

Population genomics

2026-01-20





2011

2018

2023

Bachelor/Masters

PhD

Postdoc

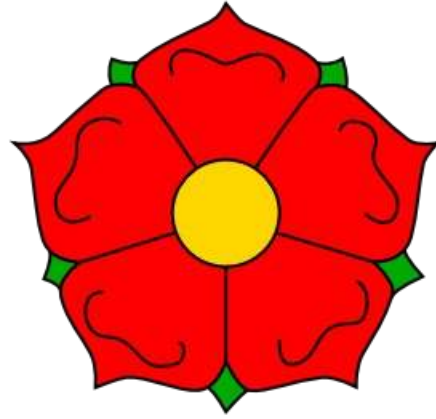


Outline for this evening

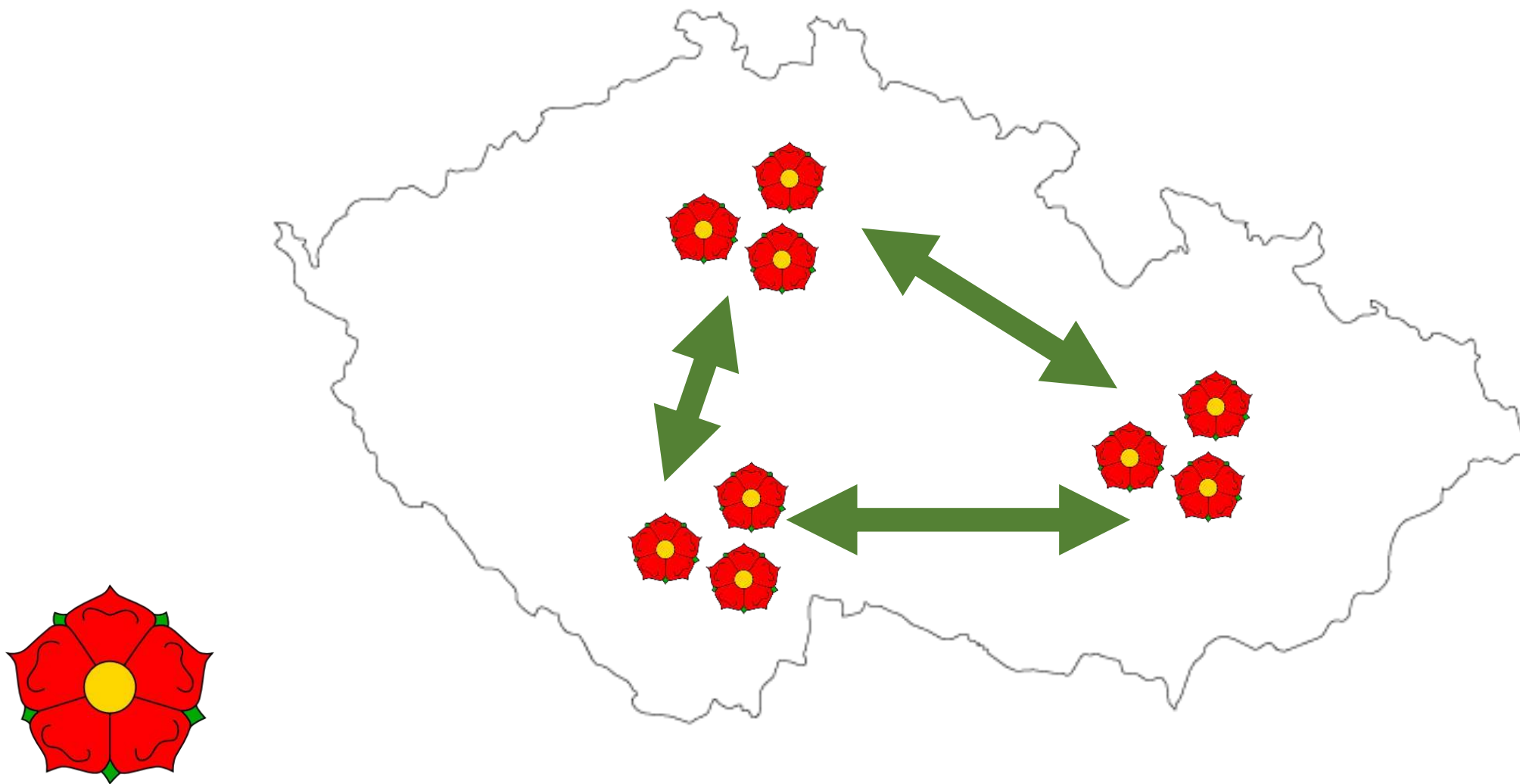
- Introduction to population genomics (~30 mins)
- Practical exercises
 - Part 1: Simulation based (~1 hour)
 - Part 2: Real world example (~1 hour)

What is population genomics?

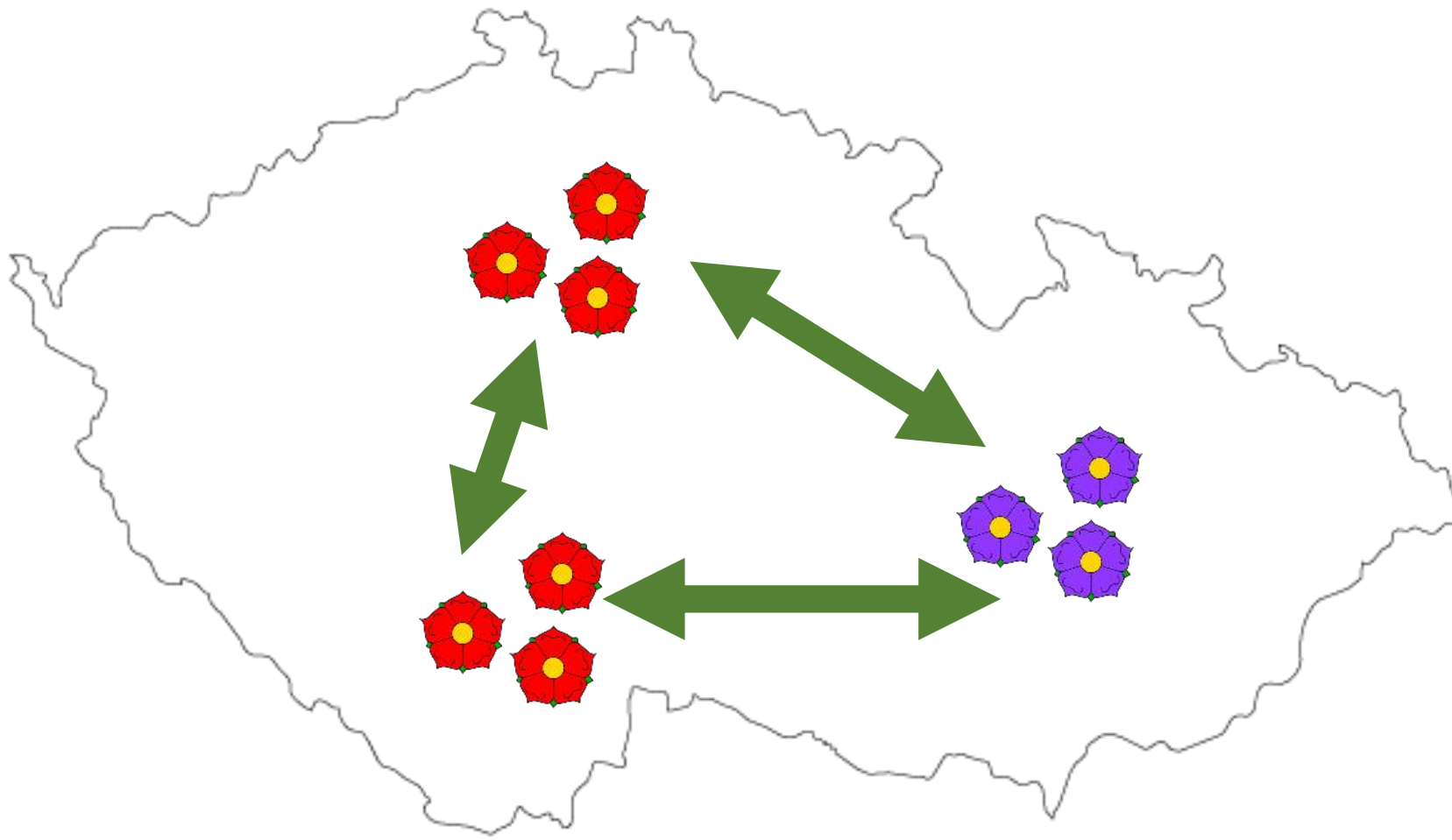
The study of how genomic variation is partitioned within and between populations, and the corresponding evolutionary causes and consequences



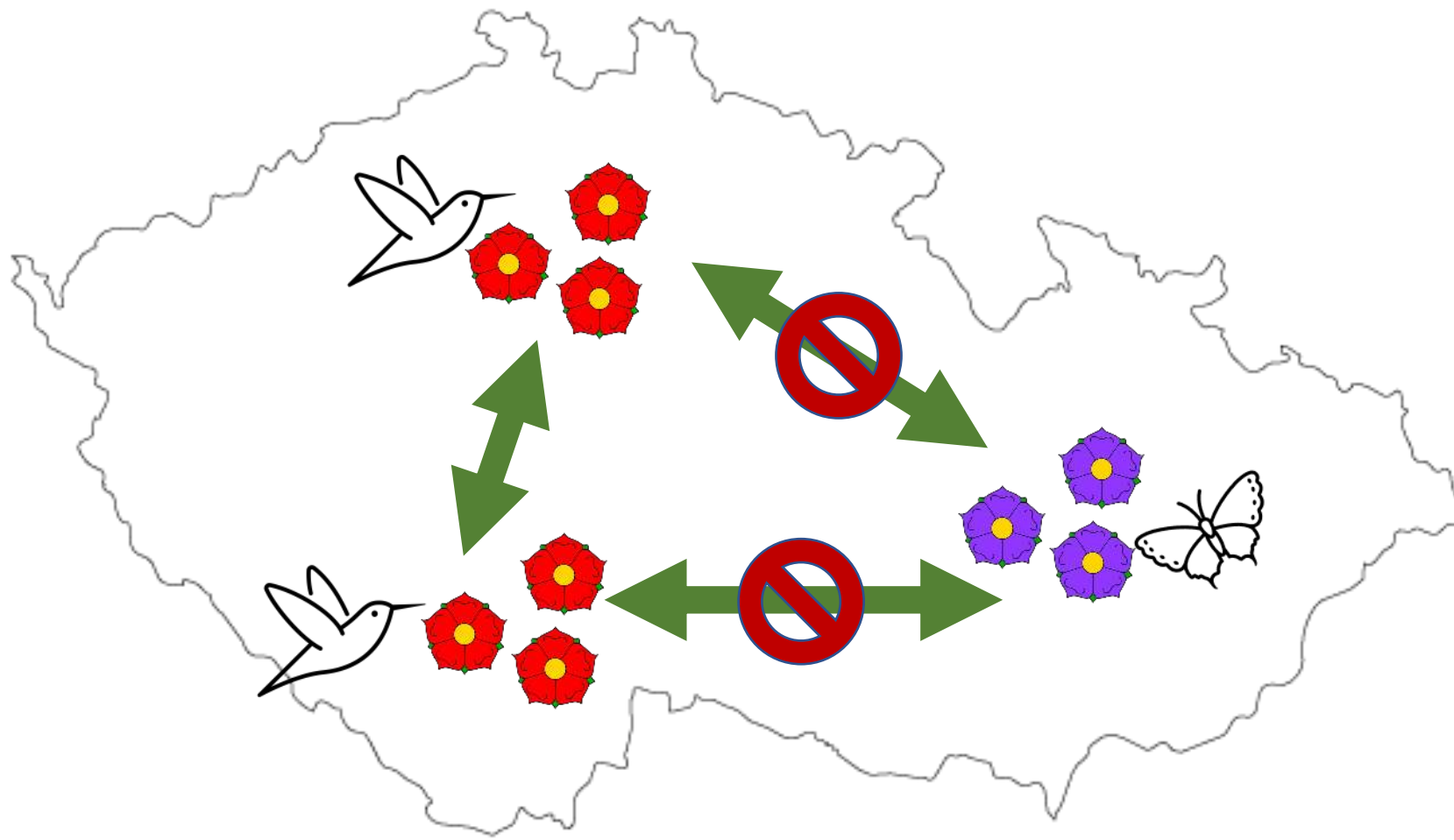
What is the genetic diversity of my species?



How much gene flow is there between populations?

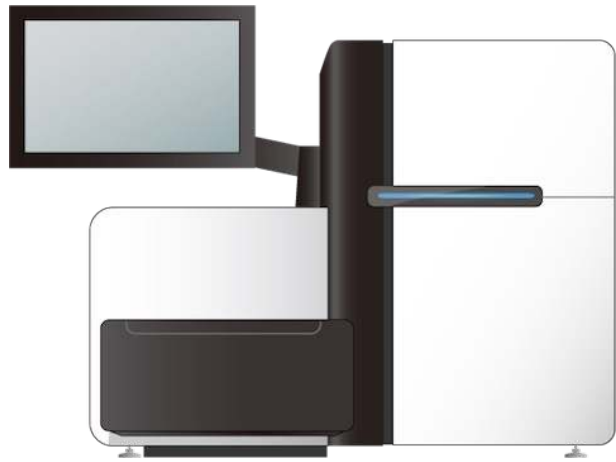


Are differences between populations due to selection?



Do trait differences restrict gene flow?

How can we address these questions?



Sequence some genomes!

