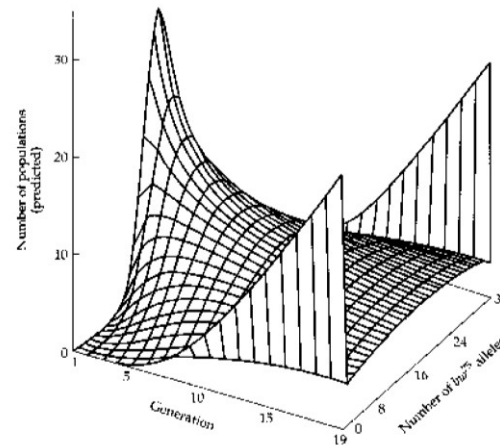
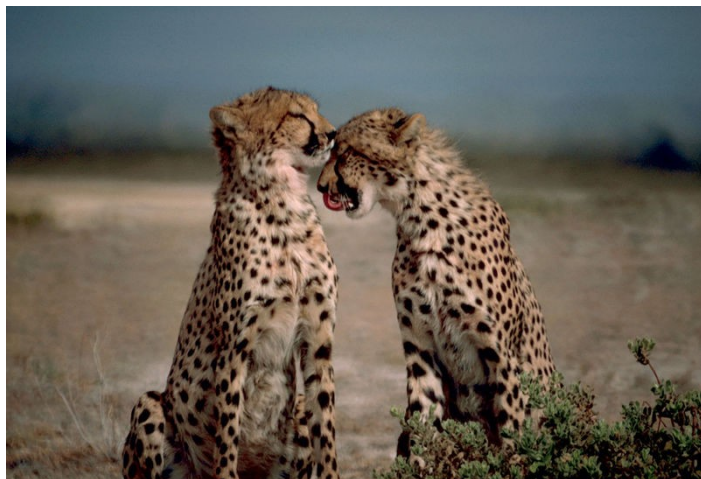
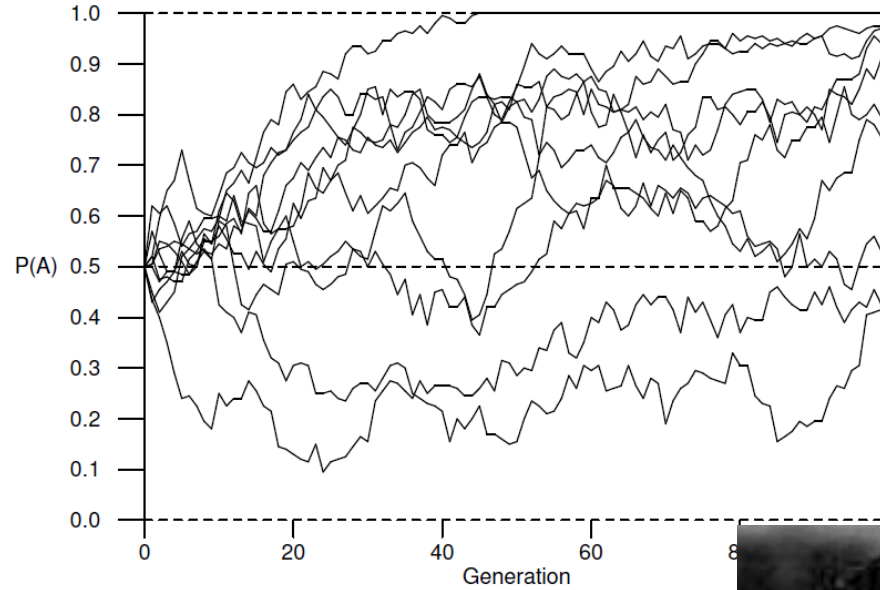


RANDOM GENETIC DRIFT



HW: infinite population but in real world population sizes finite
⇒ random processes, nonadaptive evolution

Why randomness?

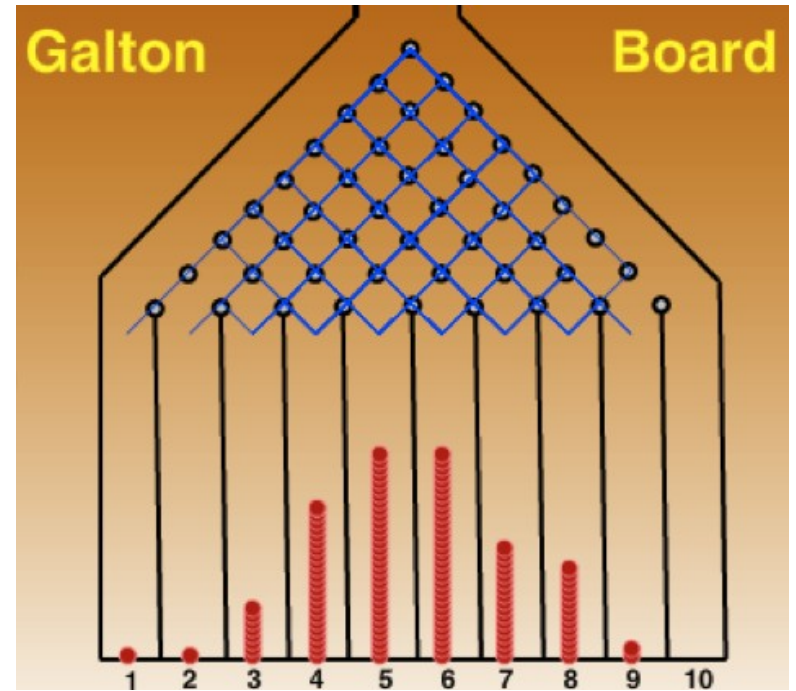
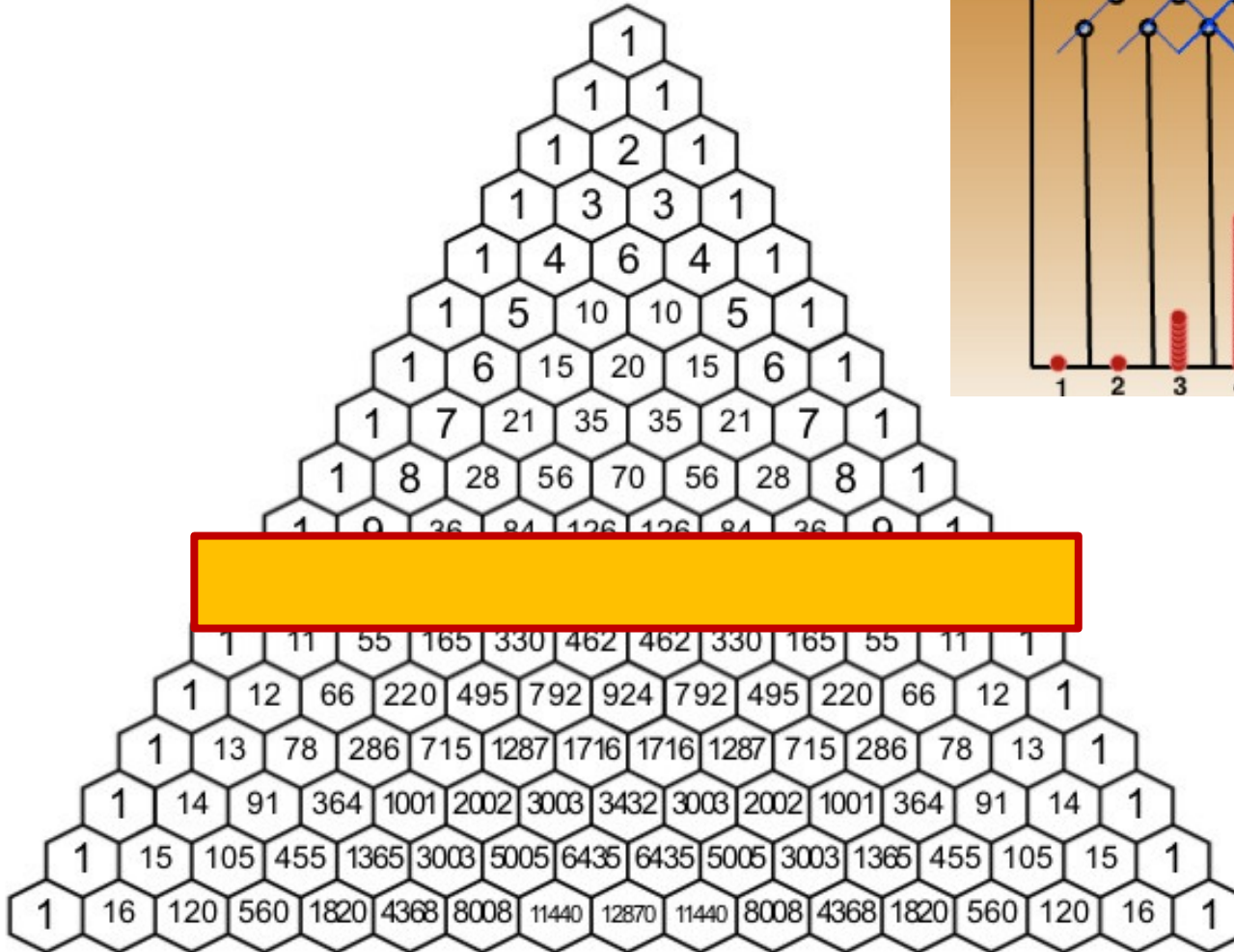
when number of repetitions finite probability of an event \neq its frequency
(cf. H-W principle)



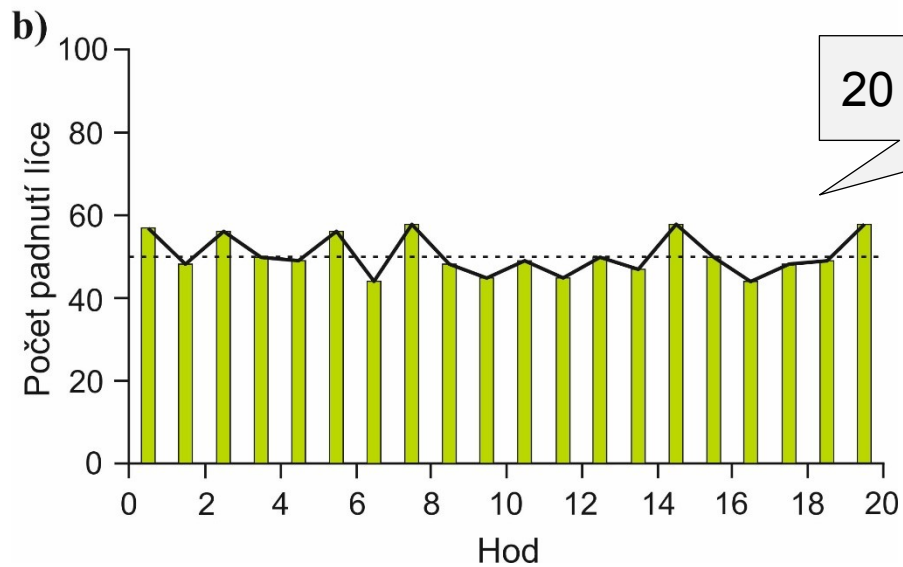
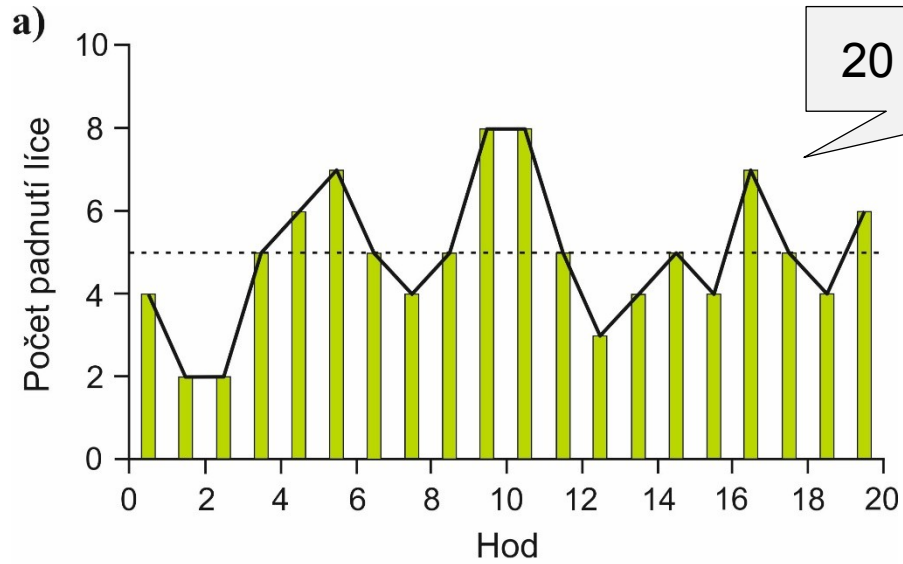
10 coins → in more than 75 % cases the ratio differs from 1 : 1

Pascal's triangle:

