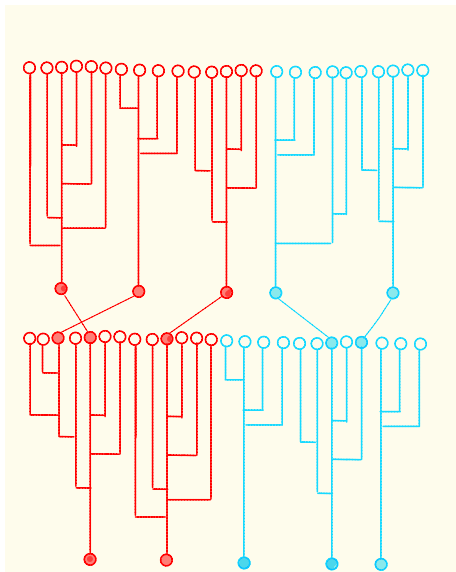
$$\{K_i\}_{i \in \mathbb{N}} := \{M_i(0)\}_{i \in \mathbb{N}},$$

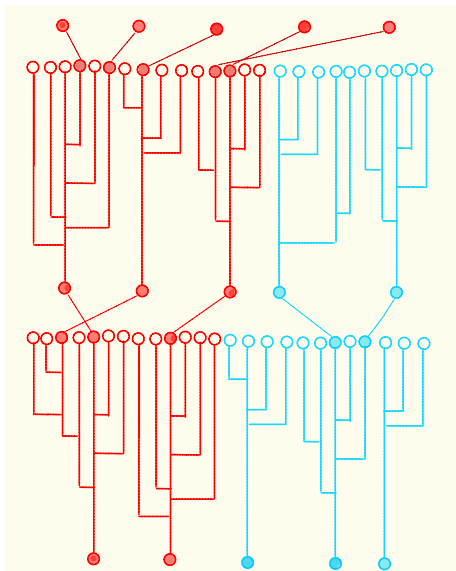
which is constructed recursively using uniform sampling.













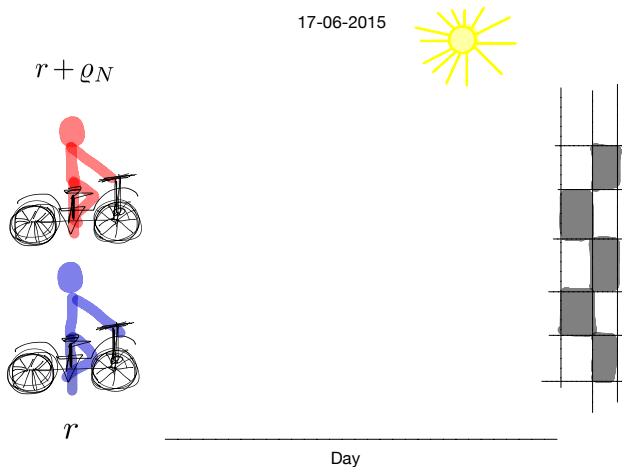
# Selective advantage

Basic population reproduces at rate  $r$ .

Mutants reproduce at rate  $r + \varrho_N$ .

$$\mathbb{E}[K_1 | K_0 = 1] = 1 + \varrho_N \frac{\log \gamma}{r} + o(\varrho_N).$$