The genome contains the signature of a population's evolutionary history – we just need to decipher it

Evolutionary processes that shape diversity

Mutation

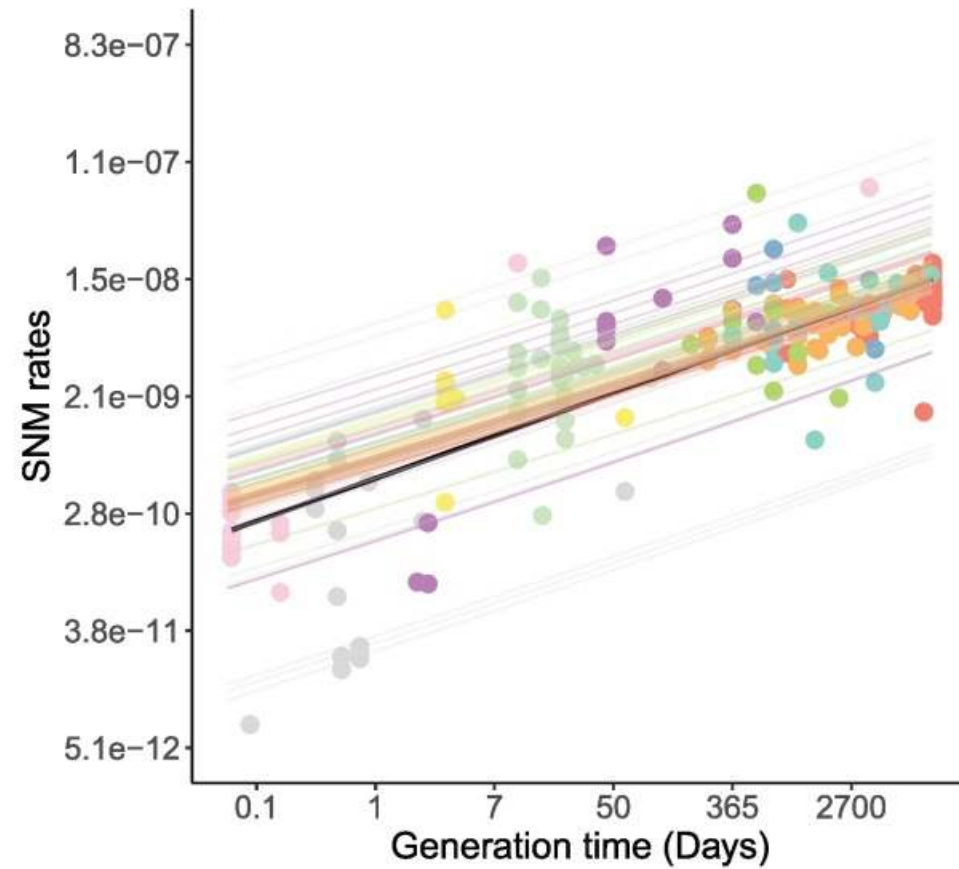
Genetic drift

Recombination

Natural selection

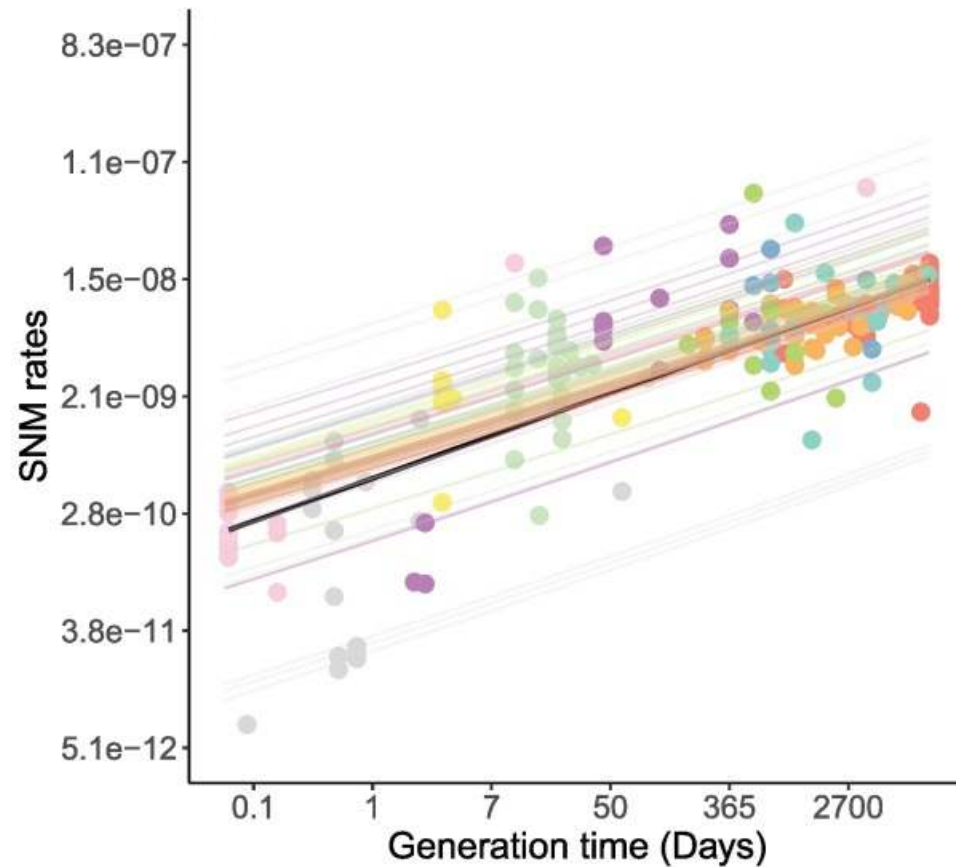
Migration

Mutation



Mutation is the original source of genetic variation

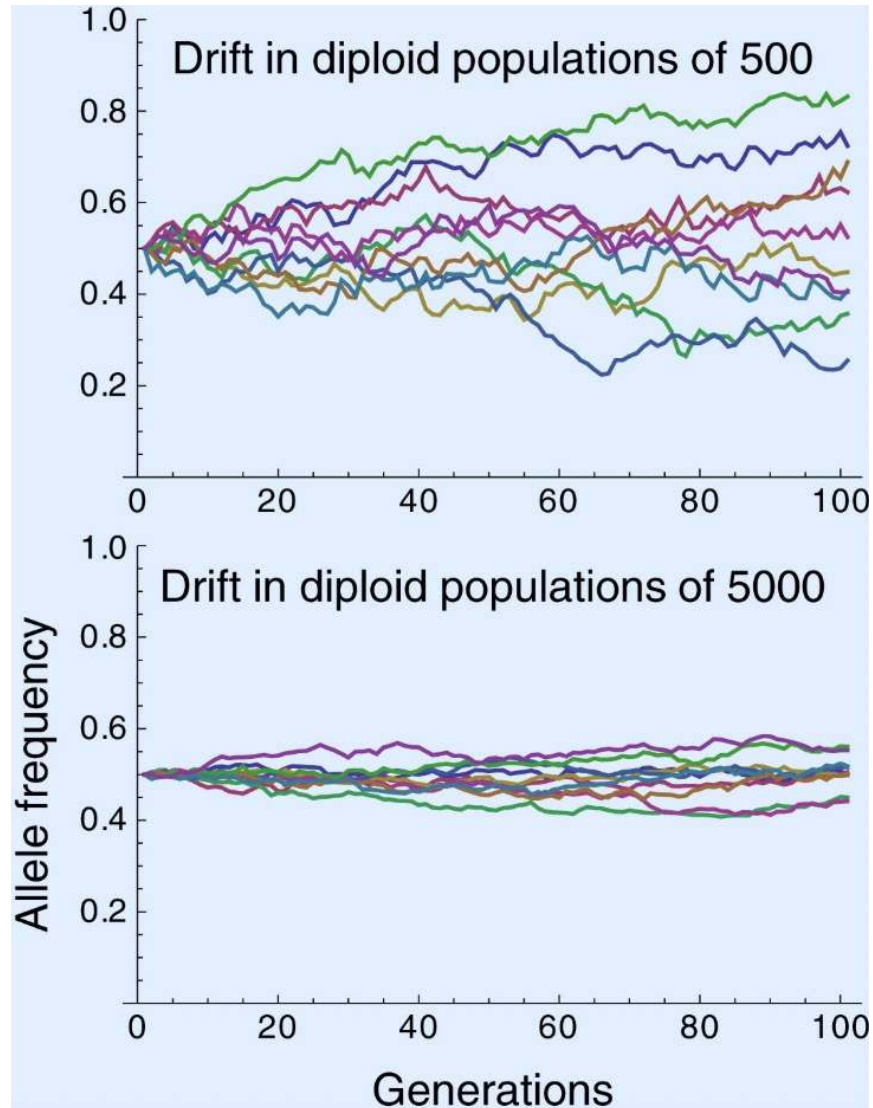
Mutation



Mutation is the original source of genetic variation

Without imperfection in DNA repair machinery we wouldn't be here – variation is the substrate of evolution!

Genetic drift

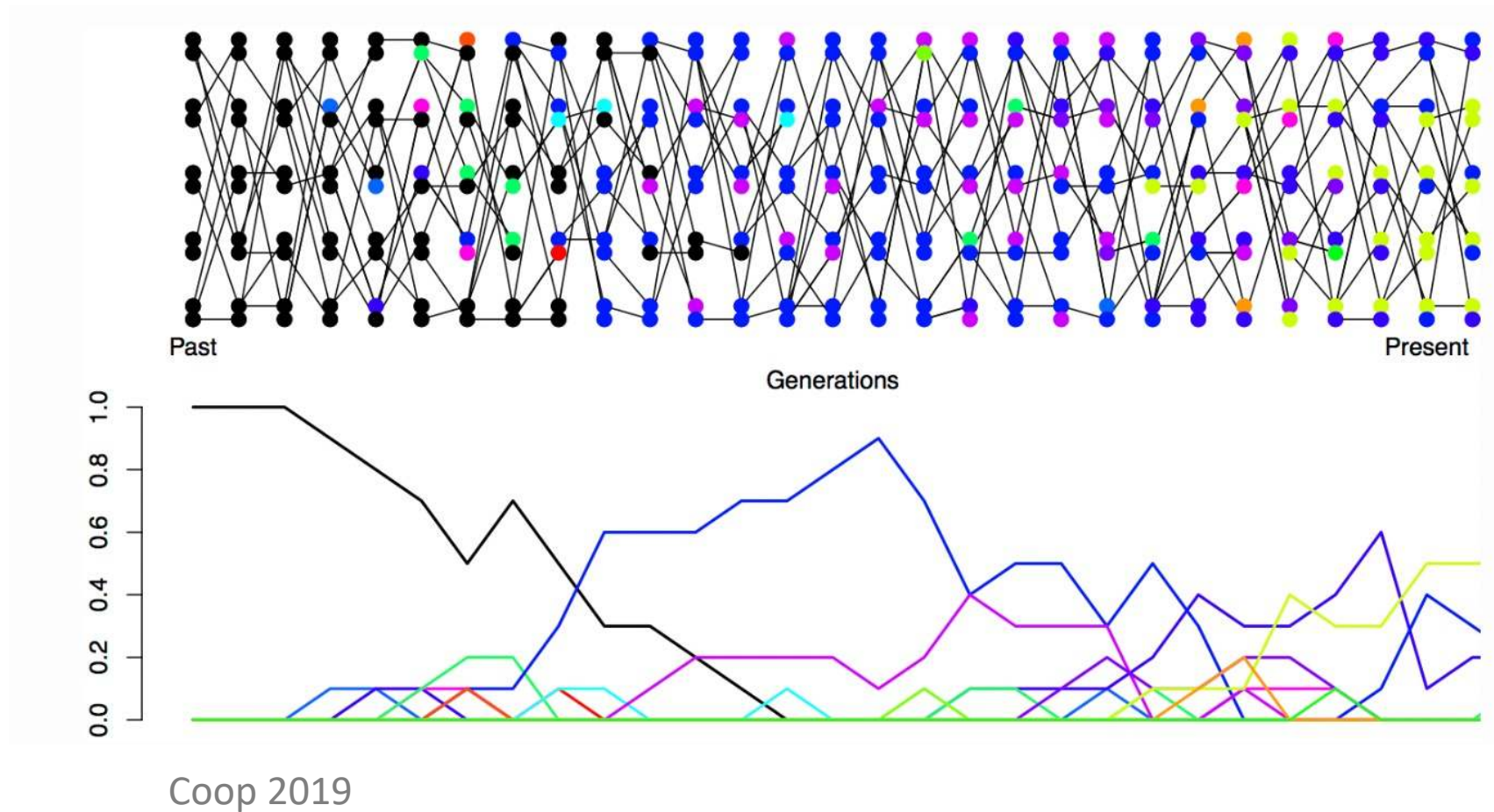


Drift leads to random fluctuations in allele frequencies

Ultimate fate of alleles is loss/fixation due to drift

Probability of fixation for a neutral mutation = $1/2N$

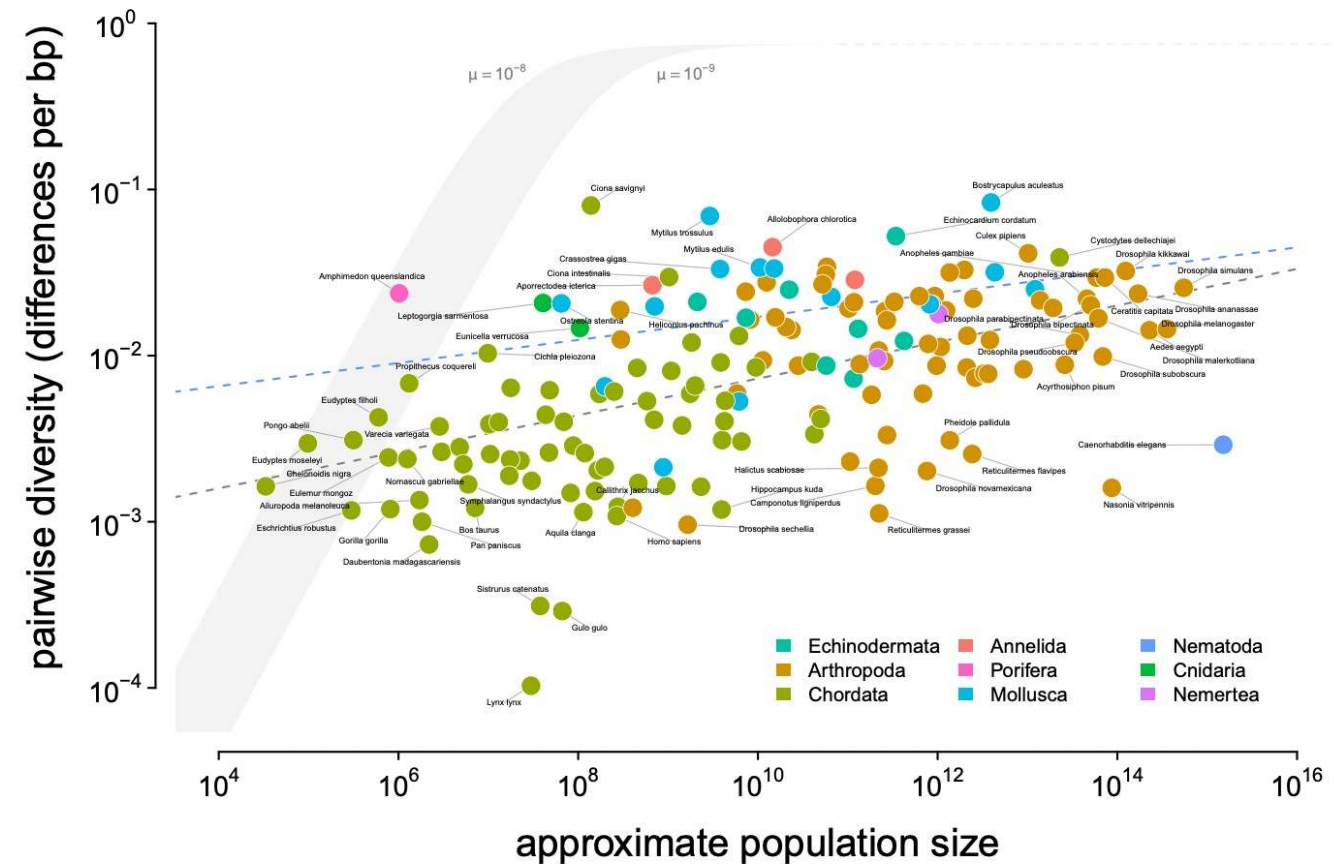
Neutral diversity is a balance between drift and mutation