11 possible results

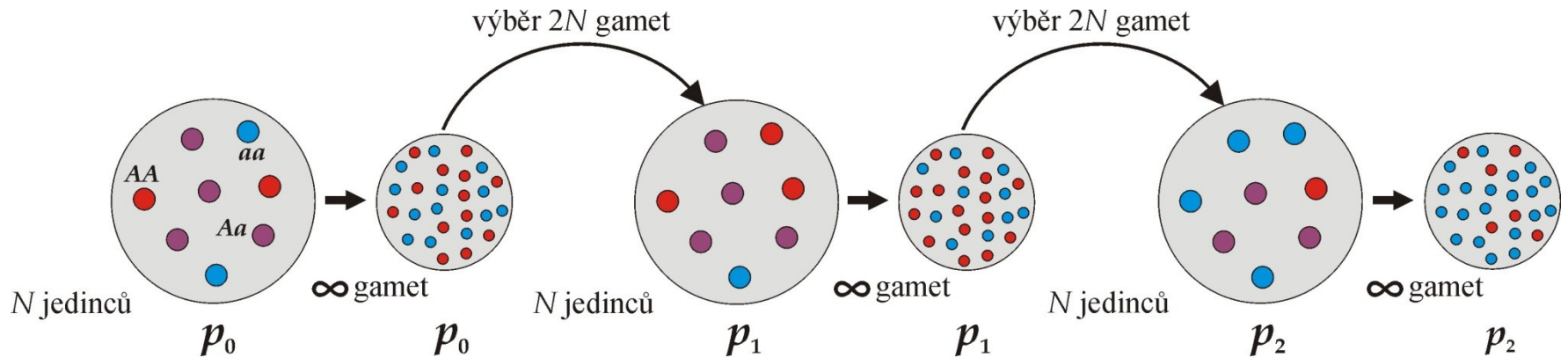


quincunx



With more coins lower variance around expected value

Random sampling from gene pool (sampling error):

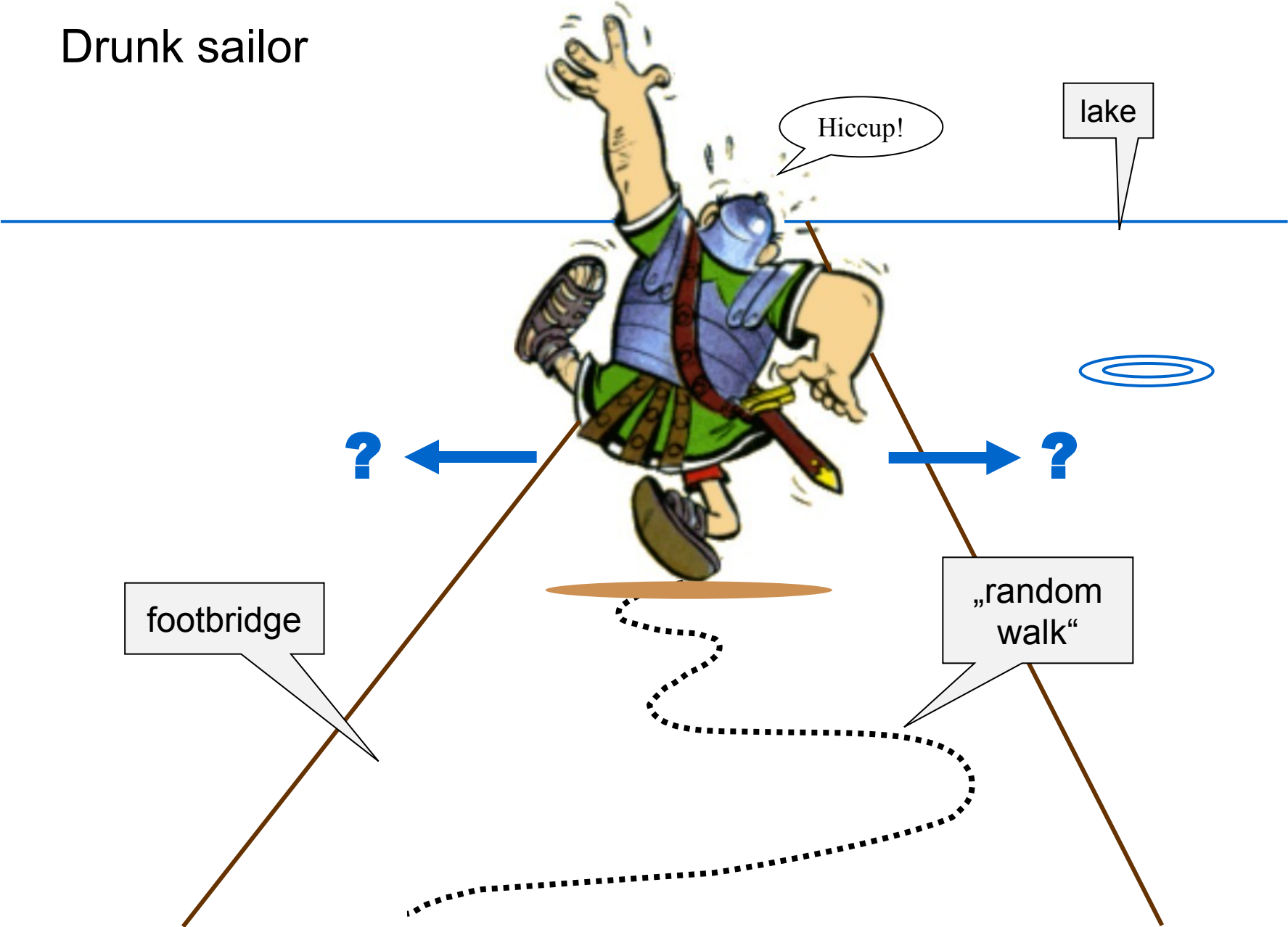


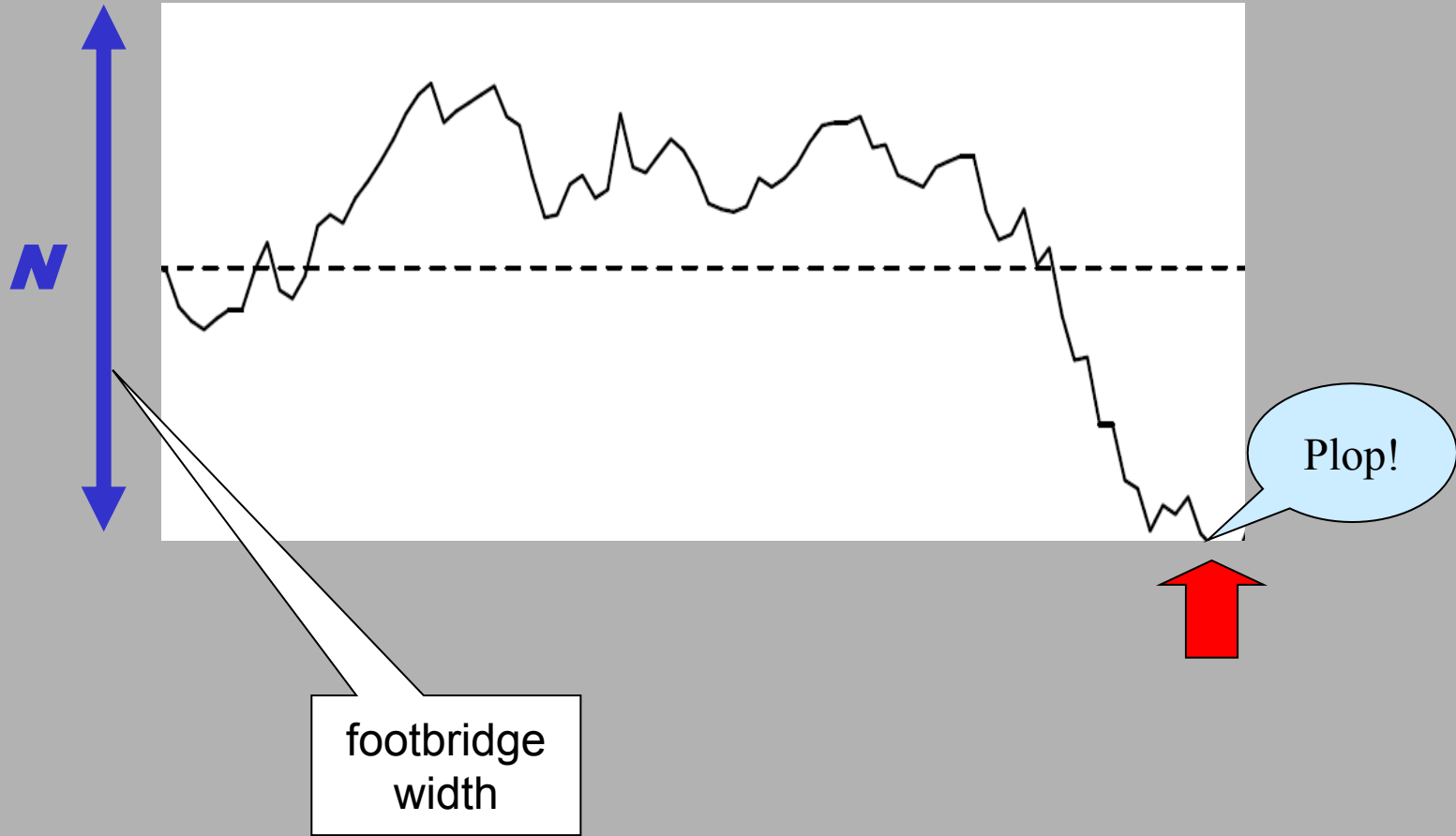
Random sampling results in fluctuations of allele frequencies across generations = „random walk“