# Race until the sun is gone

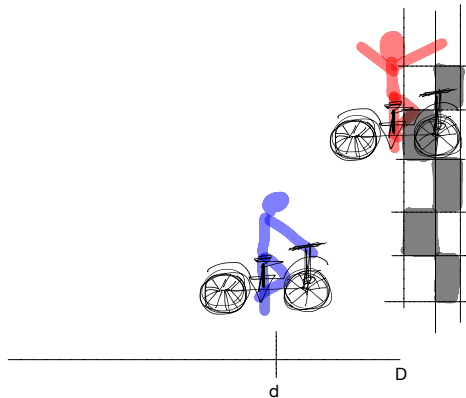


# Race until the sun is gone

17-06-2015



$$r + \varrho_N$$

 $r$ 


# Race until the sun is gone

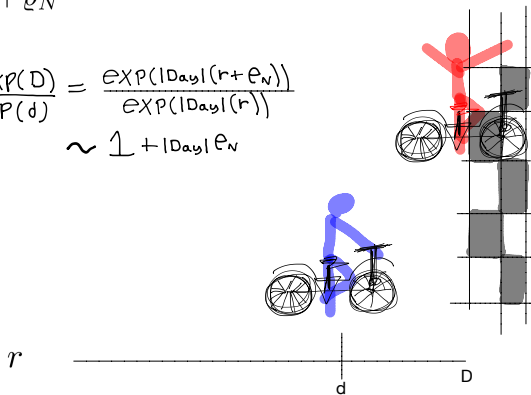
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$$r + \varrho_N$$

$$\frac{\exp(D)}{\exp(d)} = \frac{\exp(|\text{Day}|(r + e_N))}{\exp(|\text{Day}|(r))}$$

$$\sim 1 + |\text{Day}| e_N$$



# Race until the sun is gone

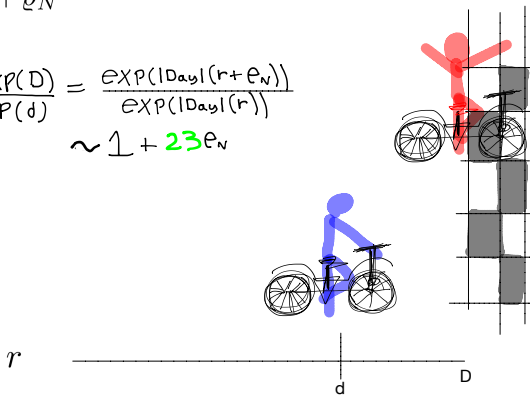
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$$r + \rho_N$$

$$\frac{\exp(D)}{\exp(d)} = \frac{\exp(|\text{Day}|(r + e_N))}{\exp(|\text{Day}|(r))}$$

$$\sim 1 + 23e_N$$

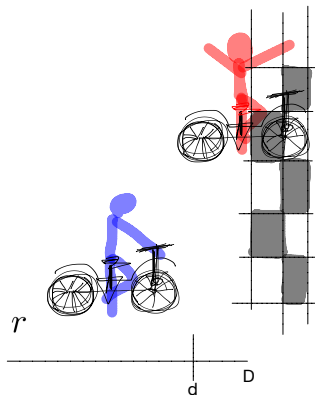


# Race until the sun is gone

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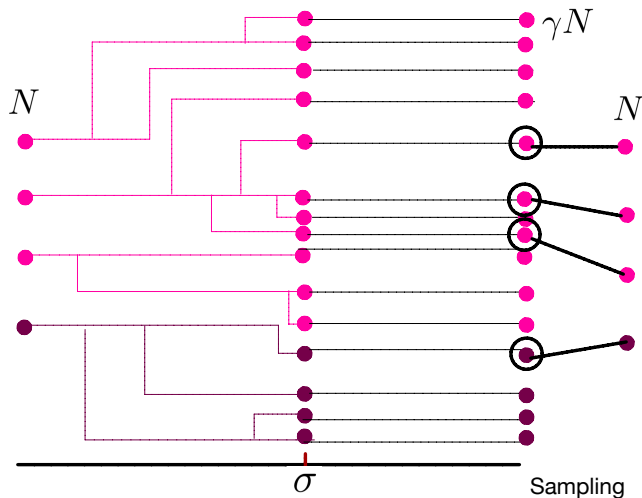
$$r + \rho_N$$



$$\frac{\exp(D)}{\exp(d)} = \frac{\exp(|\text{Day}|(r + \rho_N))}{\exp(|\text{Day}|(r))}$$

$$\sim 1 + \rho_N$$

# The *effective competition time*, and its dependence on $r$



Let  $\pi_N := \mathbb{P}(\exists i \in \mathbb{N} : K_i = N \mid K_0 = 1)$ , and  $\tau^N := \tau_{\text{fix}}^N \wedge \tau_{\text{ext}}^N$ .