Population scaled
mutation rate =
 $4N\mu$

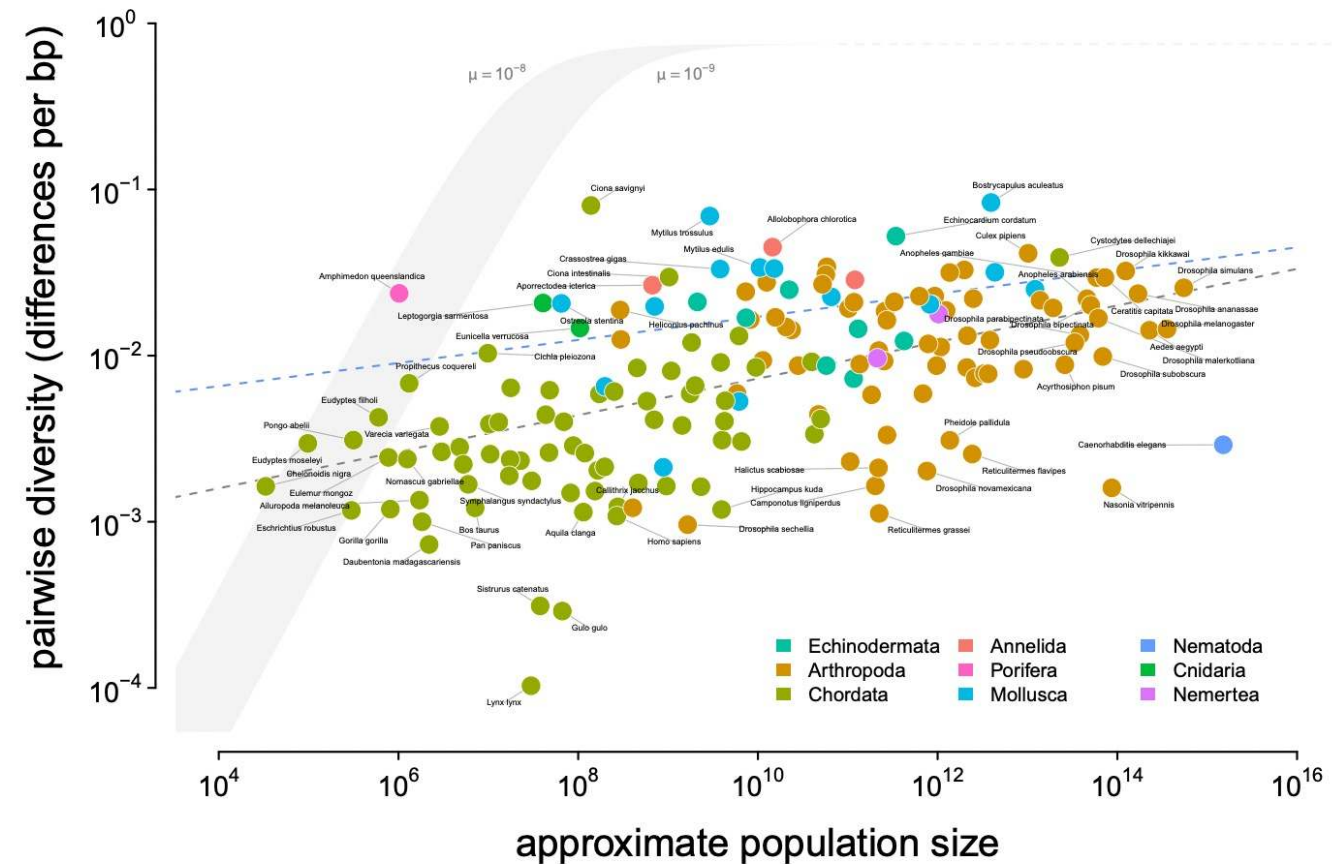
Effective population size: N_e

- Variation in census size is much greater than variation in genetic diversity

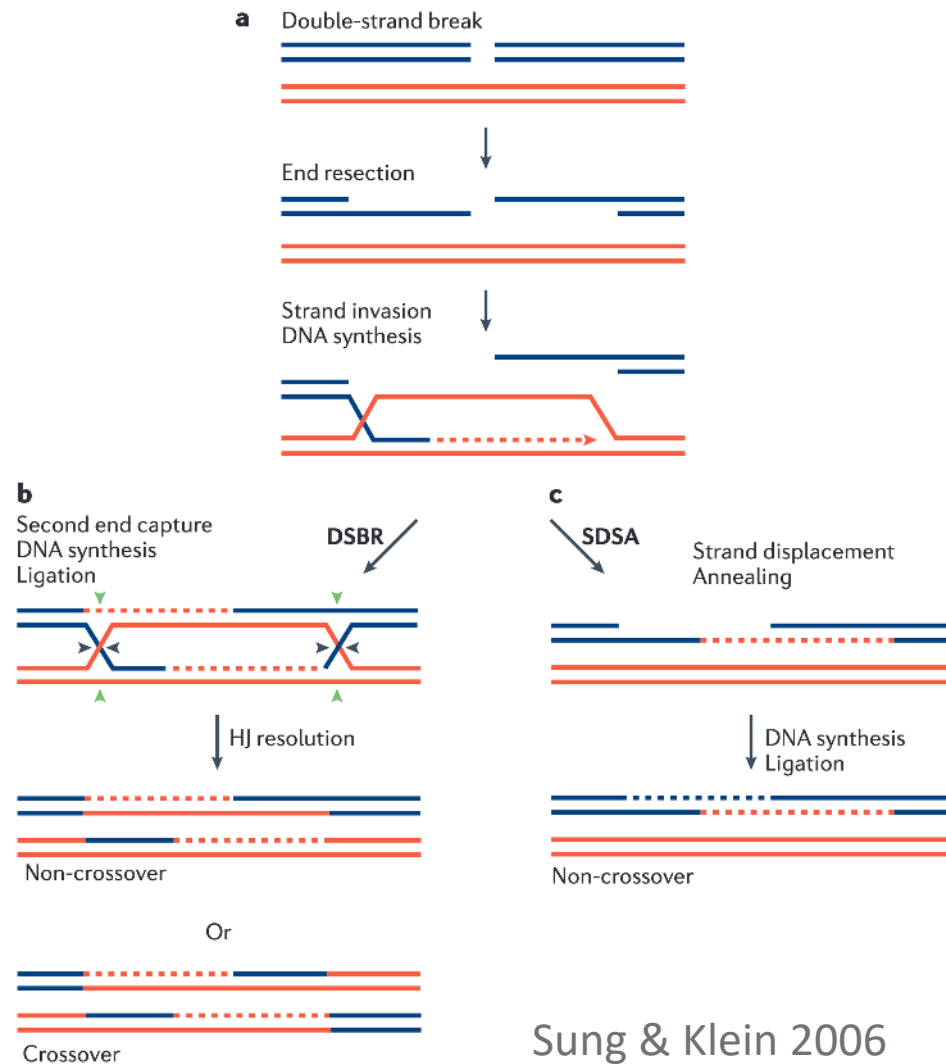


Effective population size: N_e

- Variation in census size is much greater than variation in genetic diversity
- Effective population size could explain this some of this disconnect
- N_e impacts the relative strengths of drift and selection

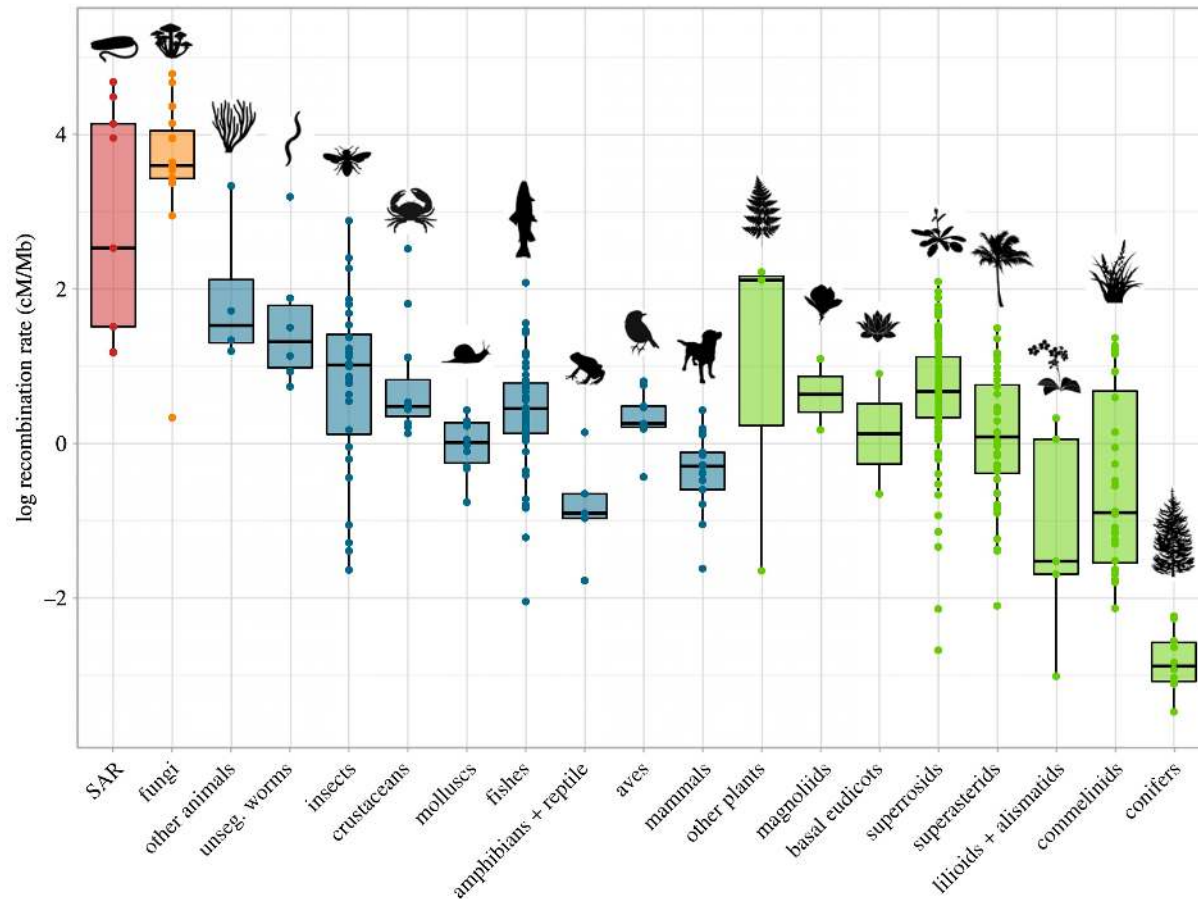


Recombination



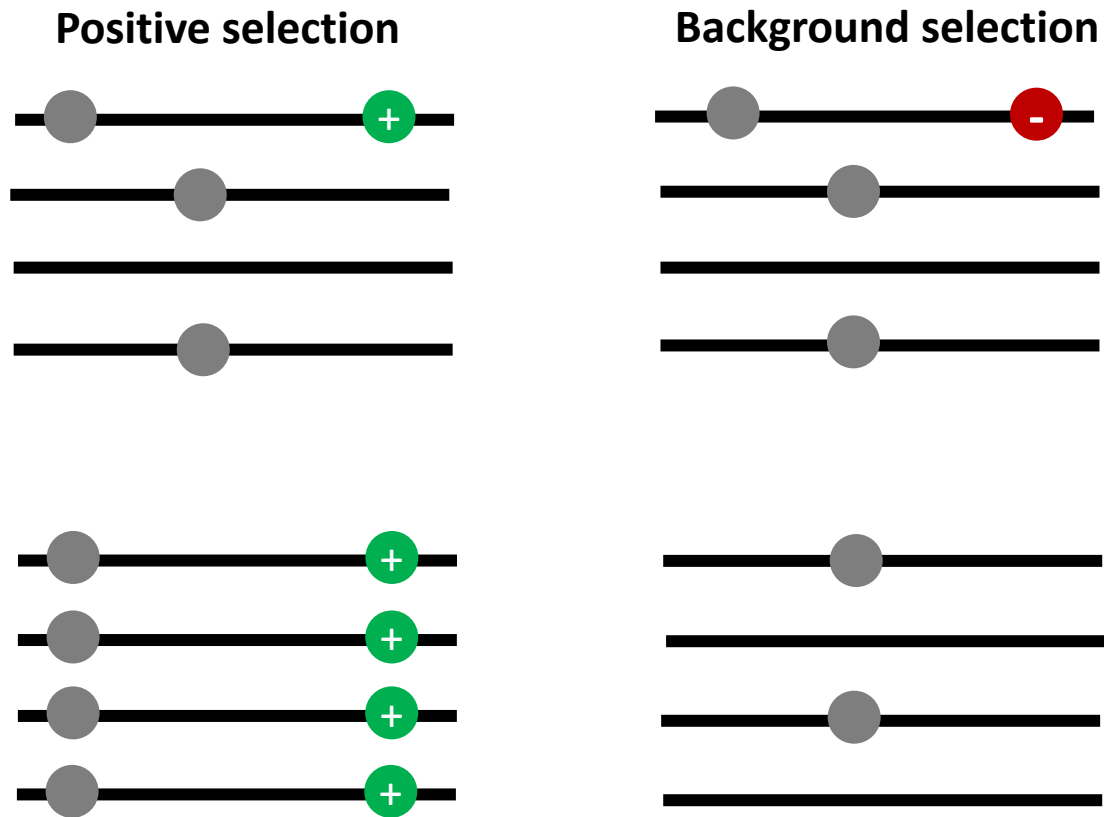
- Repair of DSBs can lead to crossover events → creates new combinations of alleles

Recombination



- Repair of DSBs can lead to crossover events → creates new combinations of alleles
- Recombination rate varies across the genome and among species

Natural selection



- Positive and background selection reduce variation in a population – both at target of selection and at linked sites
- Balancing selection maintains variation
- Methods for detecting selection often depend on this linked signature!