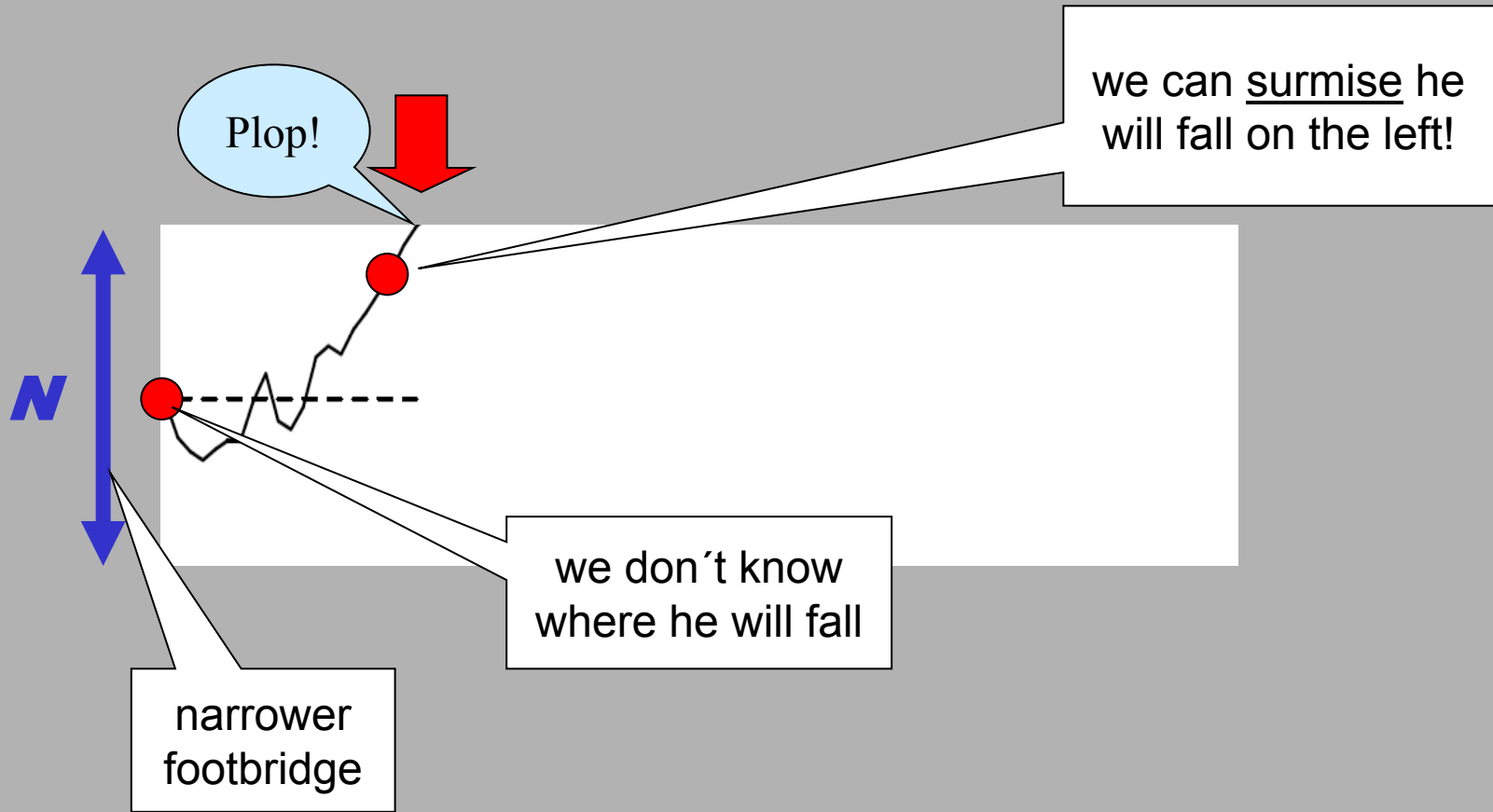
Wright-Fisher model

\approx Hardy-Weinberg model for finite populations

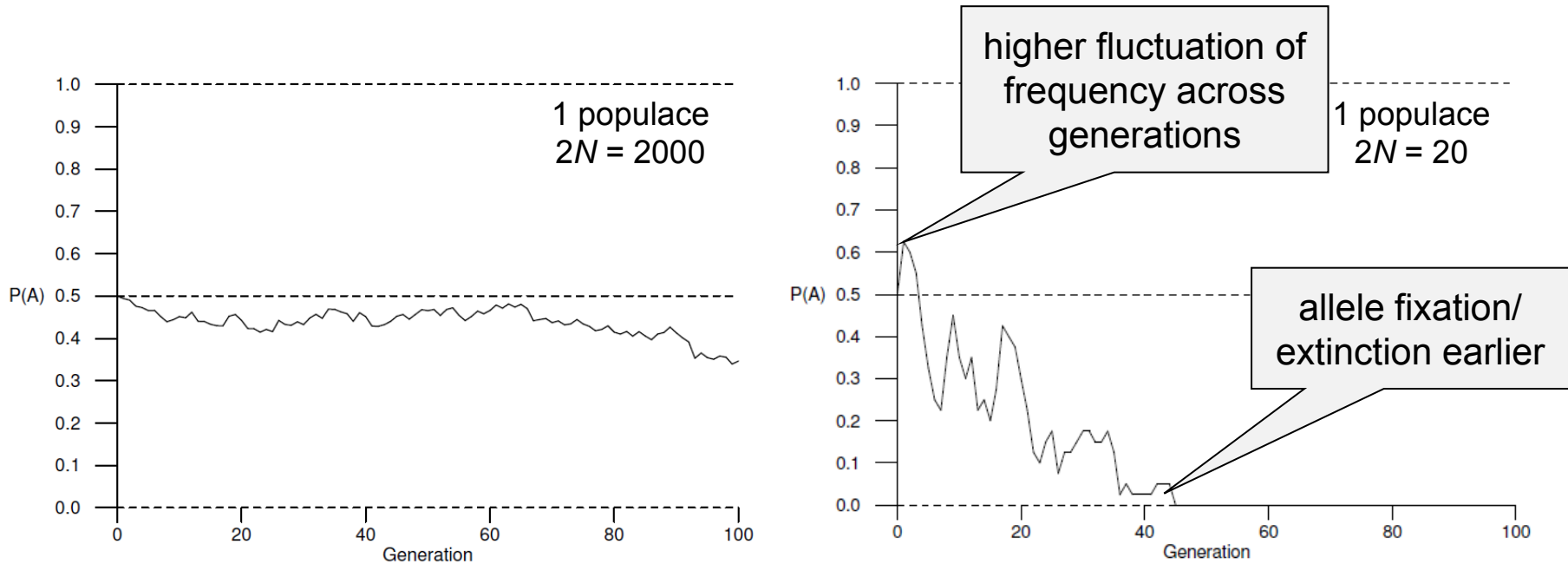
Drunk sailor



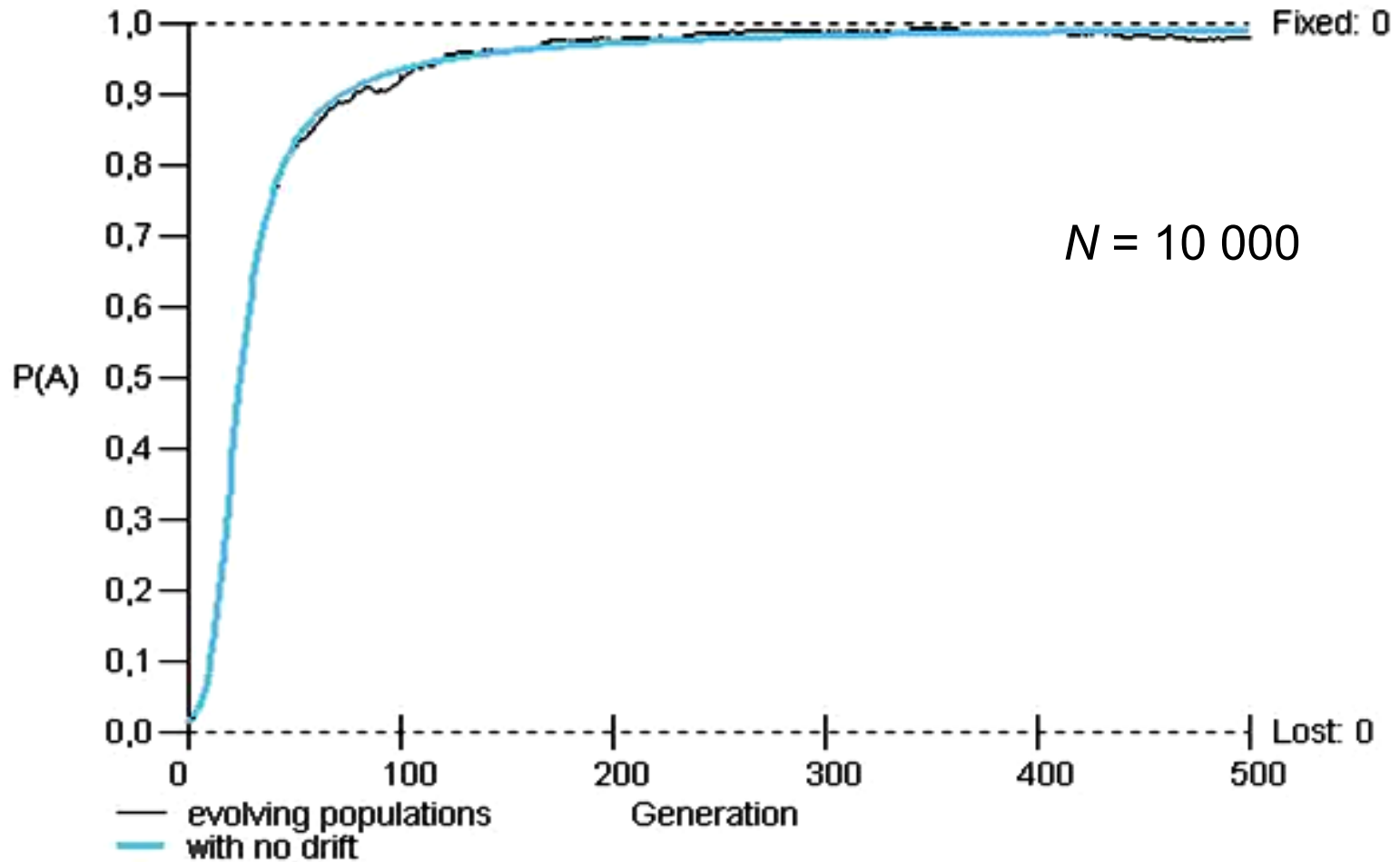




Modelling drift:

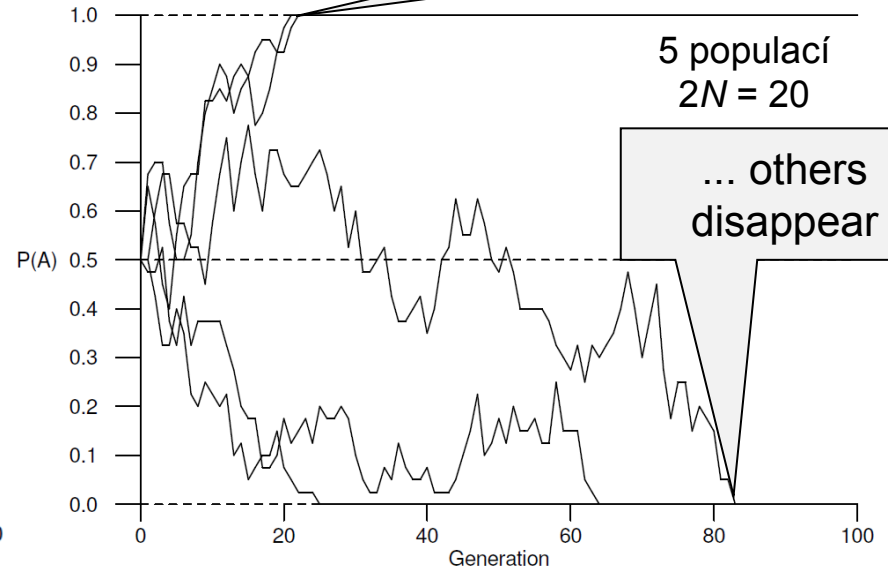
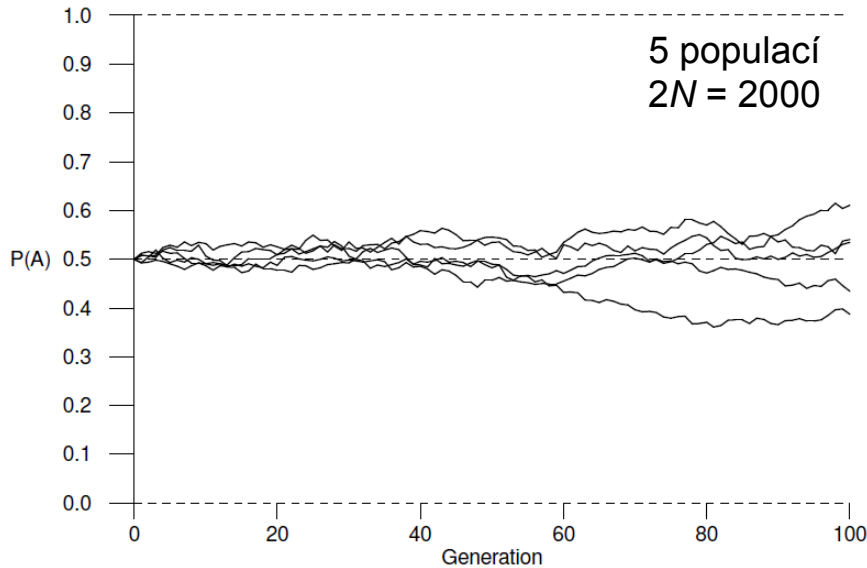


Fluctuation of frequencies across generations stronger in small populations (~ drunker sailor).



Frequency fluctuates even in very large populations!

Modelling drift:



Conclusion 1: Drift results either in allele fixation or allele extinction.

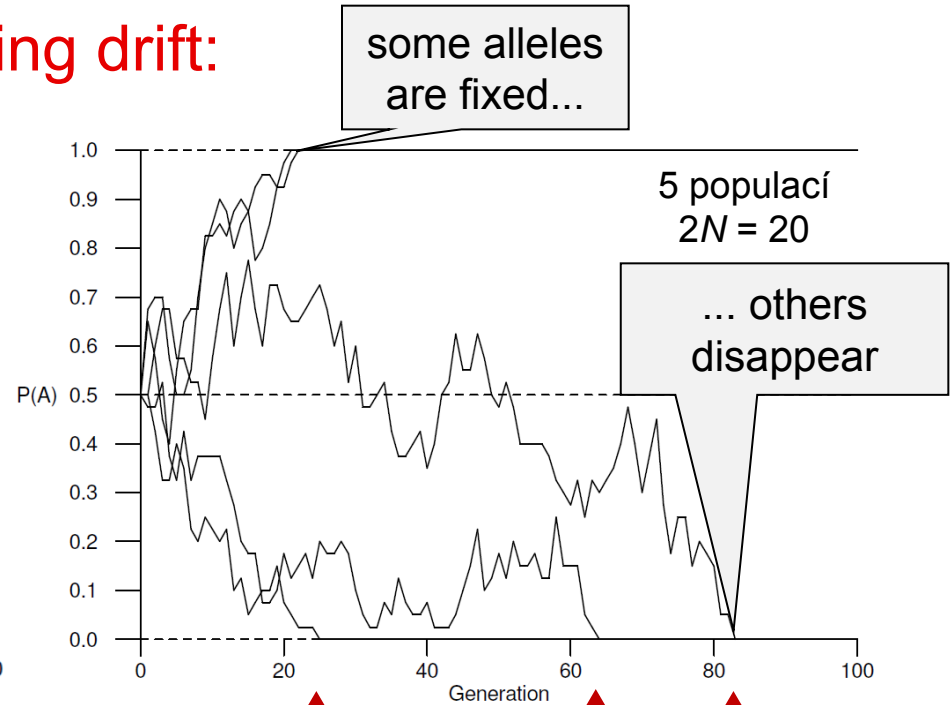
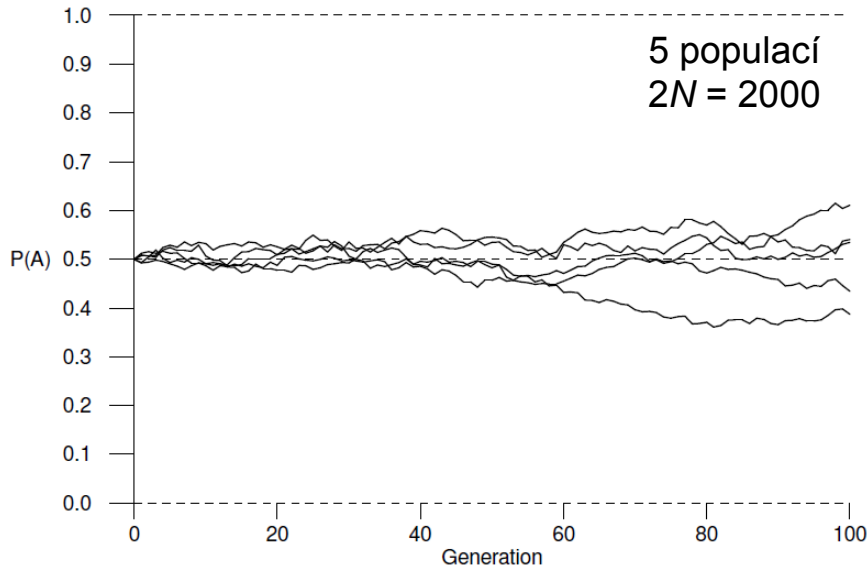
Conclusion 2: Drift results in loss of variation in demes.

Conclusion 3: Probability of allele fixation equals its frequency.

