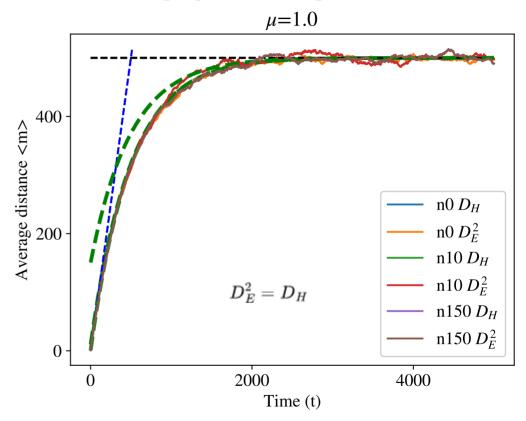


$$D_H = \sum_{i=1}^L |u_i - v_i|$$

$$D_E = ||\vec{u} - \vec{v}|| = \sqrt{\sum_{i=1}^{L} (u_i - u_j)^2}$$

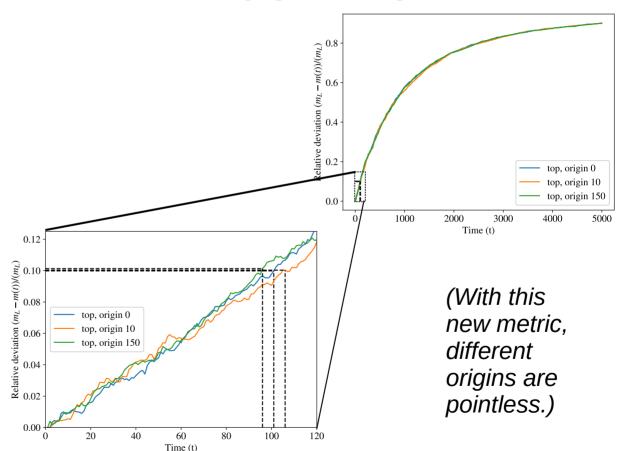


Using this metric only shifts the origin

When can we say that linear approximation is valid?

$$arepsilon := rac{m_L - m(t)}{m_L} = 0.1$$

$$m_L = \mu t$$



Developing m(t) to second order, one can obtain that

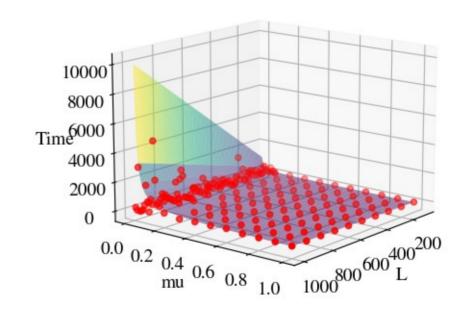
$$t_{arepsilon} = rac{2\lambdaarepsilon}{\mu}$$

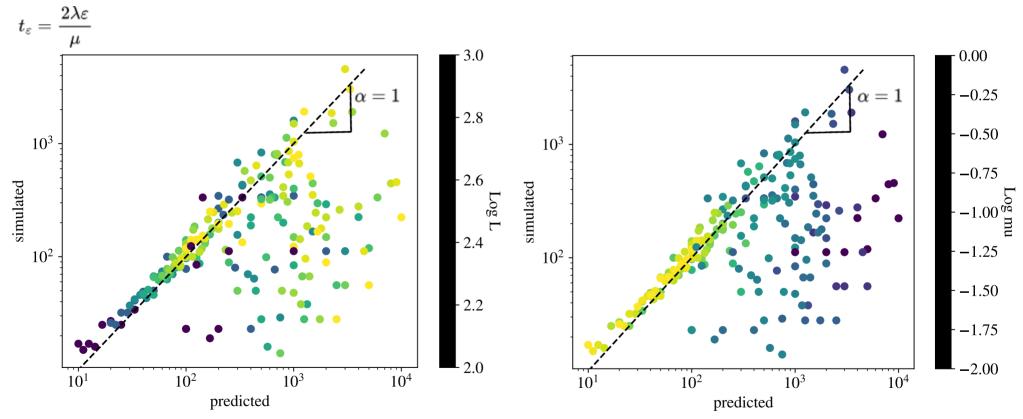
Simulations were carried and stopped when the mean number of mutations of the population was

$$m(t) \leq (1-\varepsilon)\mu t$$

then saving the value of t.

Surface → Theoretical result Points → Simulated





Although we have a good agreement, prediction fails for small values of μ

Open questions

The upcoming phase involves the duplication of sequences with a specified probability, denoted as "p." In instances where the probability is 1-p, sequences do not duplicate but may undergo mutations.

I have a struggle deciding which implementation we want for the process.

1-) Does each sequence undergo a predetermined duplication? (my most voted)

This scenario suggests that certain sequences possess advantages for replication based on their genotype.

2-) Is the duplication of each sequence a random event (with probability p)?

In this case, the replication process appears to be a stochastic one within the population, regardless of the specific genotypes present.