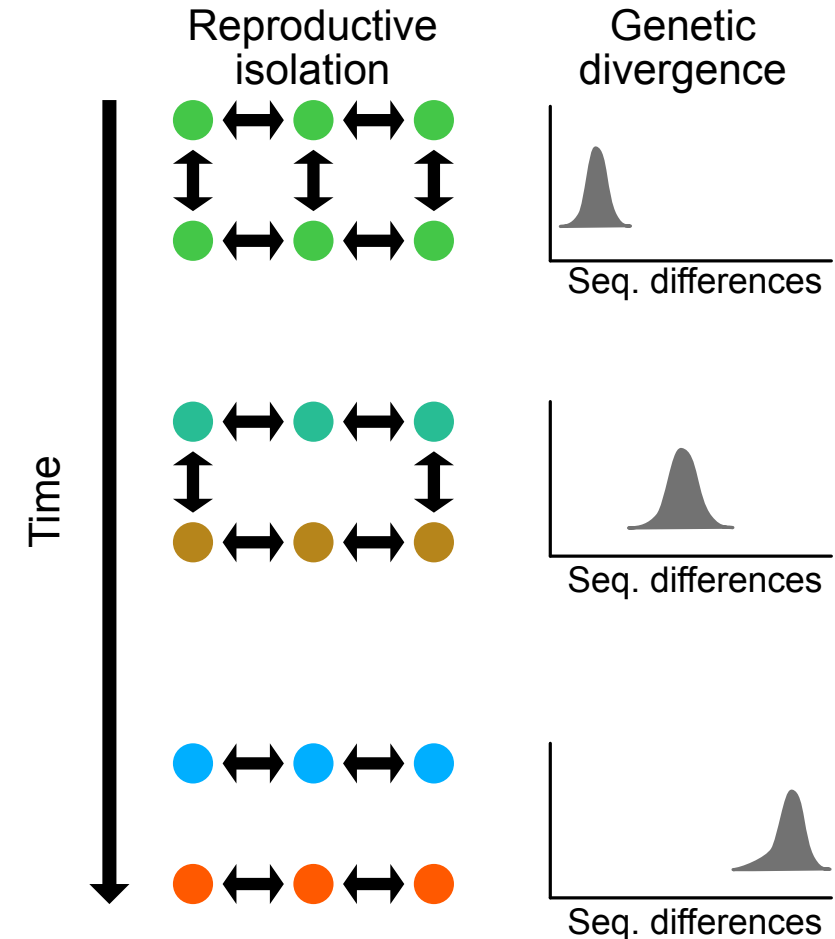
Lower RR creates **stronger** signature of selection

Gene flow

- Gene flow homogenizes populations

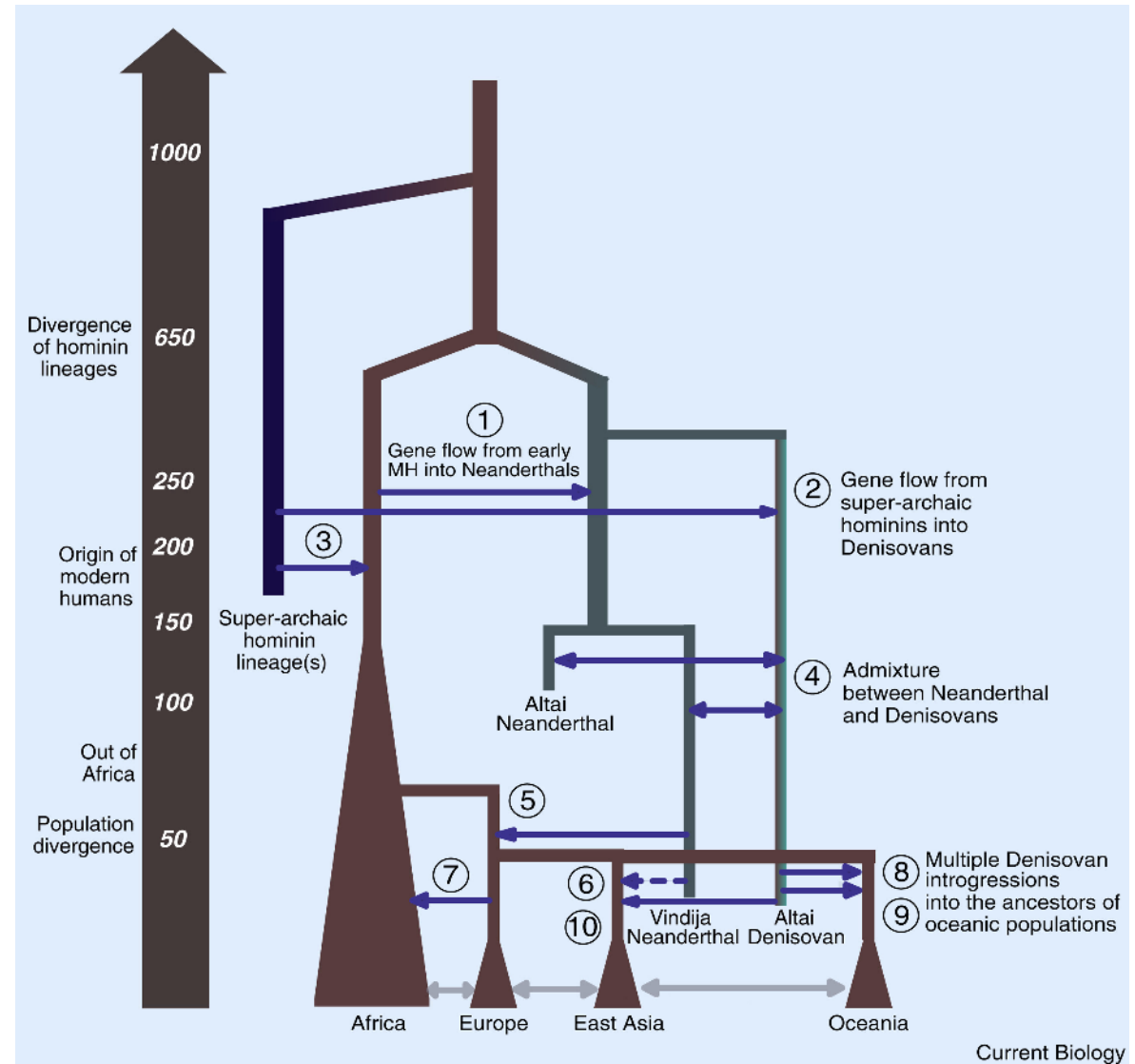
Gene flow

- Gene flow homogenizes populations
- Reduction in gene flow (reproductive isolation) can lead to speciation

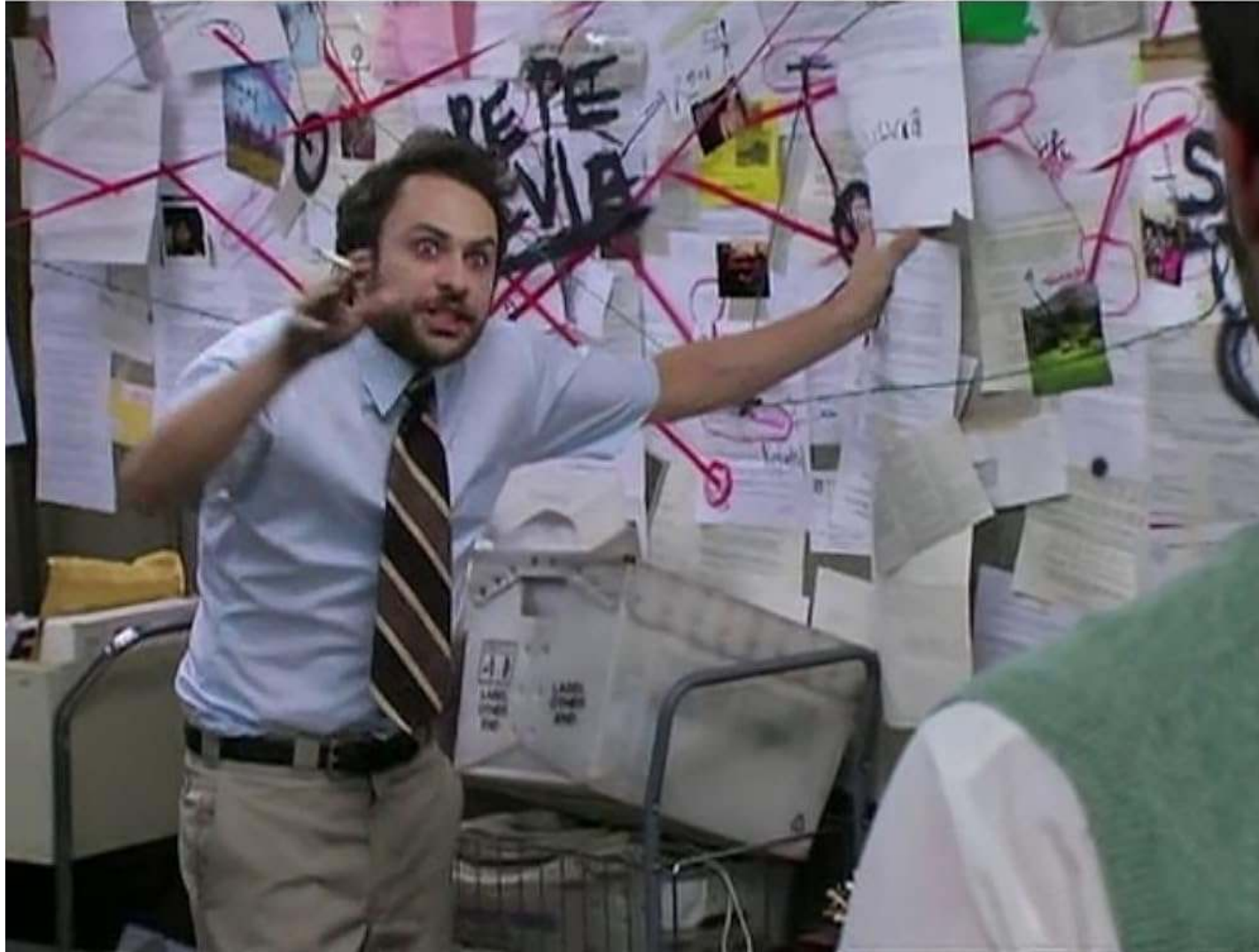


Gene flow

- Gene flow homogenizes populations
- Reduction in gene flow (reproductive isolation) can lead to speciation
- Gene flow can also be an important source of adaptive mutations!



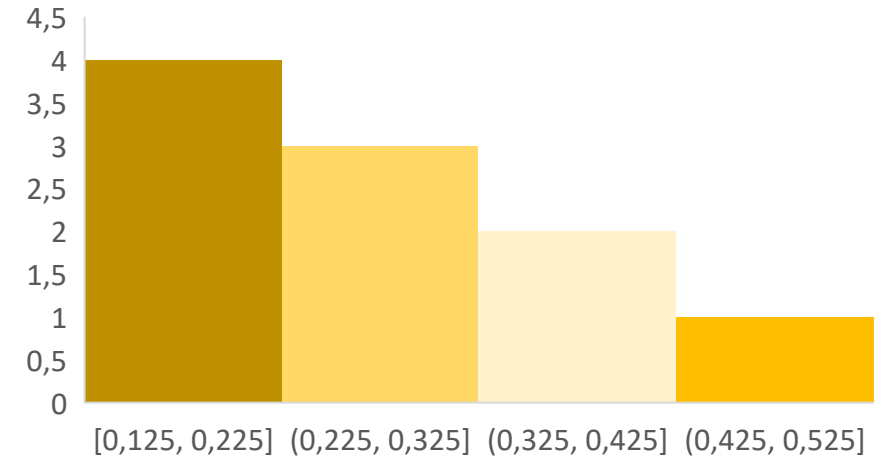
Determinants of genetic diversity



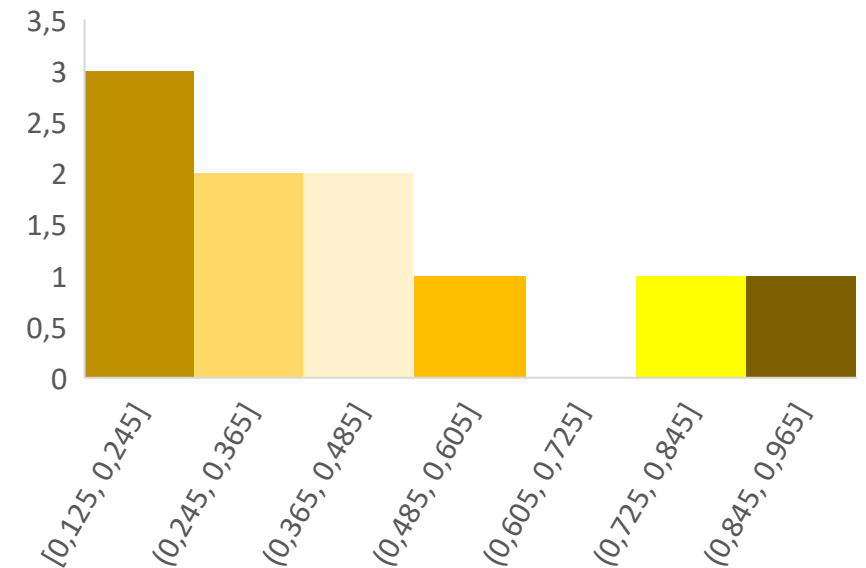
The site frequency spectrum

Outgroup	C	C	T	G	T	C	A	G	T	T
Sample 1	C	C	T	G	T	C	A	G	T	T
Sample 2	C	T	T	G	T	A	A	C	T	A
Sample 3	G	T	T	C	C	C	A	C	T	T
Sample 4	C	T	T	G	T	C	A	C	A	A
Sample 5	G	T	T	C	C	A	G	C	T	T
Sample 6	C	T	T	G	C	A	A	C	A	T
Sample 7	C	T	A	G	T	C	A	G	A	T
Sample 8	C	T	T	G	T	A	A	C	T	T