Probability of fixation of a new allele in diploids = $1/(2N)$

Conclusion 4: Mean time to fixation of a new allele $\approx 4N$.

Modelling drift:

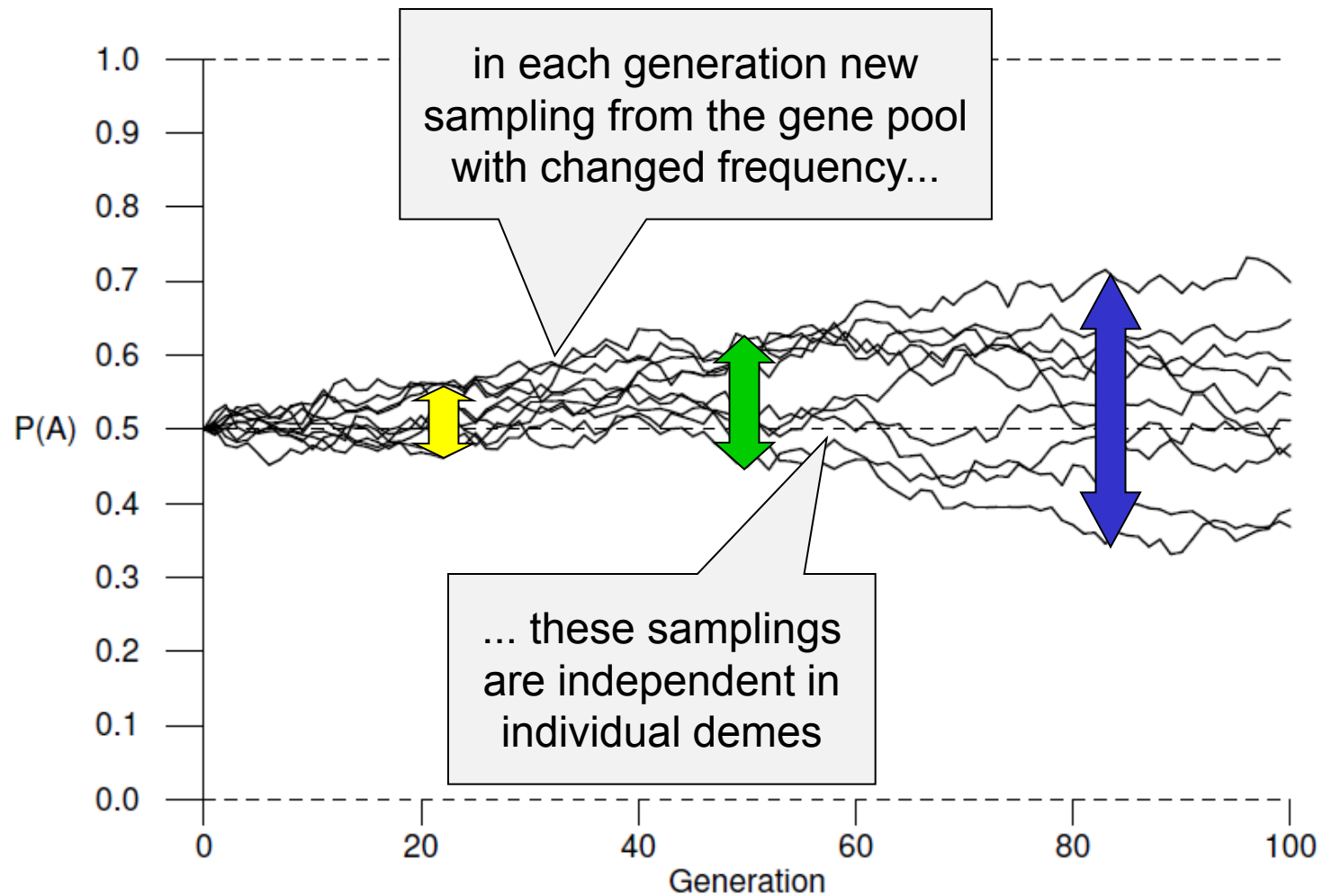


Time to fixation:

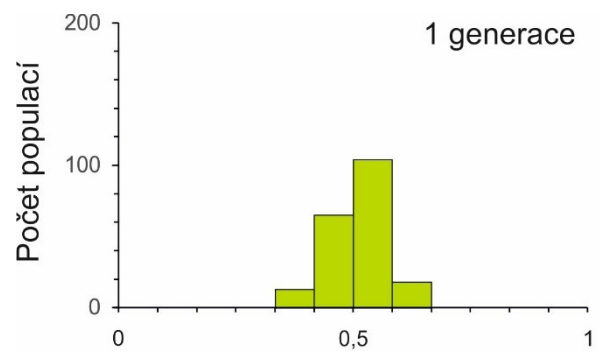
larger population \Rightarrow longer process

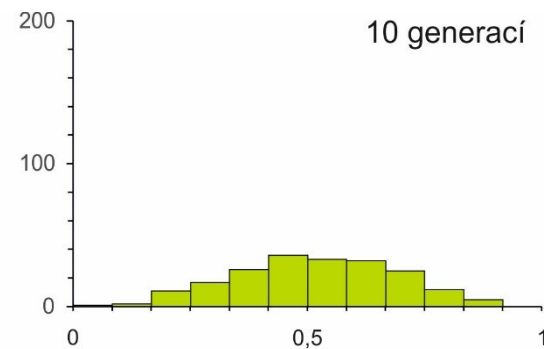
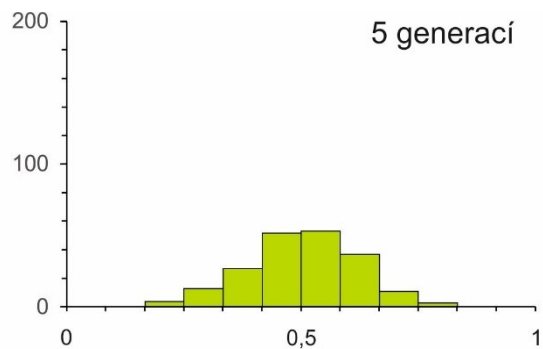
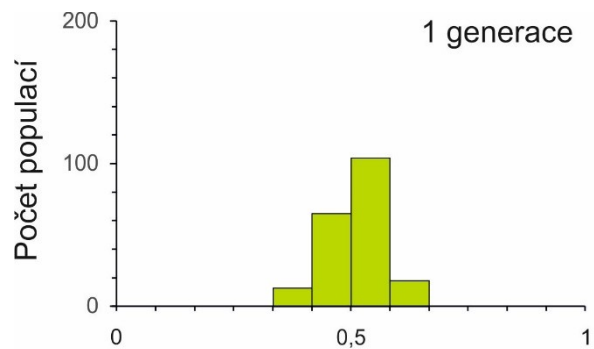
but large variance in the times!!

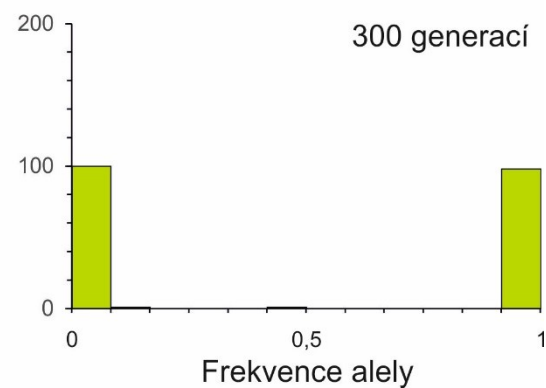
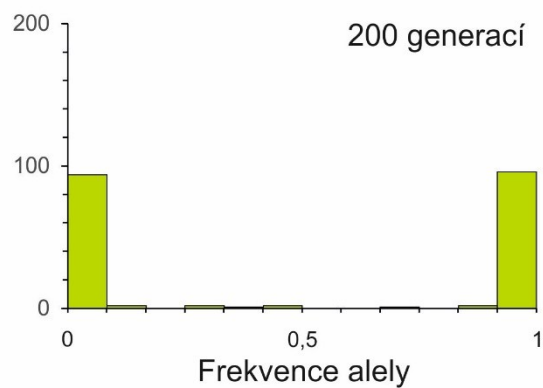
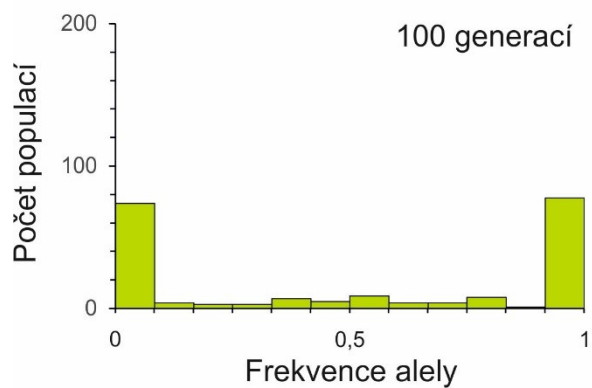
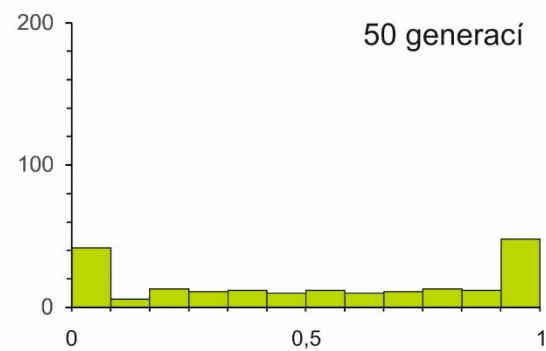
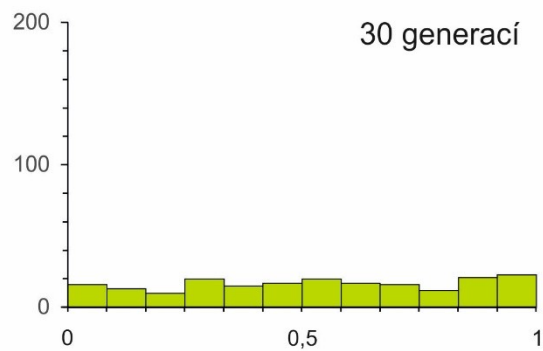
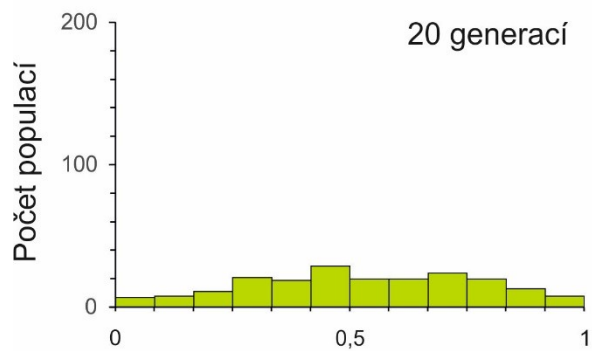
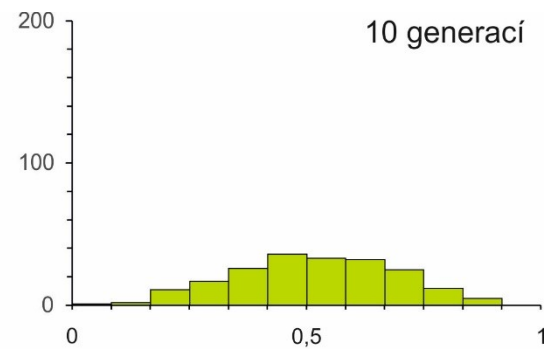
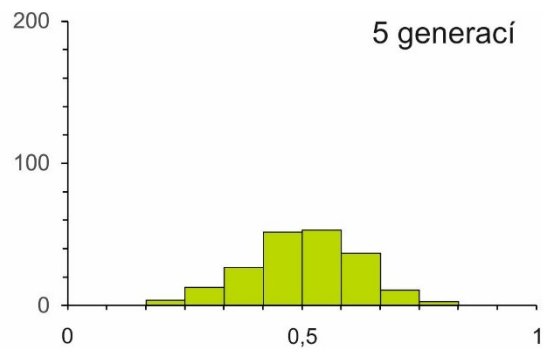
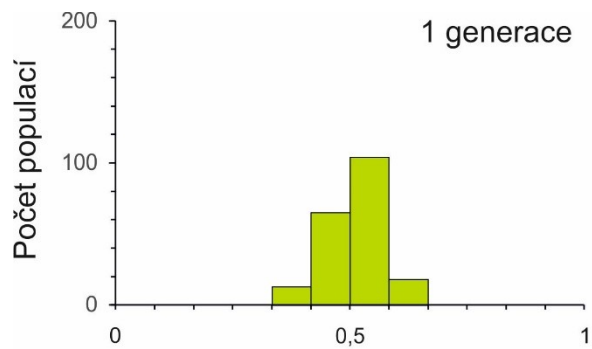
Conclusion 4: Mean time to fixation of a new allele $\approx 4N$.



Conclusion 5: Drift results in divergence among demes.