### Theorem (Probability and speed of fixation)

*Under the assumptions of our model, as  $N \rightarrow \infty$ ,*

$$\pi_N \sim \frac{\gamma}{\gamma - 1} \frac{\varrho_N \log \gamma}{r}.$$

*Moreover, for any  $\delta > 0$  there exists  $N_\delta \in \mathbb{N}$  such that for all  $N \geq N_\delta$*

$$\mathbb{P}(\tau^N > \varrho_N^{-1-3\delta}) \leq (7/8)^{\varrho_N^{-\delta}}.$$



## The weak mutation - moderate selection model (Assumption A)

- i) Beneficial mutations add  $\varrho_N$  to the reproduction rate of the individual that suffers the mutation.
- ii) In each generation, with probability  $\mu_N$  there occurs a beneficial mutation. The mutation affects only one (uniformly chosen) individual, and every offspring of this individual also carries the mutation.
- iii) There exists  $0 < b < 1/2$ , and  $a > 3b$ , such that  $\mu_N \sim N^{-a}$  and  $\varrho_N \sim N^{-b}$  as  $N \rightarrow \infty$ .

$$\mu_N \ll \varrho_N$$

We define the fitness of the population at the beginning of day  $i$  with respect to that at the beginning of day 0 as

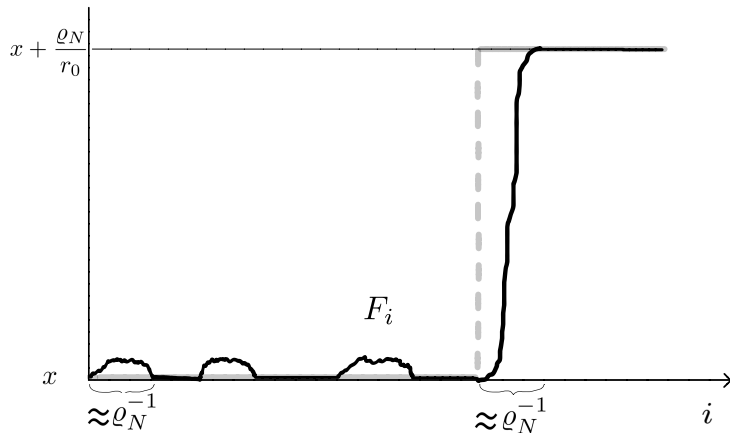
$$F_i := \frac{\log \frac{1}{N} \sum_{j=1}^N e^{R_{i,j}t}}{\log e^{r_0 t}}$$

where  $R_{i,j}, j = 1, \dots, N$  are the reproduction rates of the individuals present at the beginning of day  $i$ , and  $t$  is a given time for which the two populations are allowed to grow together.

If the whole population reproduces at the same rate ( $R_i$ ), then

$$F_i = \frac{R_i}{r_0}$$

where  $r_0 := R_0$ .



**Figure:** The number of attempts to go to fixation, when the reproduction rate of the basic population is  $x$ , is distributed Geometric with parameter  $\pi_N \sim \varrho_N \frac{C(\gamma)}{x}$ .

### Theorem (Convergence of the relative fitness process)

*Assume  $R_{0,j} = r_0$  for  $j = 1, \dots, N$ , and let  $(F_i)_{i \in \mathbb{N}_0}$  be the process of relative fitness. Then under Assumption A, the sequence of processes  $(F_{\lfloor (e_N^2 \mu_N)^{-1} t \rfloor})_{t \geq 0}$  converges in distribution as  $N \rightarrow \infty$  locally uniformly to the deterministic function*

$$f(t) = \sqrt{1 + \frac{\gamma \log \gamma}{\gamma - 1} \frac{2t}{r_0^2}}, \quad t \geq 0.$$

Table: Our model compared with Wiser et al.

	Our model	Wiser et al
Clonal interference	No	Yes
Epistasis	No	Yes
Design of the experiment	Yes	No
	$f(t) = (1 + \frac{2C(\gamma)t}{r_0^2})^{1/2}$	$w(t) = (1 + ct)^{1/2g}$

Thank you!