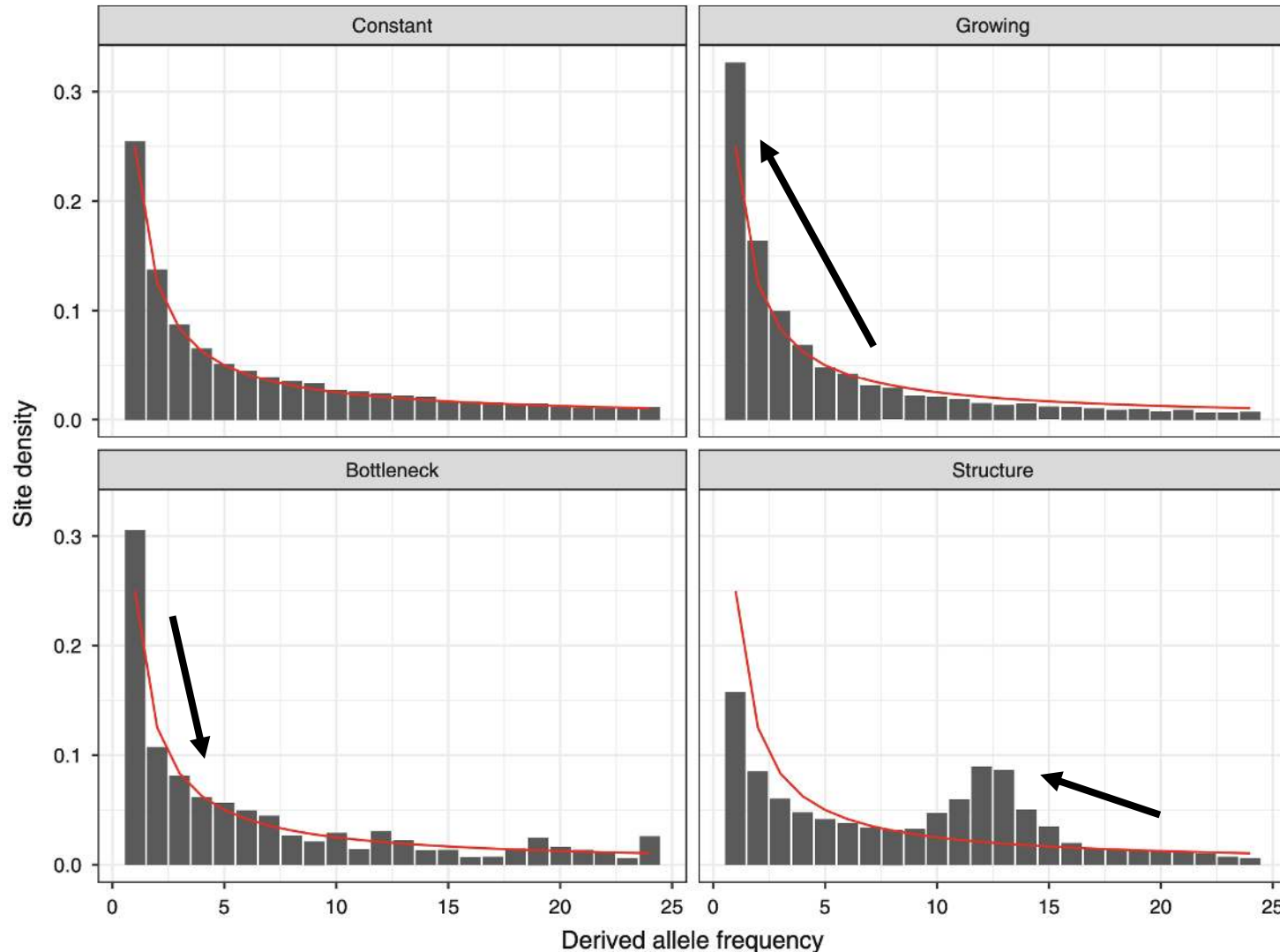
Folded SFS



Unfolded SFS



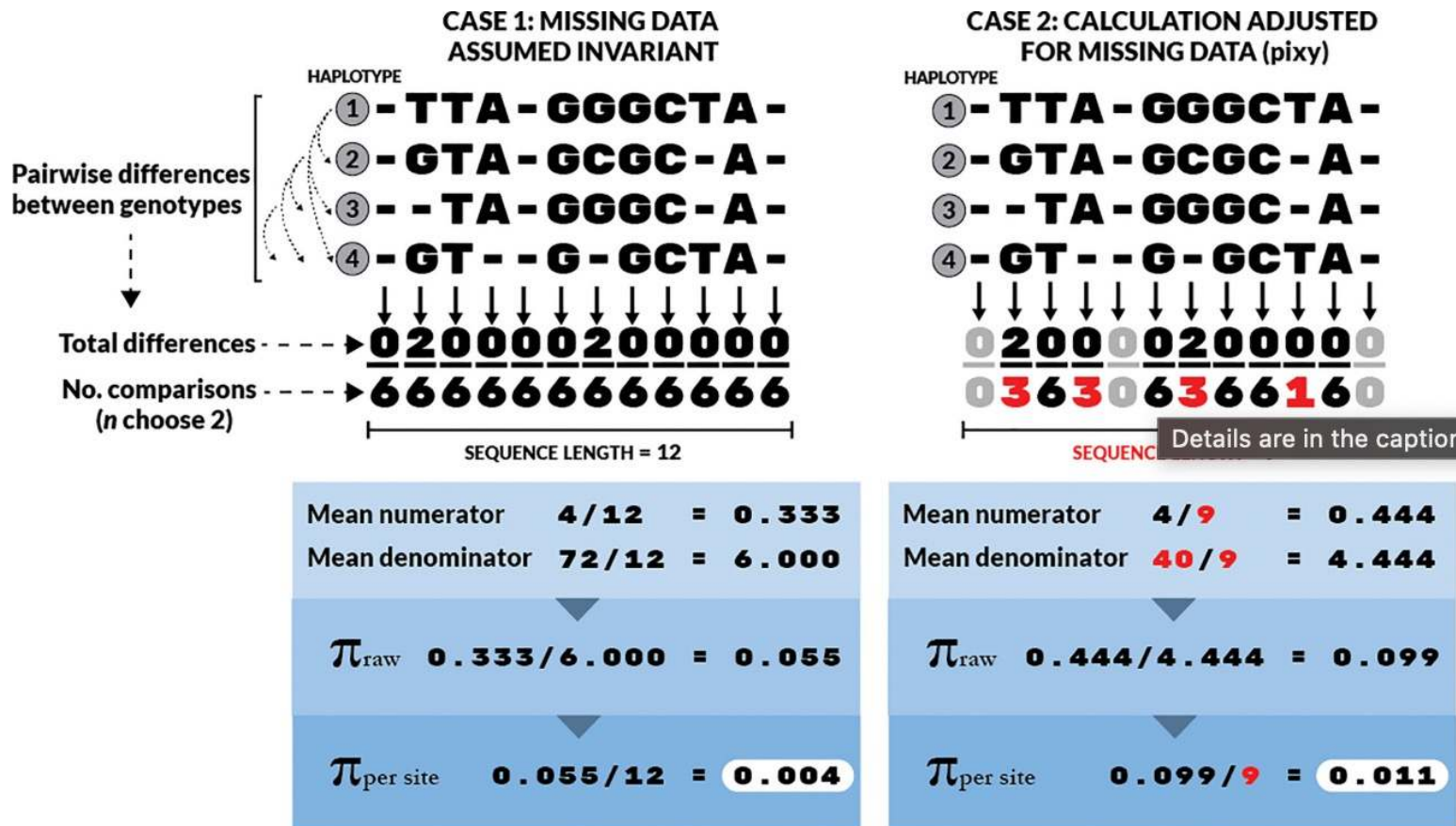
The site frequency spectrum



Neutral expectation for the density of sites in a frequency class = $1/f$

The genome-wide SFS is impacted by the demographic history of a population

Estimating genetic diversity

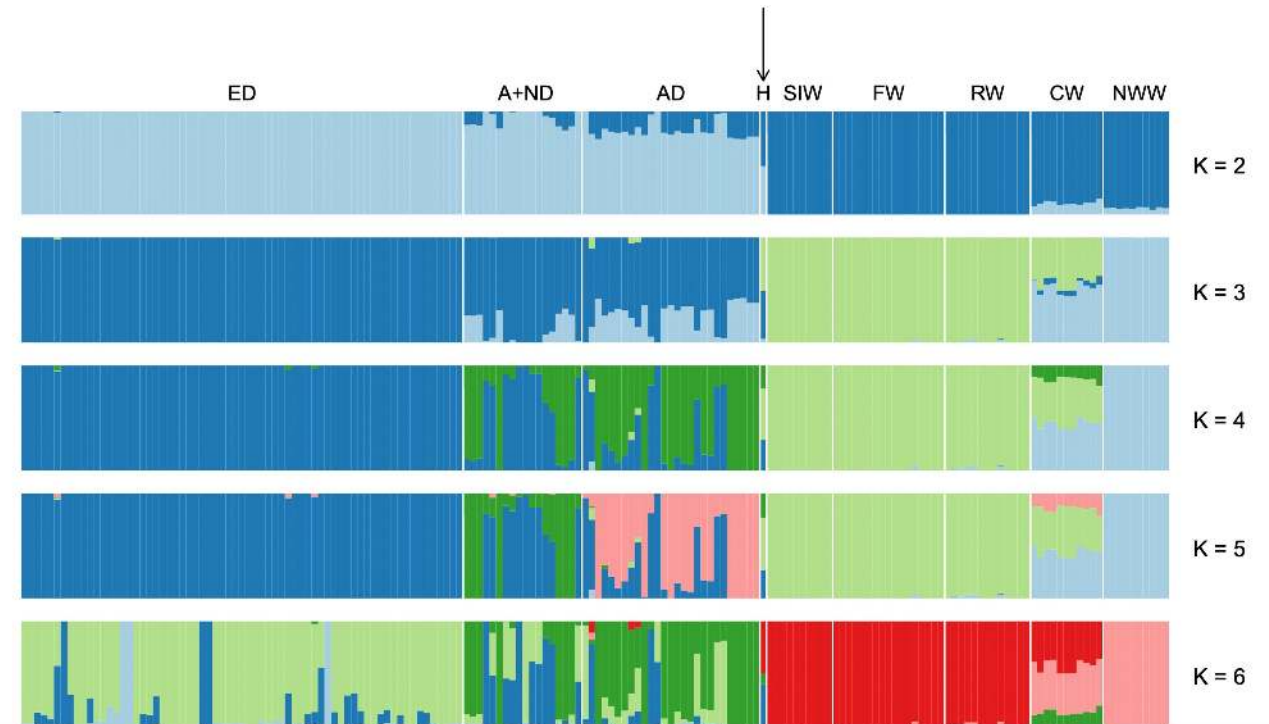
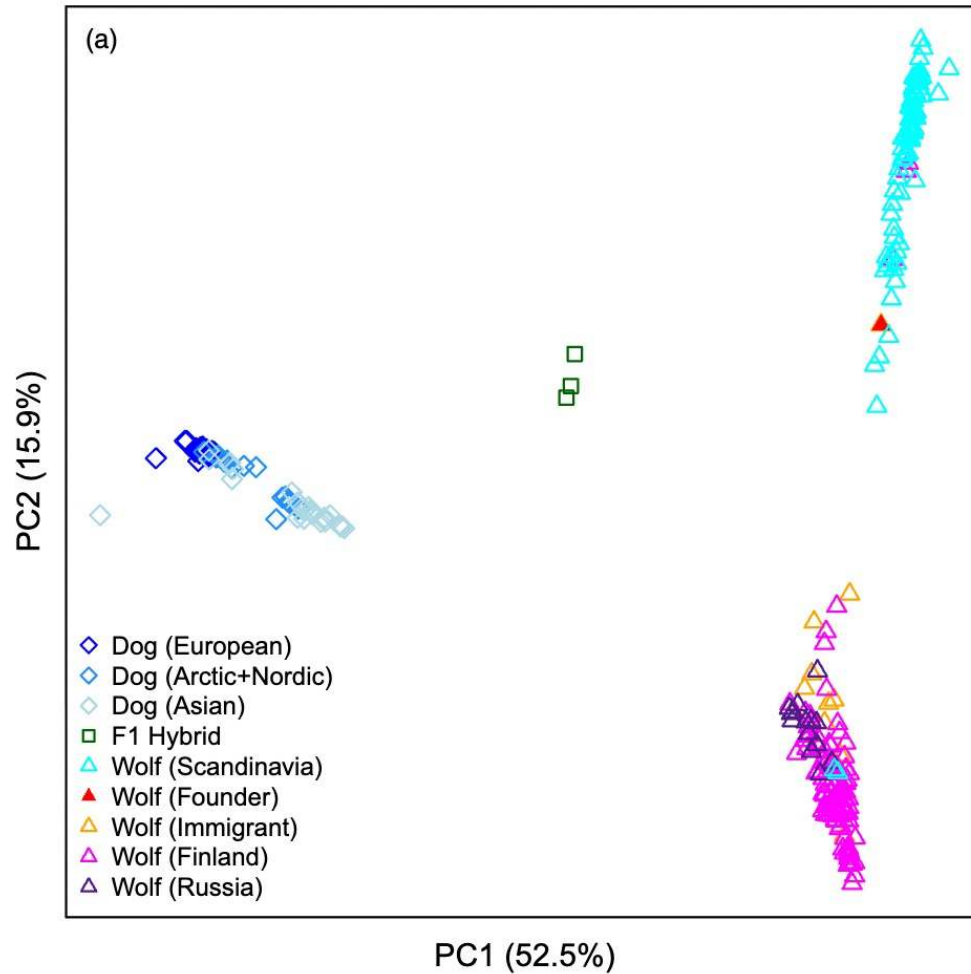