Peter Buri (1956):

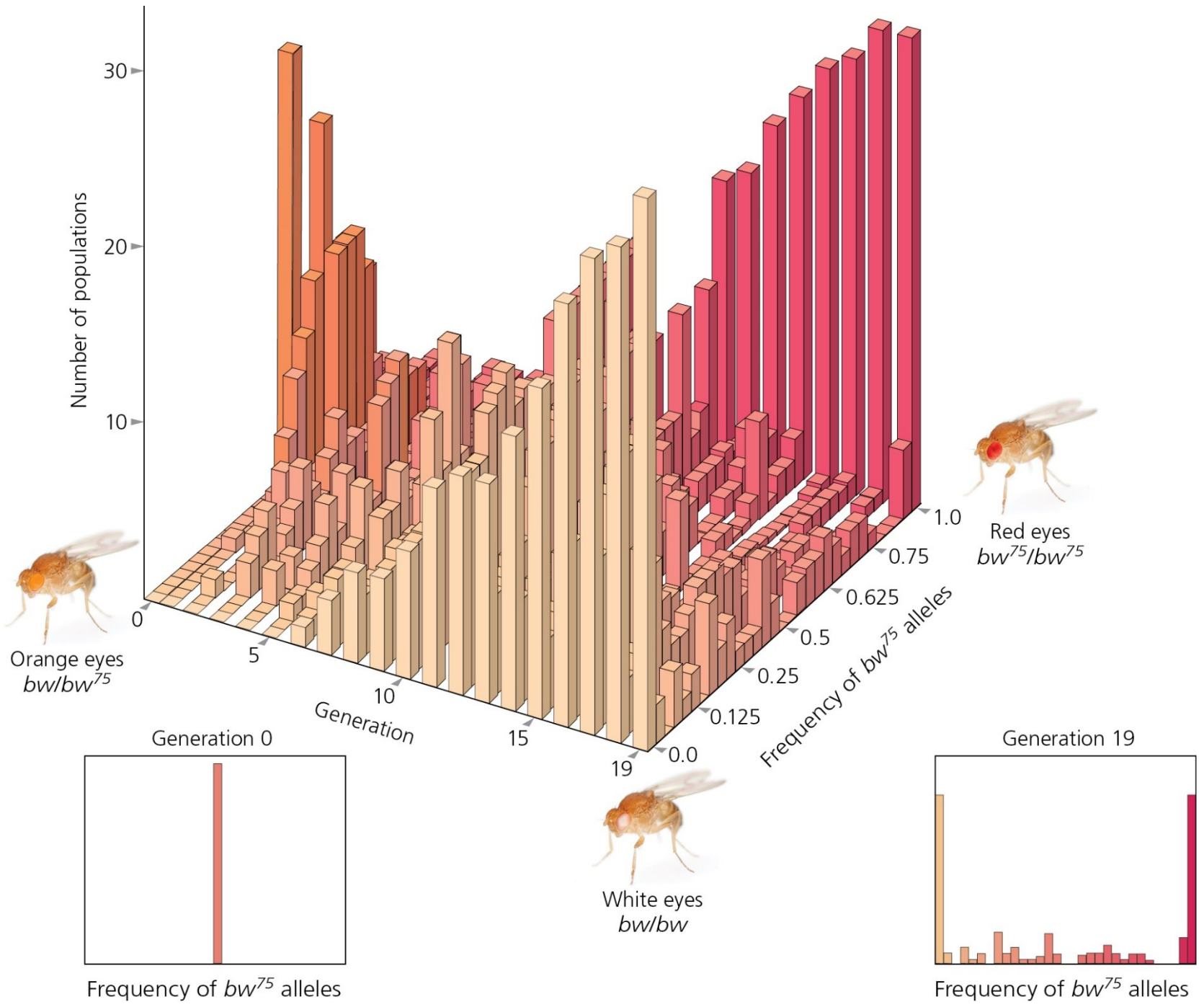
107 populations of *D. melanogaster*

zeroth generation: 16 heterozygous bw^{75}/bw individuals in each population

in each generation random sampling of 8 males and 8 females

19 generations





Number of populations

30

20

10

0

5

10

15

19

0.0

0.125

0.25

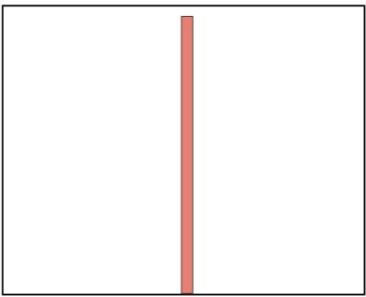
0.5

0.625

0.75

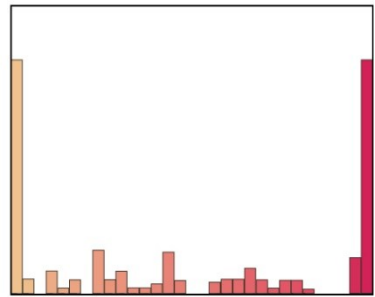
1.0

Generation 0



Frequency of bw^{75} alleles

Generation 19

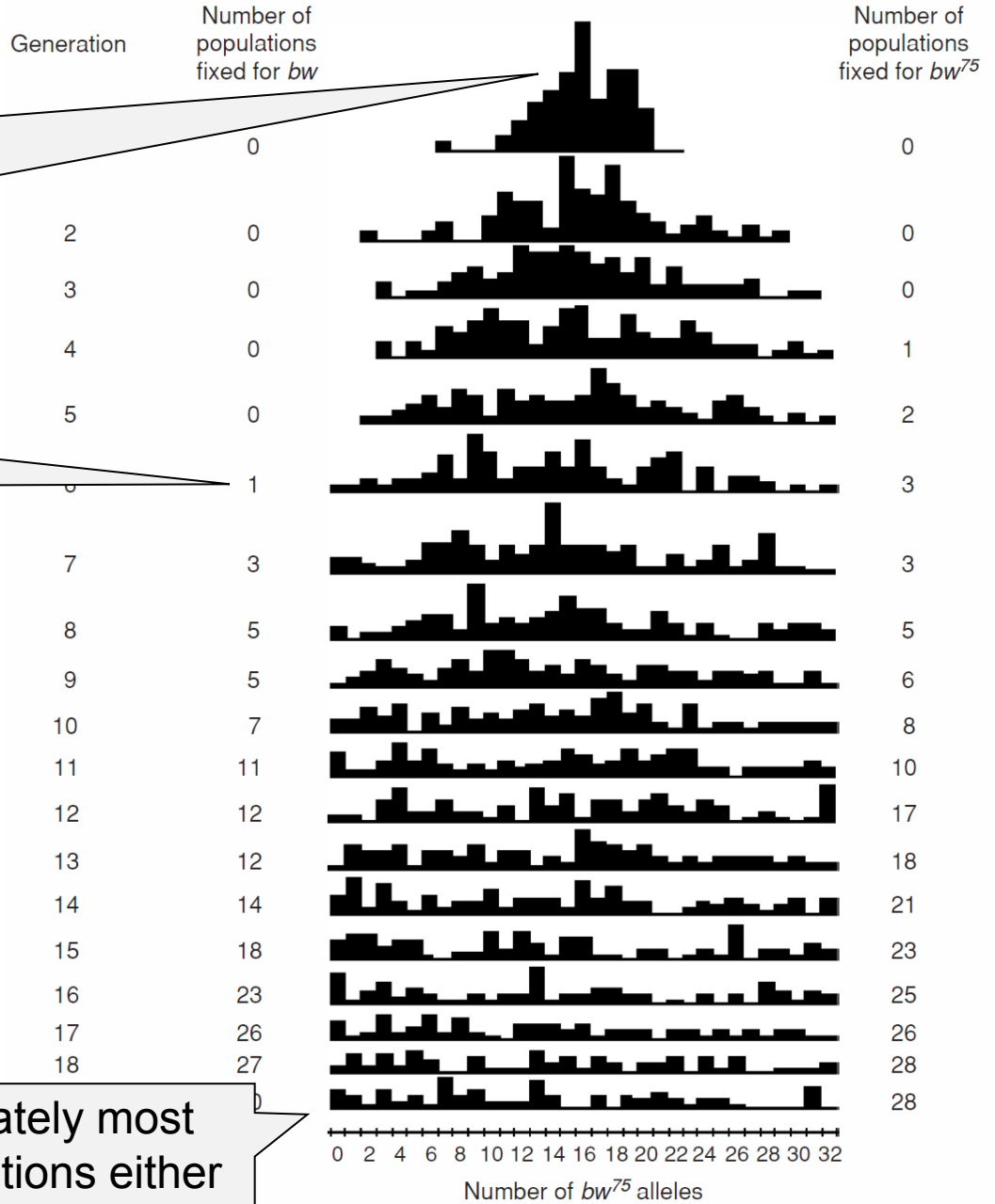
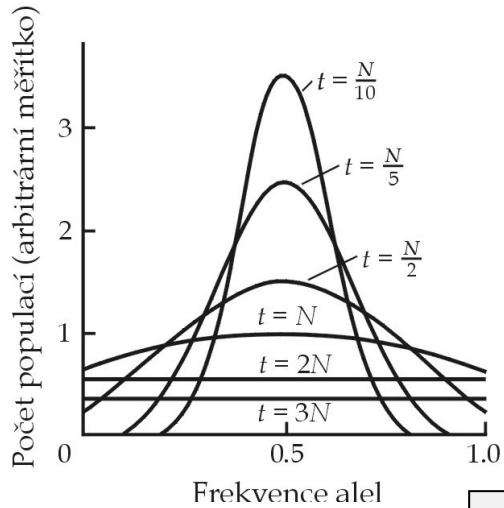


Frequency of bw^{75} alleles

Buri (1956):

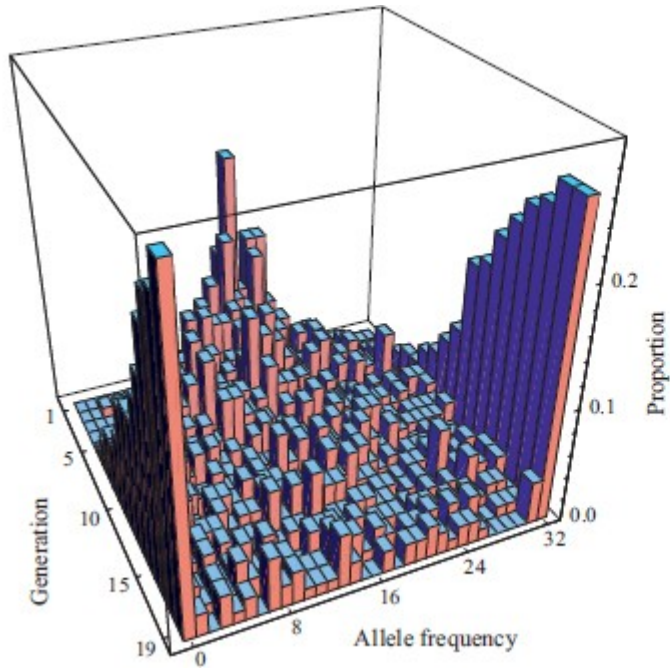
in the first generation most populations around $p = 0,5$