Pairwise nucleotide diversity: π

Mean number of pairwise differences in a population

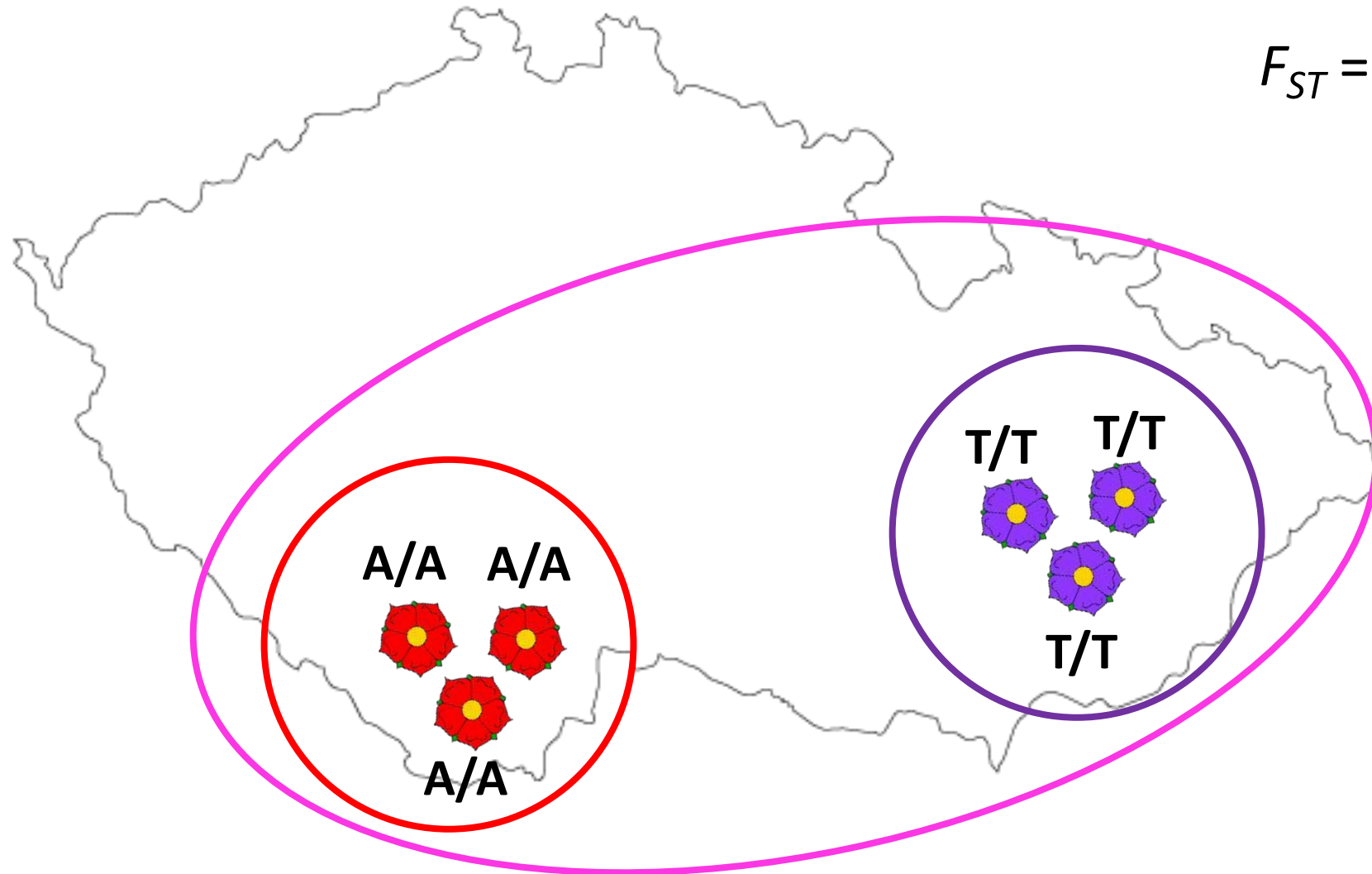
$$\pi = 4N_e \mu$$

Population structure



Measures of genetic differentiation

$$F_{ST} = \frac{\pi_{BETWEEN} - \pi_{WITHIN}}{\pi_{BETWEEN}}$$

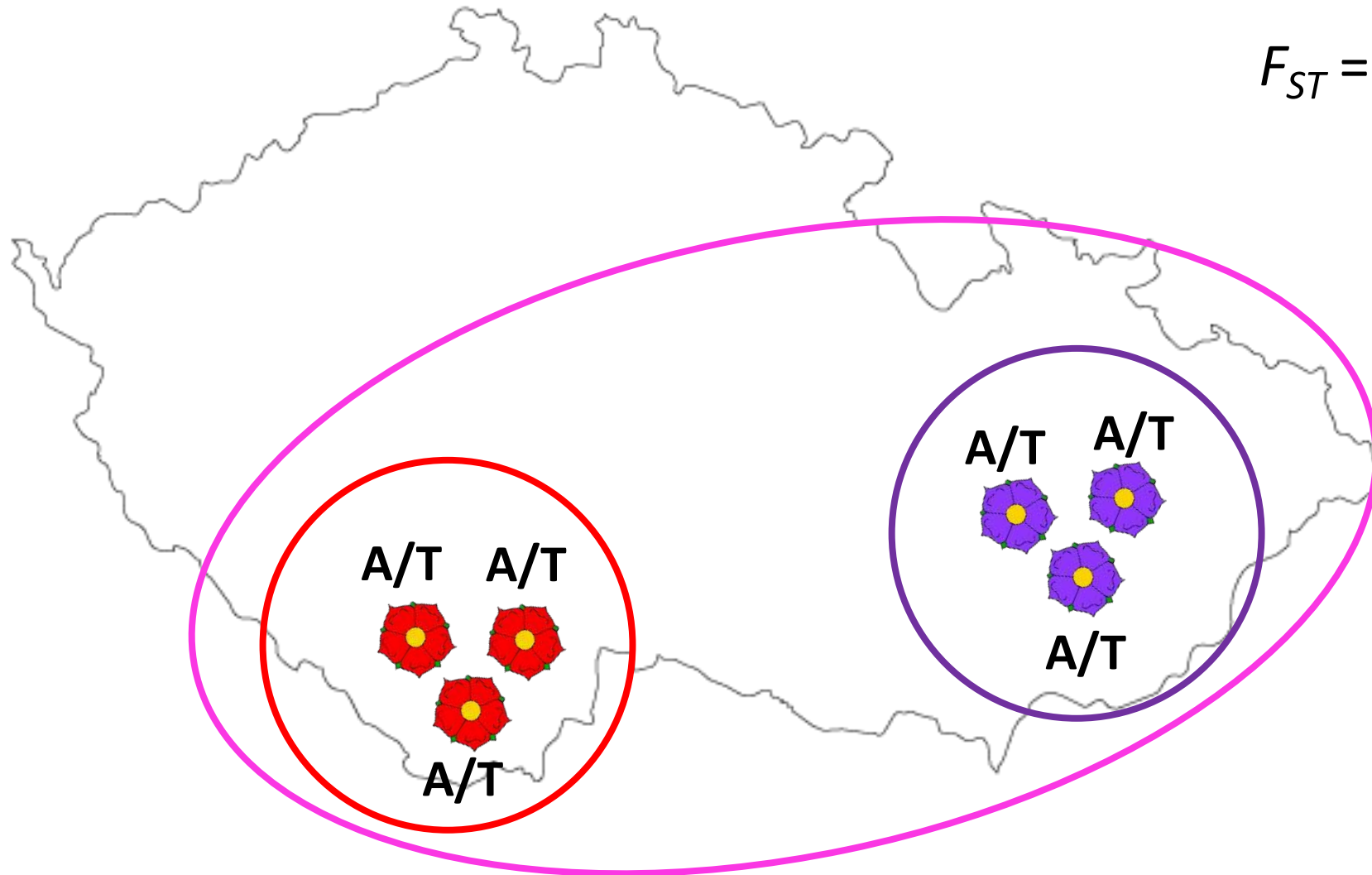


$$F_{ST} = \frac{0.5 - 0}{0.5}$$

$$F_{ST} = 1$$

Measures of genetic differentiation

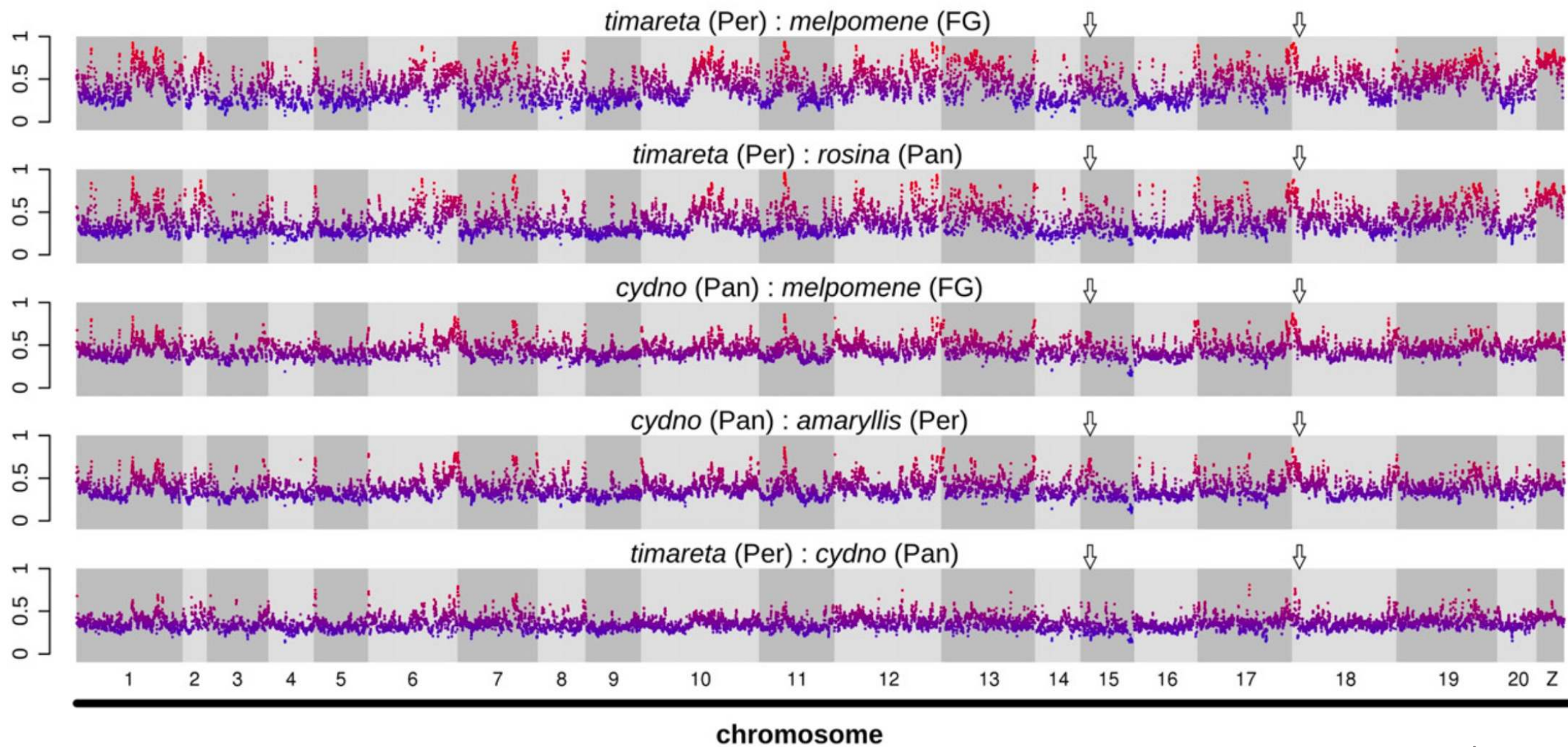
$$F_{ST} = \frac{\pi_{BETWEEN} - \pi_{WITHIN}}{\pi_{BETWEEN}}$$



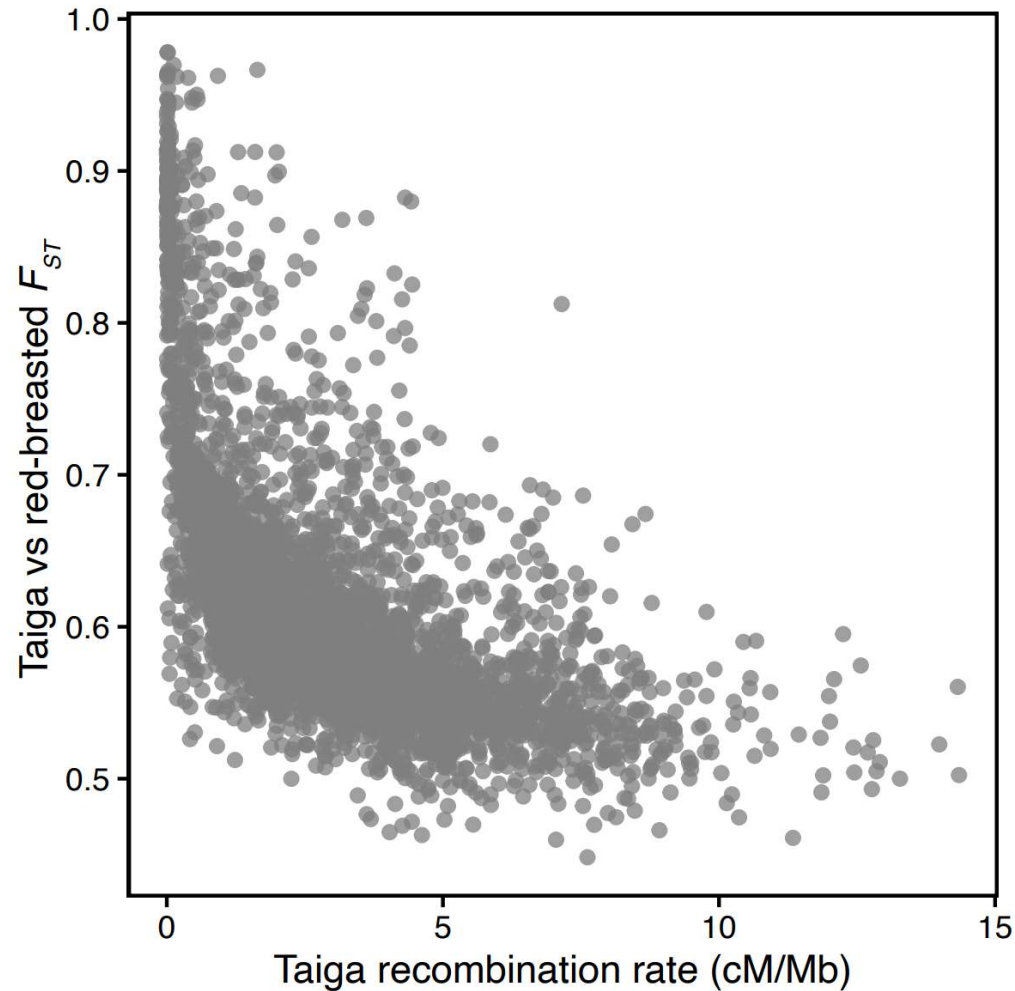
$$F_{ST} = \frac{0.5 - 0.5}{0.5}$$

$$F_{ST} = 0$$

Differentiation varies across the genome

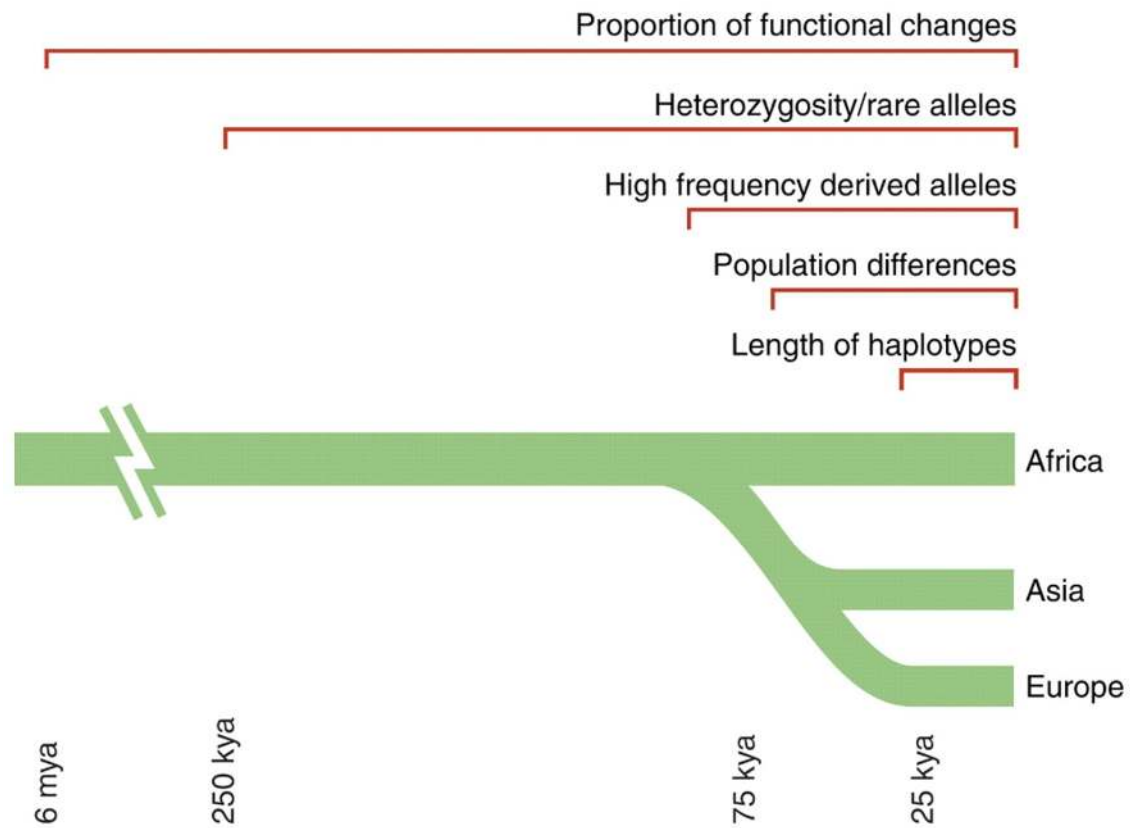


Differentiation varies across the genome

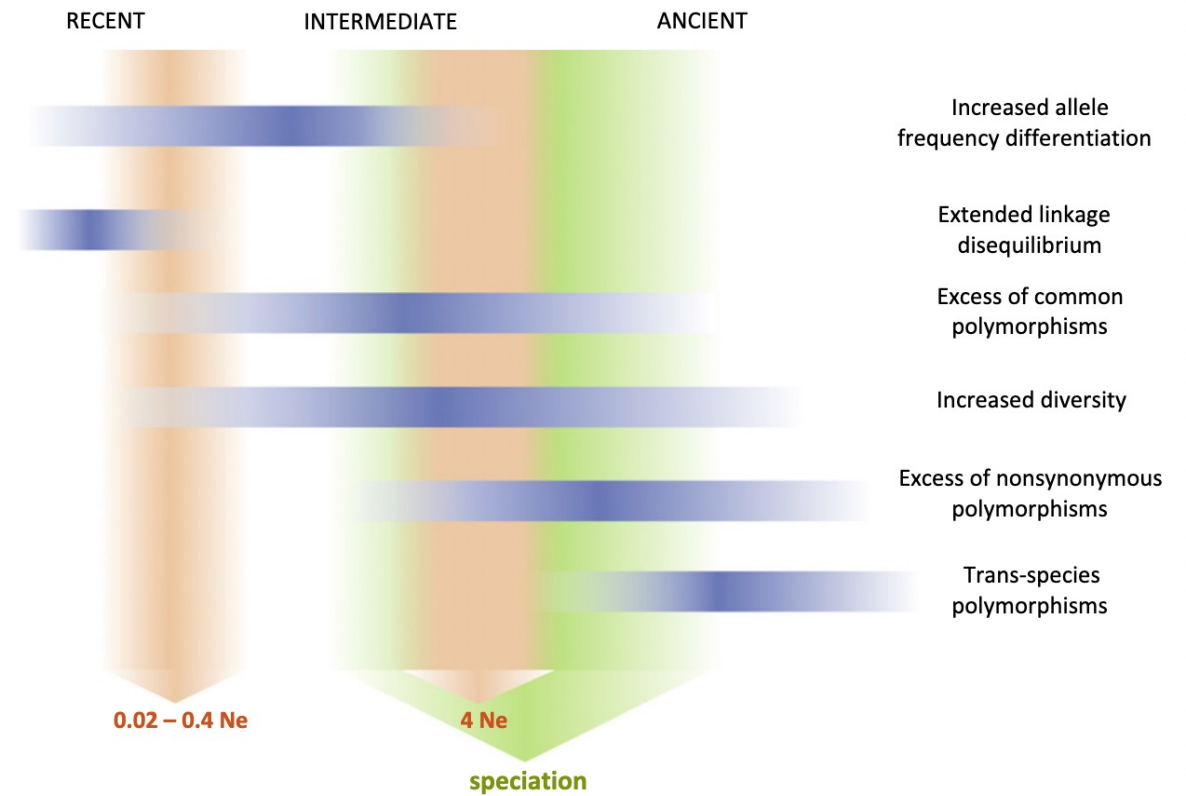


Differentiation is strongly correlated with recombination rate across a wide range of organisms

Detecting natural selection



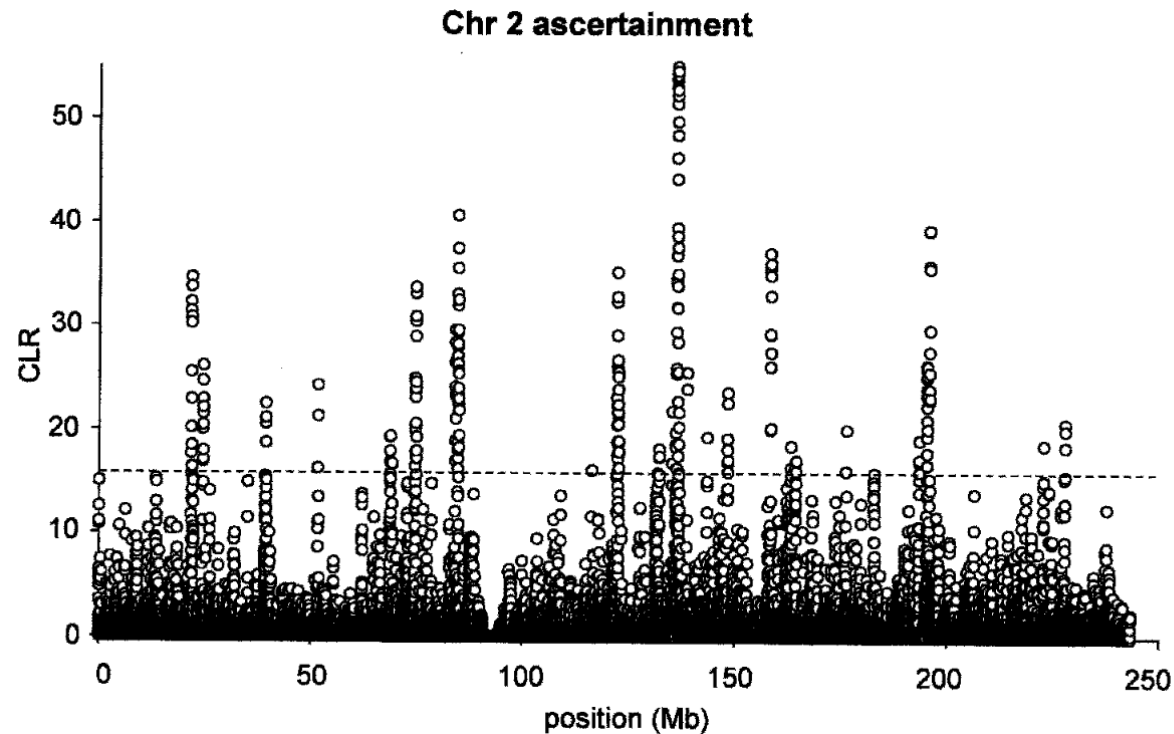
Sabeti et al. 2006, Science



Fijarczyk & Babik 2015, Mol. Ecol.