population divergence

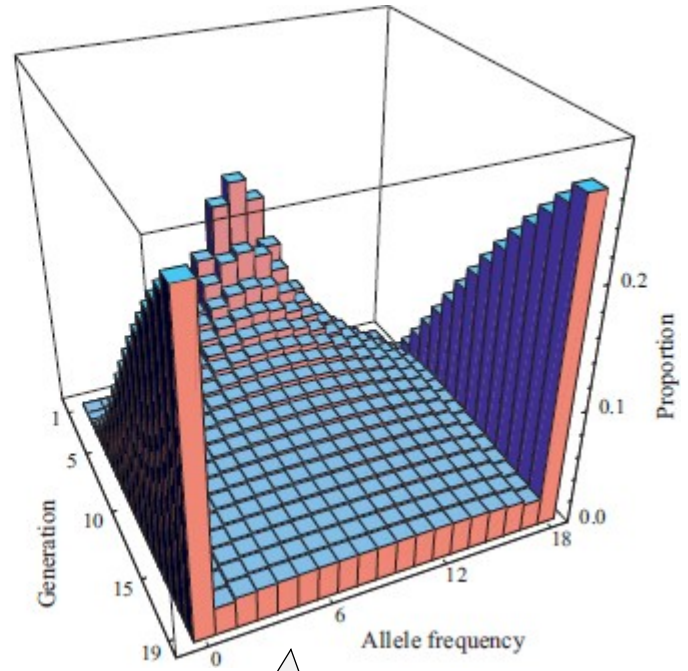


ultimately most populations either $p = 0$ or $p = 1$

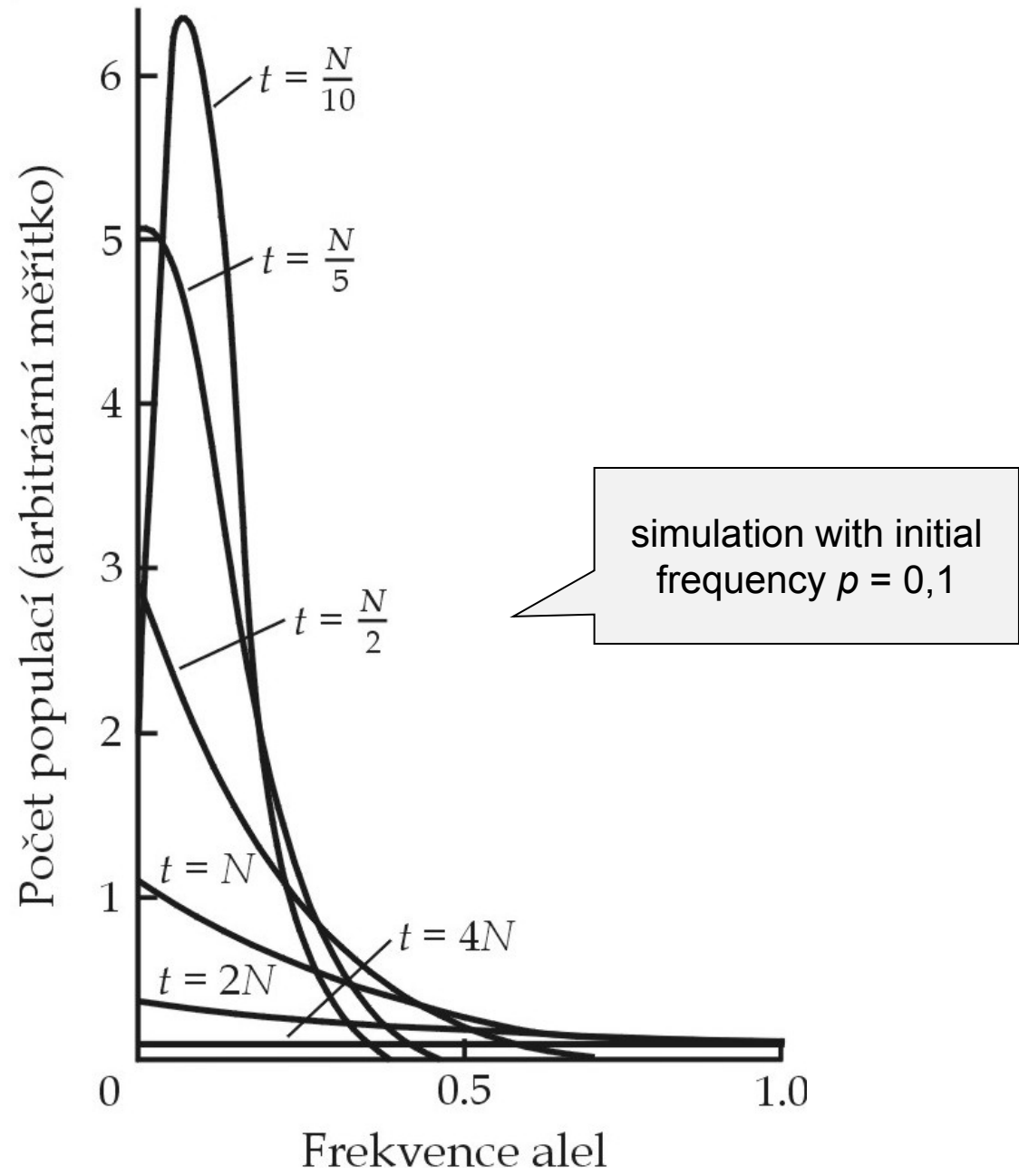
A



B



mathematical
simulation (diffusion
approximation)



Eg.: Galapagos lava lizard (*Microlophus albemarlensis*)

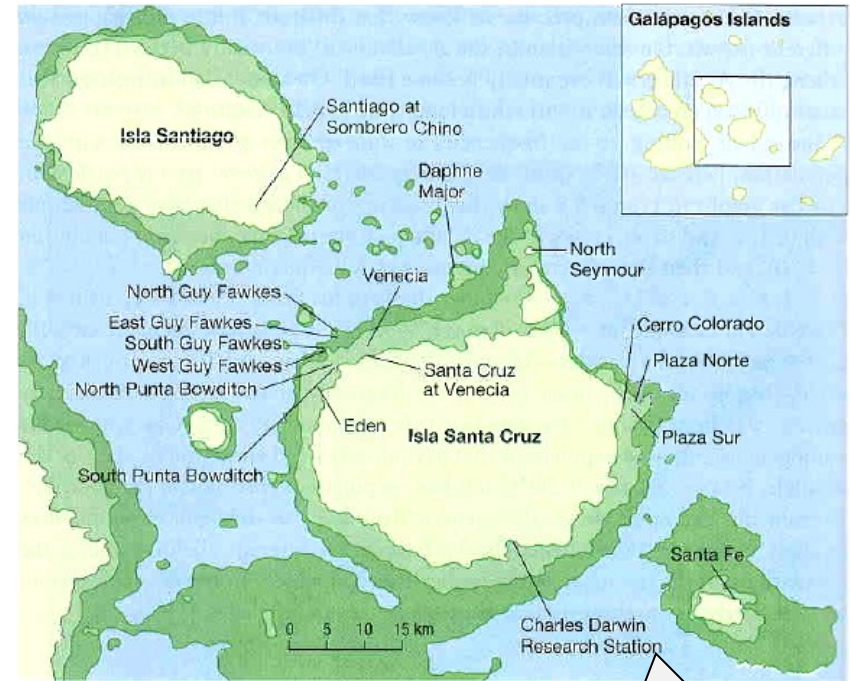
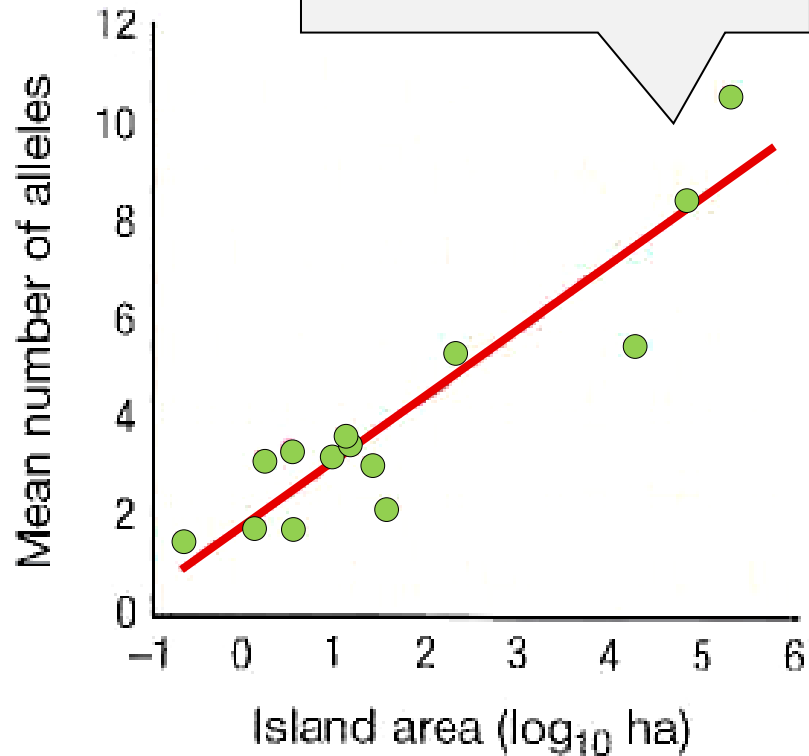


M. Jordan, H. Snell (2002):

17 populations

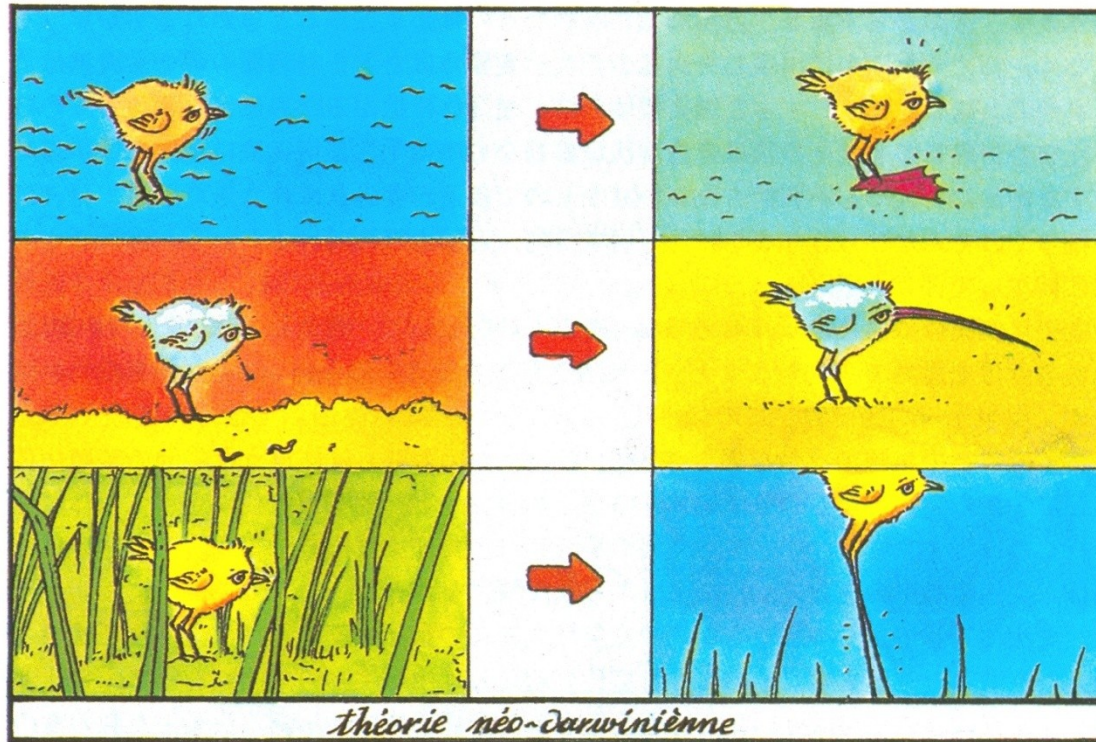
11 microsatellite loci

lizards on larger islands have higher variation

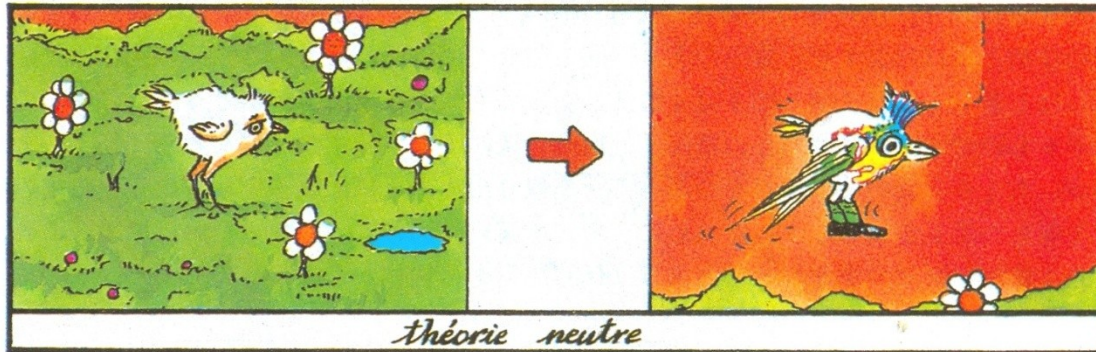


sea level 17 and 12 thousand years ago and nowadays

Evolution of selectively neutral traits is **random**



Darwinian evolution:
„survival of fittest“



neutral evolution:
„survival of luckiest“

Effective population size

Real populations differ from the WF model (fluctuations of N , different reproductive success and mortality, unequal sex ratio,)

→ effective population size N_e allows us to measure drift in non-ideal populations

Effective population size = the number of individuals of ideal Wright-Fisher population displaying the same rate of drift as the studied non-ideal population

Like in the inbreeding coefficient there is no single effective population size!!

Some factors decrease N_e relative to N :

overlapping generations

fluctuating population size across generations

different number of breeding males and females

high variation of the number of offspring within populations

**Caution! Under some circumstances
the effective population size can be higher than N !!**

Effect of fluctuating population size:

effective size can be approximated as harmonic mean \Rightarrow strong influence of small N !!

$$N_e = \frac{t}{\frac{1}{N_1} + \frac{1}{N_2} + \dots + \frac{1}{N_t}}$$

harmonic mean

mean much closer to the lower value

Find HARMONIC MEAN
of 1 AND 100

$$H = \frac{2}{\frac{1}{a_1} + \frac{1}{a_2}}$$
$$= \frac{2}{\frac{1}{1} + \frac{1}{100}}$$

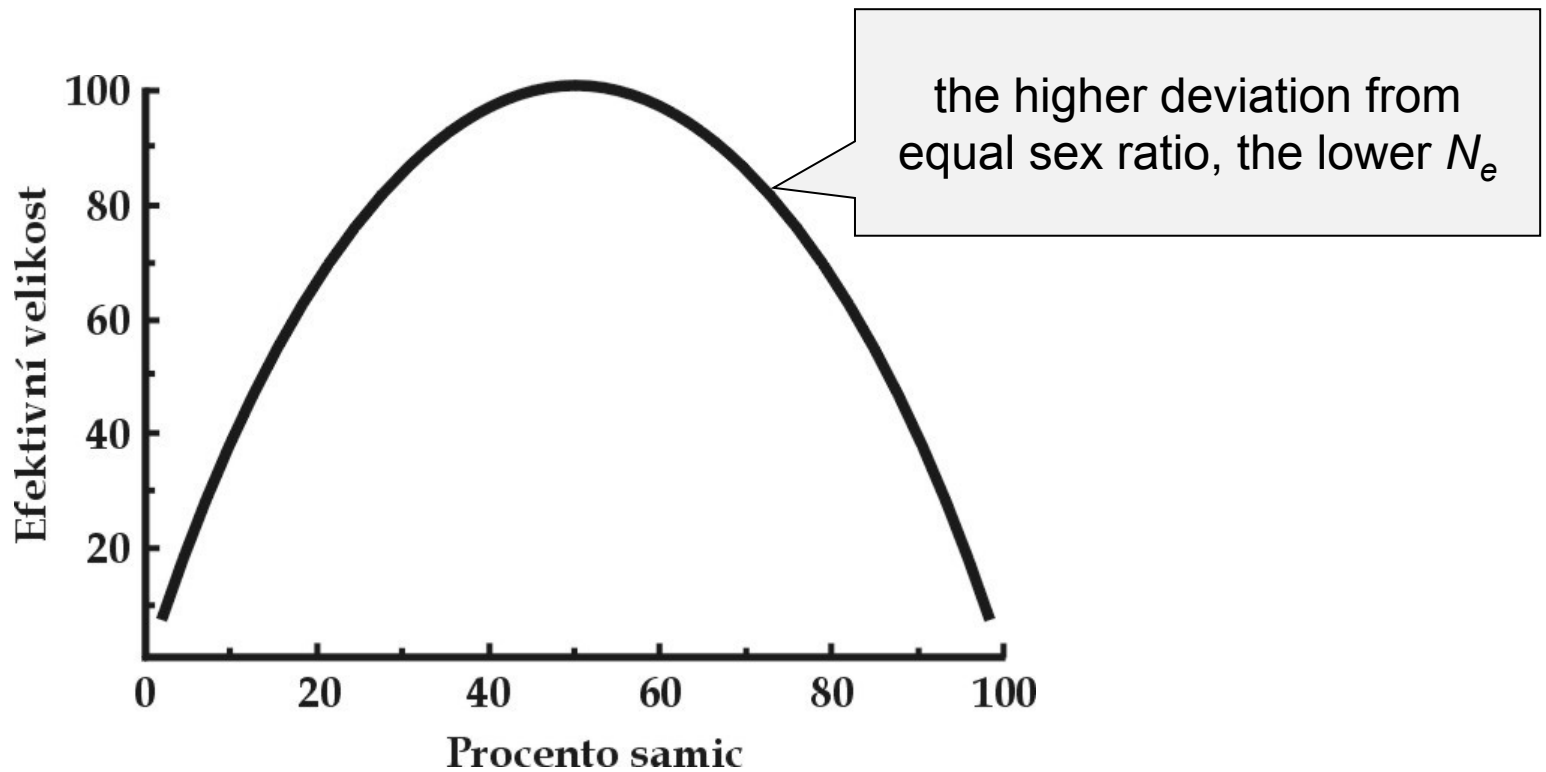
$$= \frac{2}{1.01}$$
$$= 1.9802$$

Harmonic Mean
of 1 and 100

Effect of biased sex ratio:

Till now we assumed the same number of breeding males and females

N_m = number of breeding males, N_f = number of breeding females



Effect of biased sex ratio:

$$N_m = 1:$$

$$N_e = \frac{4N_m N_f}{n + c}$$



$$N_e = \frac{4N_f}{1 + c} \approx 4$$

it follows from this formula that if there is only a single breeding male in the population $N_e \approx 4$ regardless of the total number of individuals

Effect of unequal reproductive success:

southern elephant seal:

sex ratio within a harem 1:40^{*)}

^{*)} effective ratio 1:4-5 due to cuckoldry and short period of male's dominance (1-2 years)



Reproductive success on the gene level:

If a gene is affected by selection variance of the number of offspring among members of a population is high (individuals with a positive allele have more offspring)

⇒ N_e for this gene is lower than for a neutral gene

Each genetic trait requires its own N_e :

For genes on autosomes, sex chromosomes, and mtDNA there are different effective population sizes:

autosomes:	N_e	$4 N_e$
X, Z:	$\frac{3}{4} N_e$	$3 N_e$
Y, W, mtDNA:	$\frac{1}{4} N_e$	$1 N_e$

COALESCENT

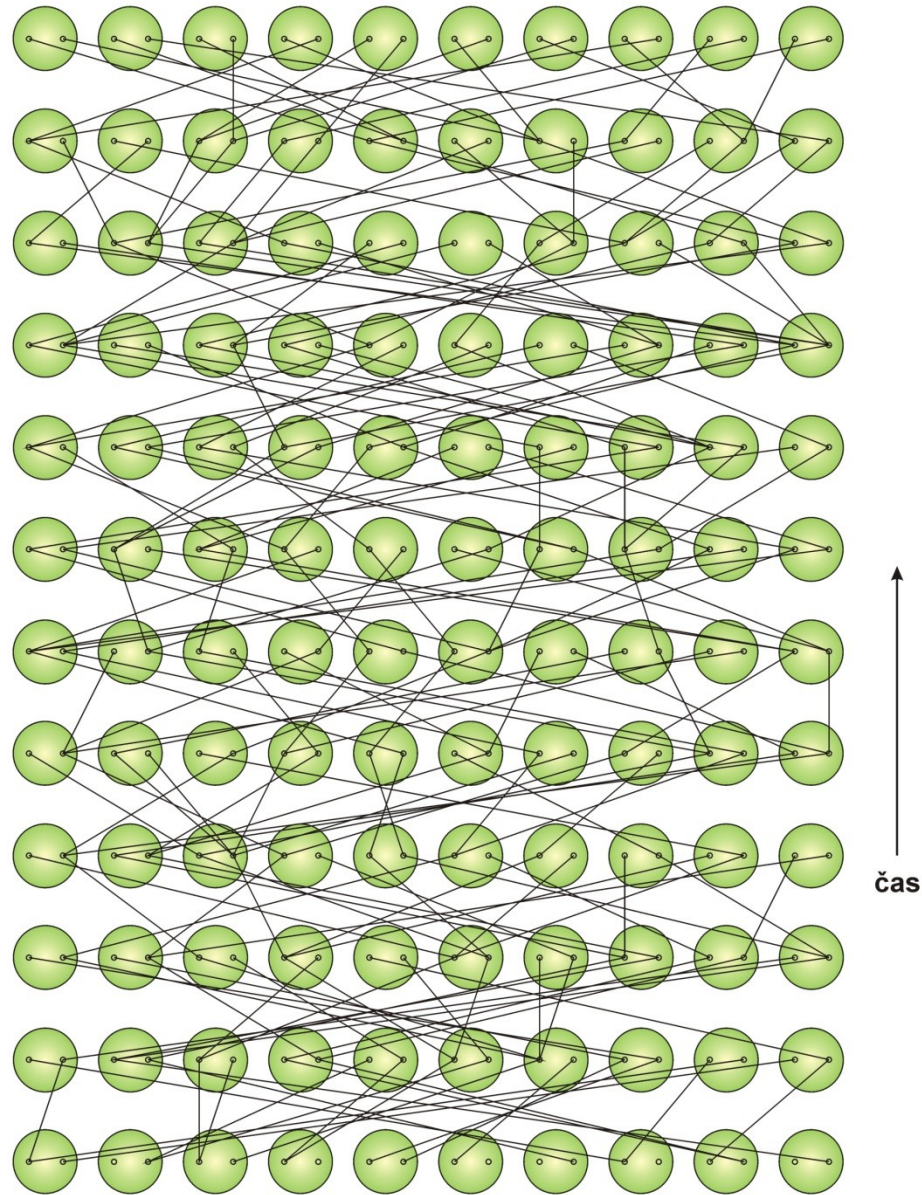
under drift some alleles disappear from a population \Rightarrow when there are no mutations ultimately all gene copies have a common ancestor

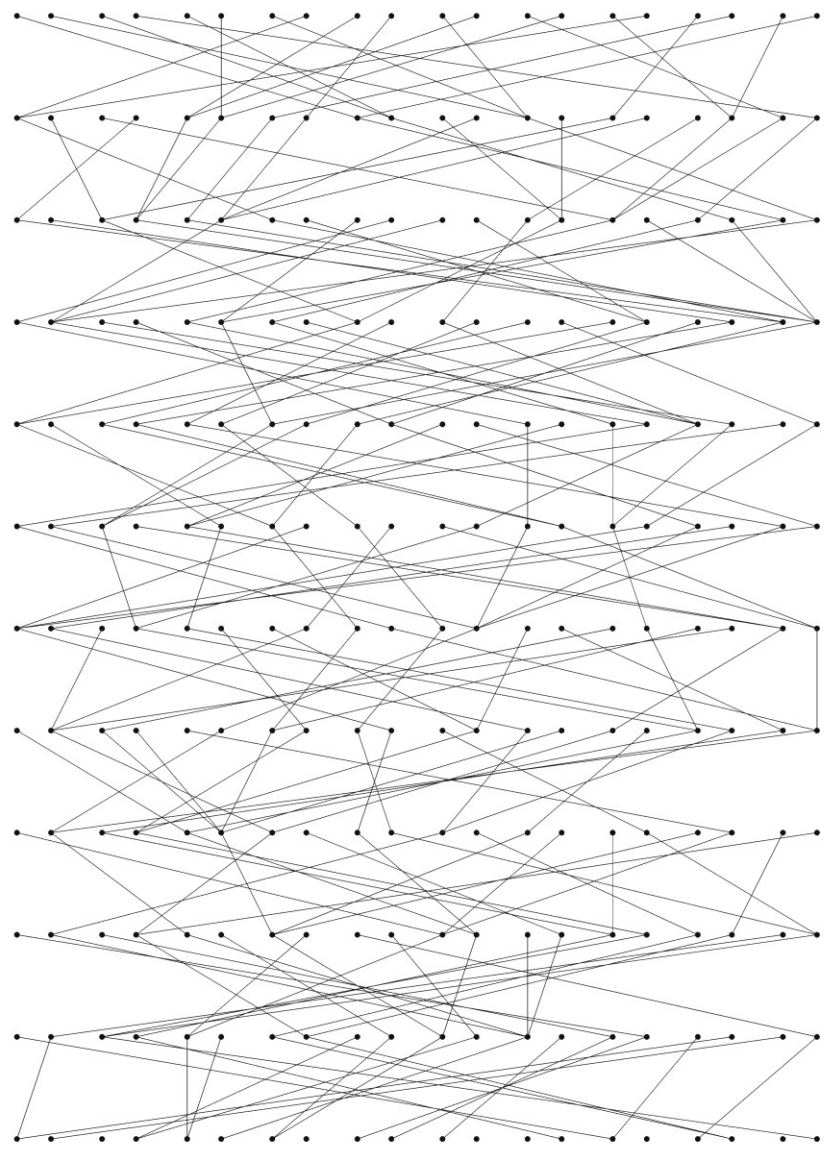
„forward“ approach

we can proceed also back in time – „backward“ approach \rightarrow
moving back in time till two or more gene copies „fuse“
= **coalescent event**

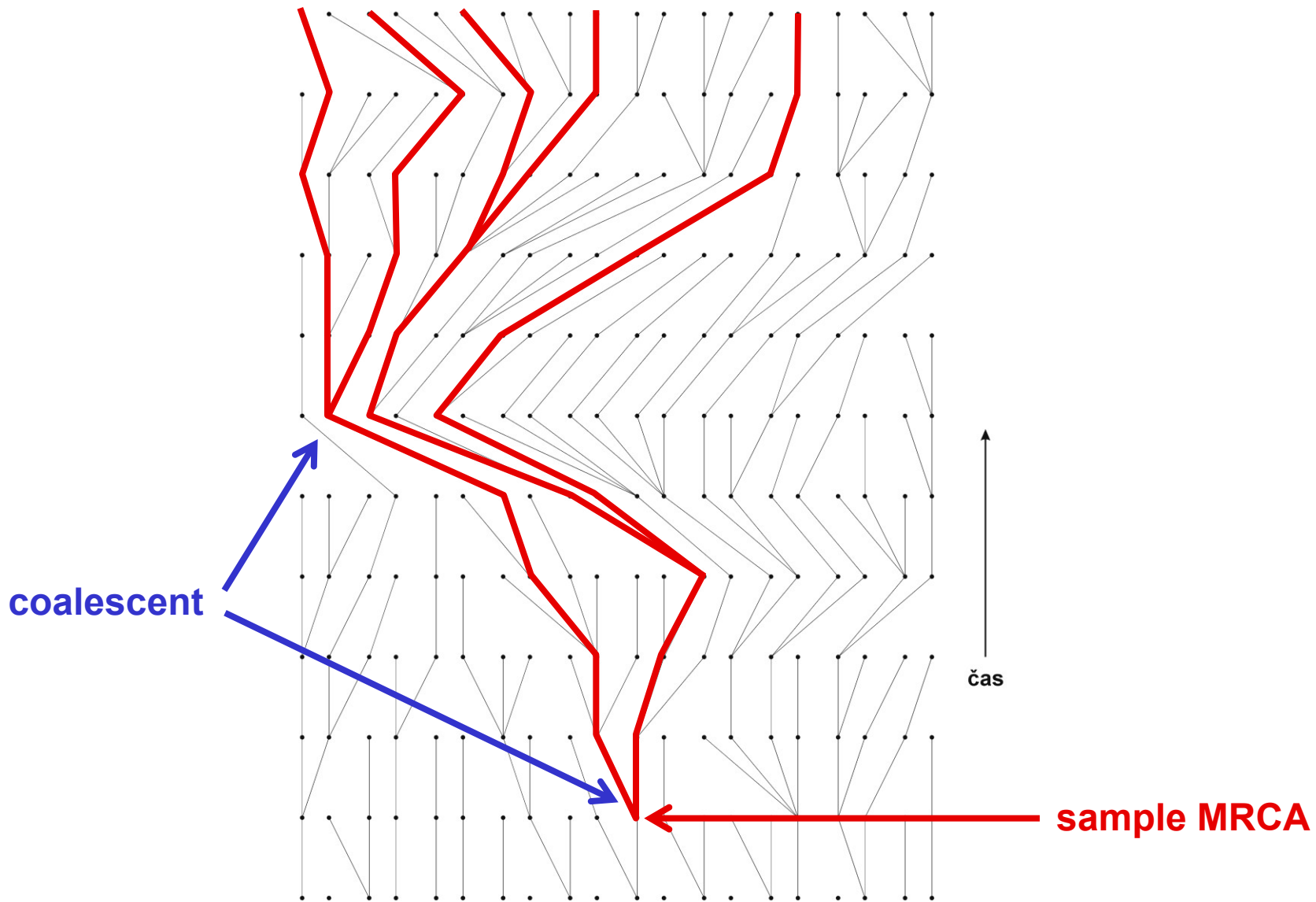
the most recent common ancestor (MRCA)