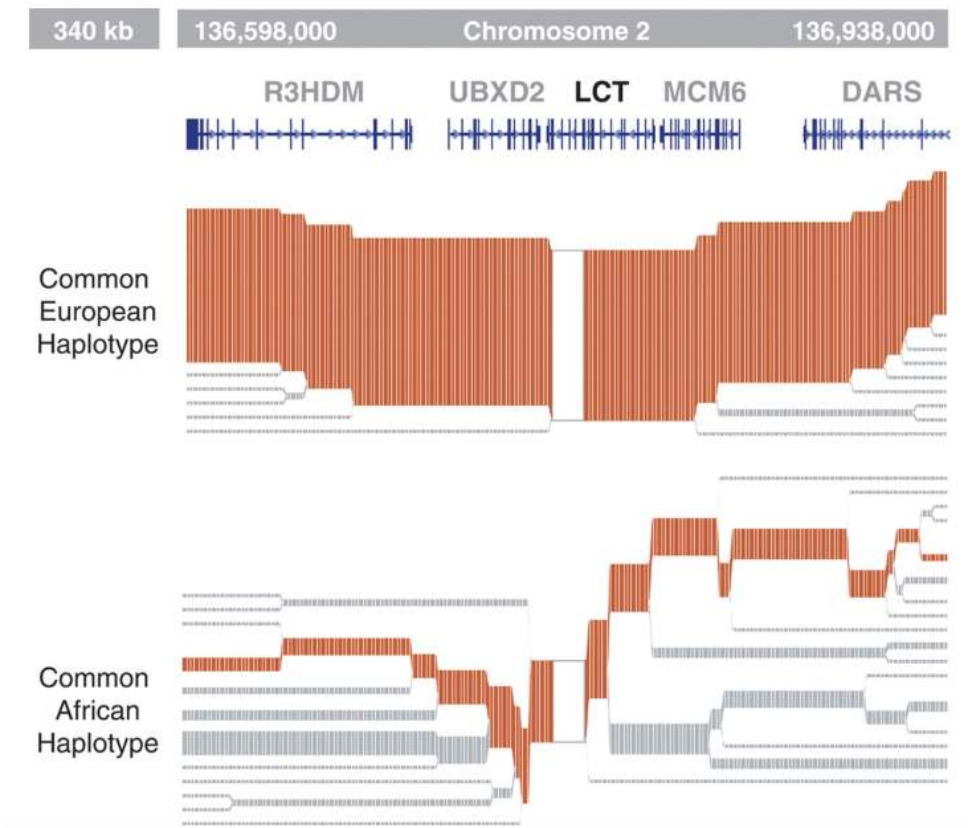
Detecting natural selection

SFS-based methods



Nielsen et al. 2005, Genome Research

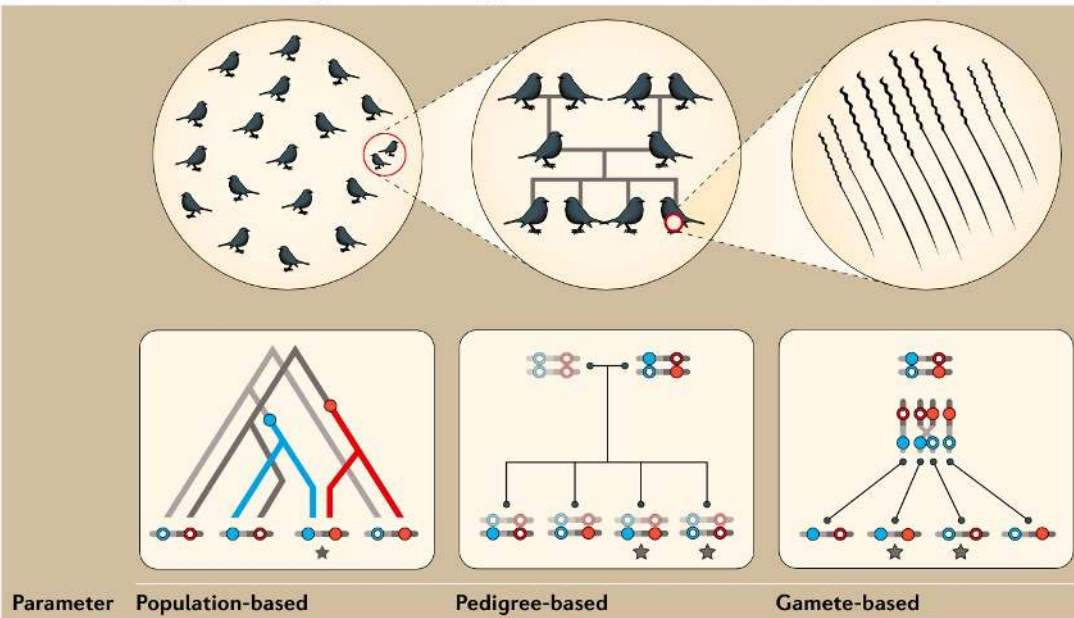
Haplotype-based methods



Sabeti et al. 2006, Science

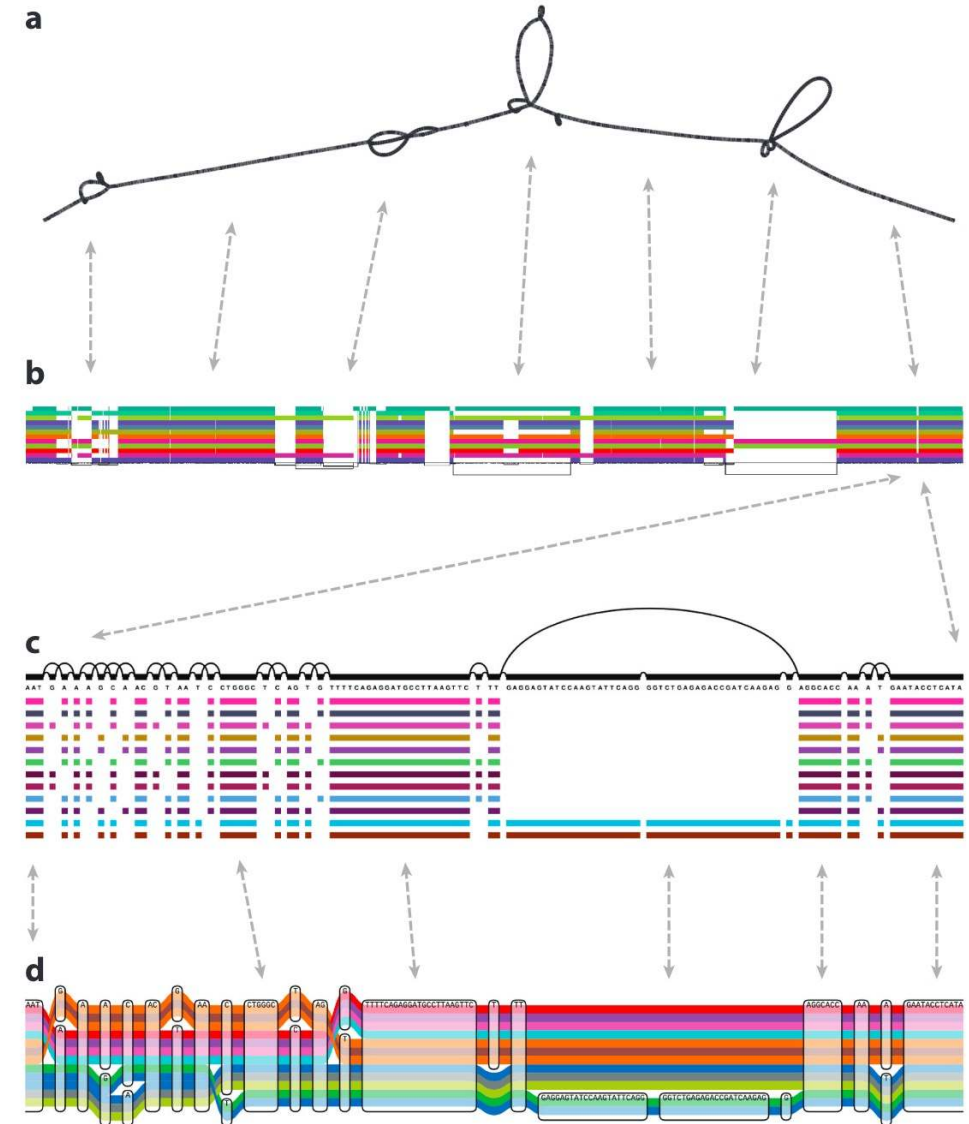
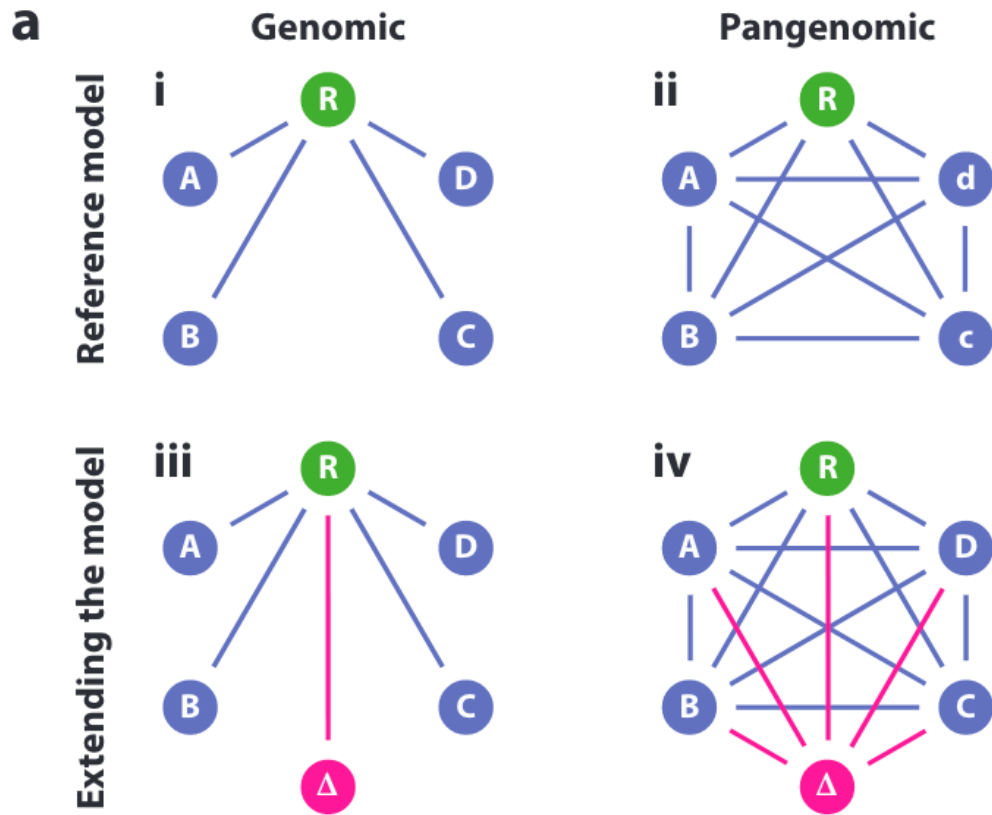
Estimating recombination rates

Table 1 | Summary of the three genomic-based approaches to infer the recombination landscape

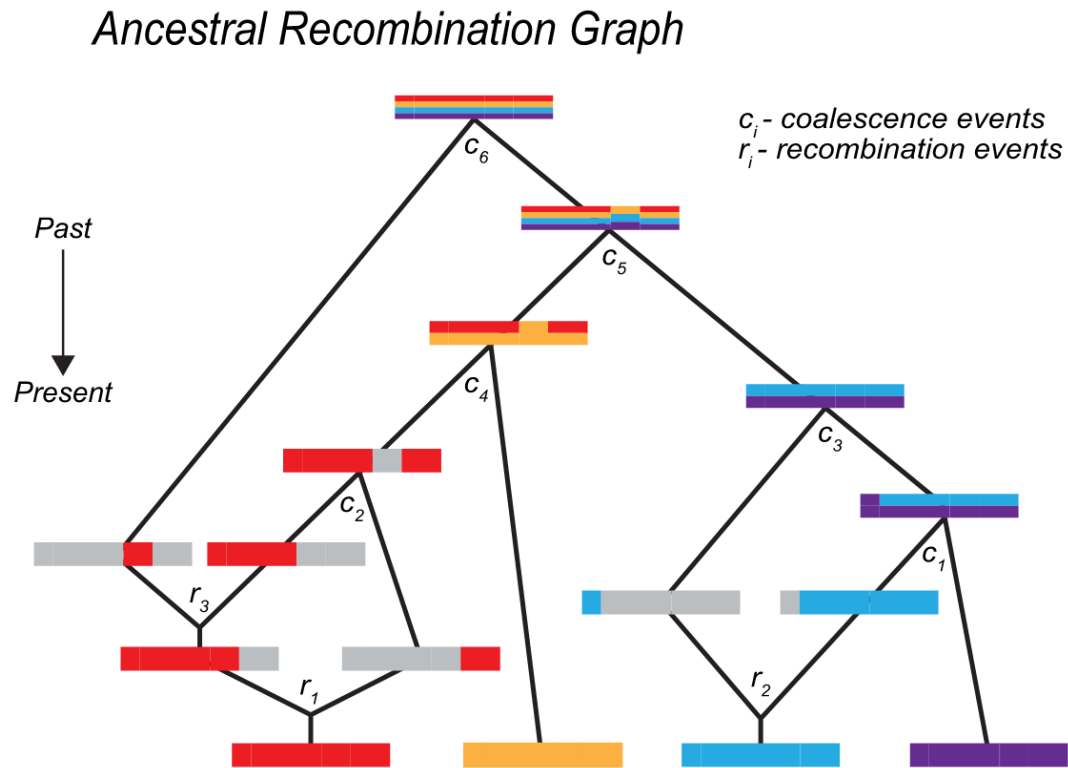


Parameter	Population-based	Pedigree-based	Gamete-based
Estimate	Population recombination rate	Genetic distance	Crossover frequency
Sample size	10–30	>100	≥1
Pros	Low to moderate sampling Moderate to high resolution Simple study design	Sex-specific recombination maps Unbiased by population level processes Variation between individuals and heritability can be measured Can improve genome assemblies	High resolution Low sampling required Variation between individuals can be compared
Cons	Average between sexes and through time Biased by effective population size, demographic history, selection and mutation Dependent on genome assembly quality	Large sampling required Complex and time-intensive study design required Low resolution (~1 Mb, 0.5–2 cM) Biased by SNP density in mapping population Only contemporary snapshot of recombination	Single-sex recombination map (usually males) Biased by individual SNP density Dependent on genome assembly quality Only contemporary snapshot of recombination

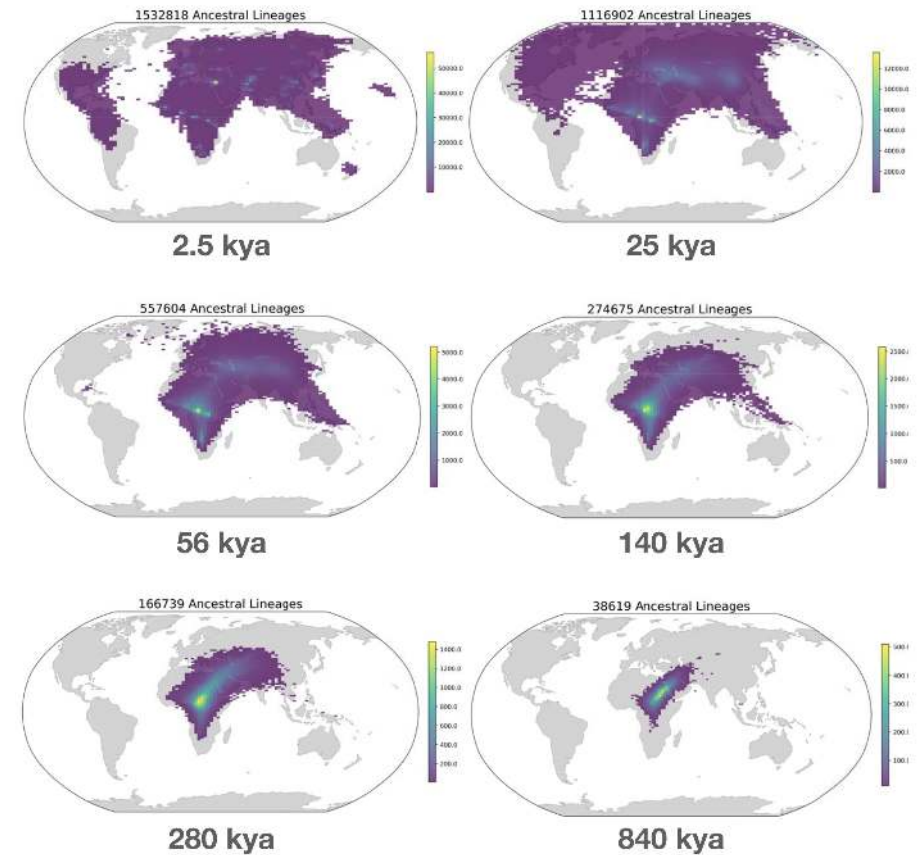
Advances in population genomics: Pan-genomics



Advances in population genomics: Ancestral recombination graphs



Shipilina et al. 2022



Wohns et al. 2022

Population genetics and scientific racism

POPULATION GENETICS

A geographic history of human genetic ancestry

Michael C. Grundler¹, Jonathan Terhorst², Gideon S. Bradburd^{1*}

Genetic variation in humans is often summarized with discrete labels, but these can be inaccurate and misleading (5, 6). Even when based on geographic history, genetic ancestry labels oversimplify a complex picture because they implicitly focus on only a single point in time. For example, based on our current understanding of human origins, all living individuals are “African” (regardless of the geography of their recent ancestors) when considering their ancestry ~200,000 years before present. Advances in the study of ancient DNA have revealed a lack of genetic continuity within geographic regions (7–10), further highlighting the shortcomings of genetic ancestry labels.

The fact that these labels are generated using statistical genetics approaches gives them the veneer of authenticity, further reifying problematic notions of race and ancestry in society (6, 11).

Reviewing Rushton's magnum opus book, Barash (1995) averred that combining “numerous little turds of variously tainted data” does not yield results, just a “pile of shit.” It is a good line, but to our argument, it speaks to the cognitive style of the human biodiversity movement: accumulate a giant corpus of decontextualized mainstream science, flawed race science, and political writing, ignore or dismiss any critiques, and then treat genetically determined racial differences in behavior as forbidden knowledge—obviously true, but suppressed by the academic left.

Panofsky et al. 2020

Further resources

- Statistical population genomics

doi.org/10.1007/978-1-0716-0199-0

- Population and quantitative genetics

- <https://cooplab.github.io/popgen-notes/>

- Work on scientific racism and genetics:

- <https://doi.org/10.1002/hast.4925>
 - <https://doi.org/10.3389/fgene.2024.1345631>
 - <https://doi.org/10.1002/ajpa.24150>
 - <https://doi.org/10.1038/s41436-021-01109-w>