Wright-Fisher model:





↑
čas



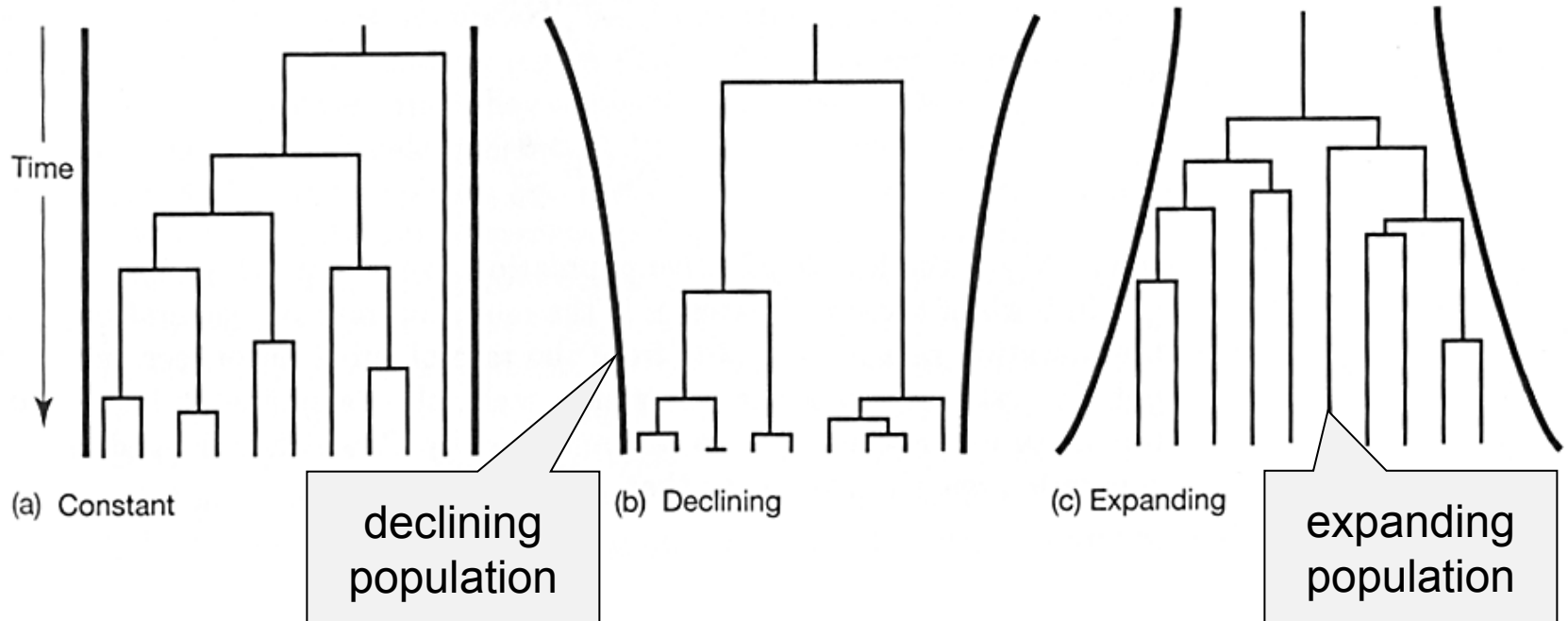
Coalescence and effective population size

from the coalescent theory several interesting consequences follow:

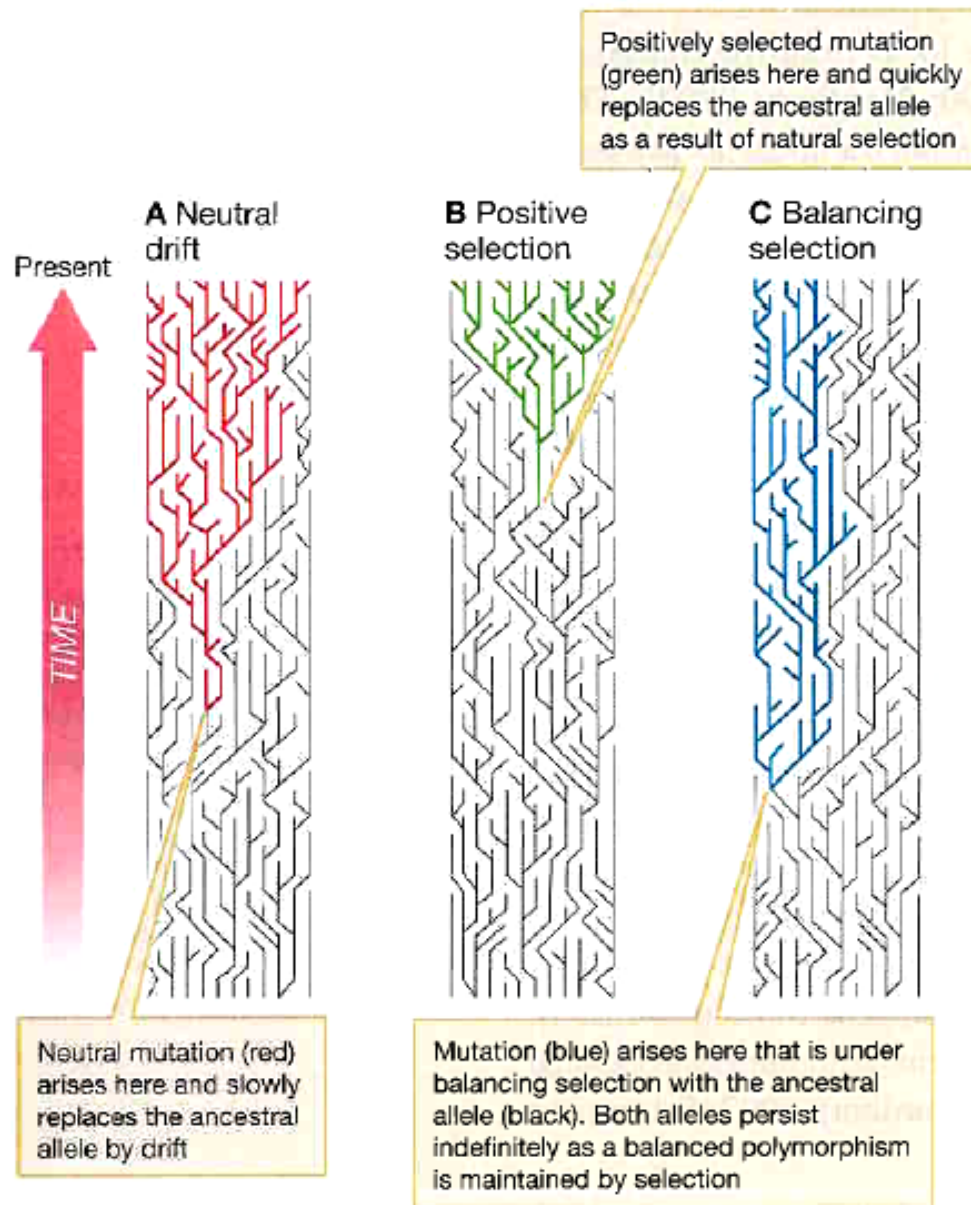
in small populations coalescent rate higher than in large populations

⇒ we can estimate N_e

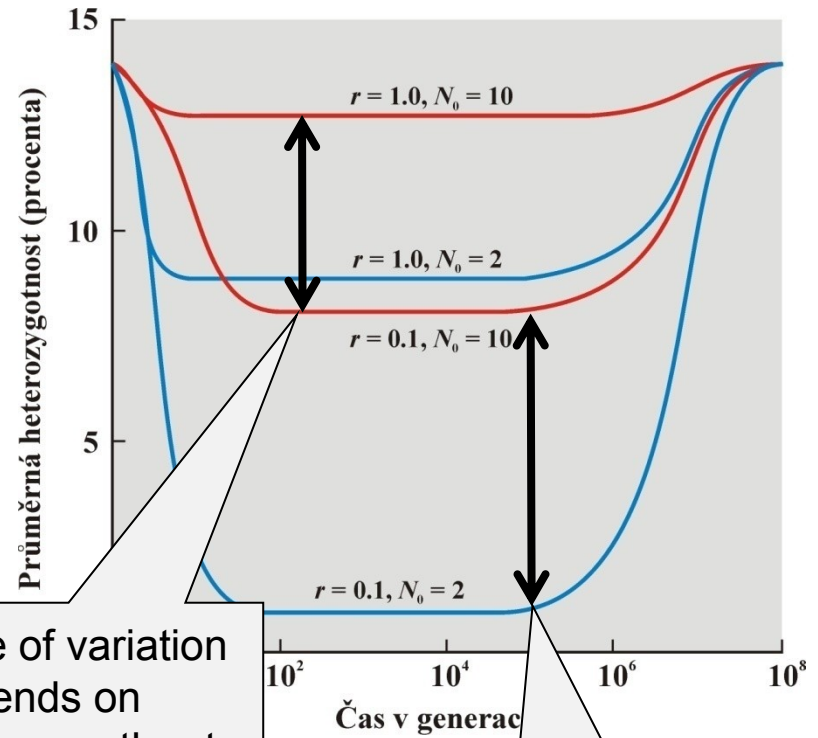
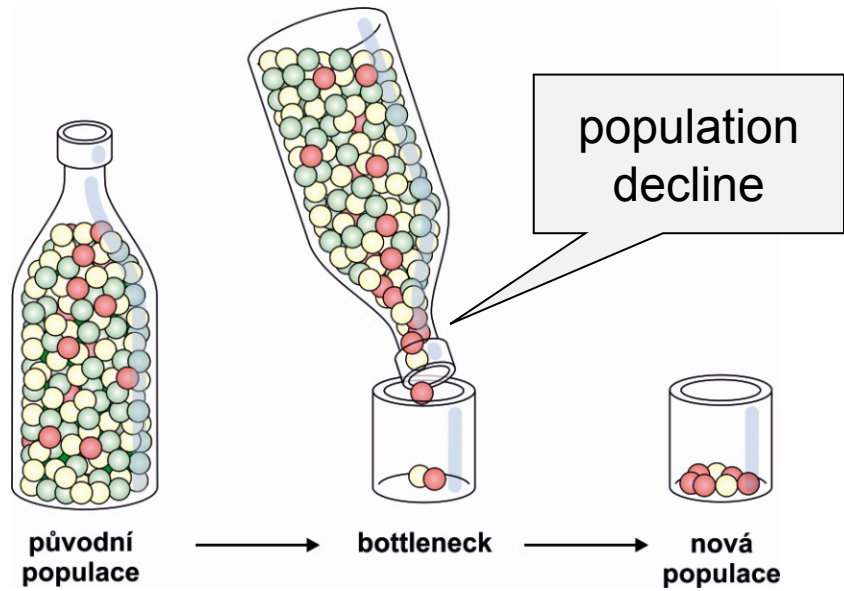
but we can estimate also changes of N_e in time



The same effect of selection on the coalescent tree shape:



BOTTLENECK and FOUNDER EFFECT



bottleneck reduces variation

magnitude of this reduction

depends on reduction of N_e and duration of bottleneck

rate of decrease of variation different for various genetic traits (autosomes, mtDNA, Y...) – different N_e !

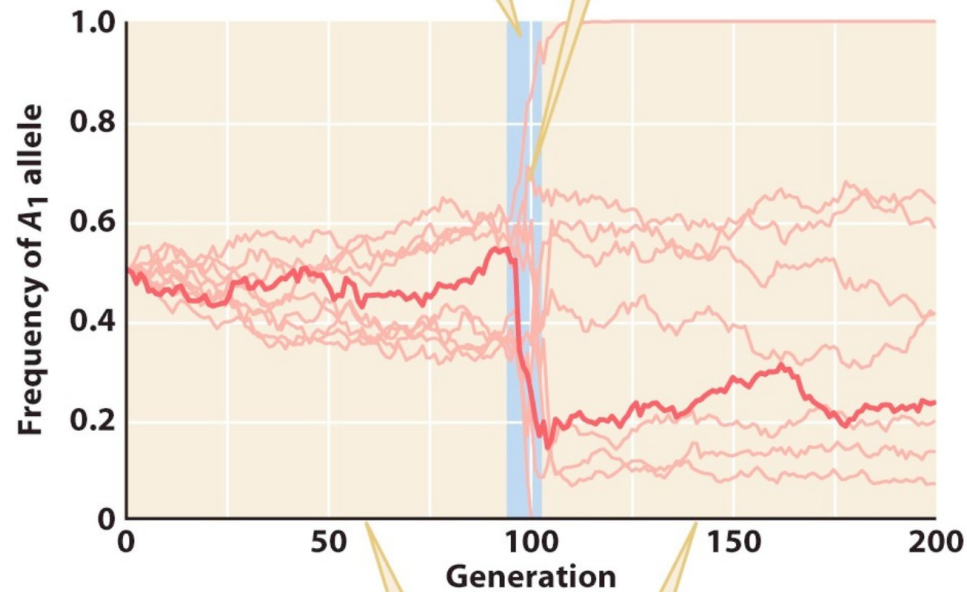
decrease of variation depends on population growth rate

variation more reduced under stronger bottleneck

Bottleneck:

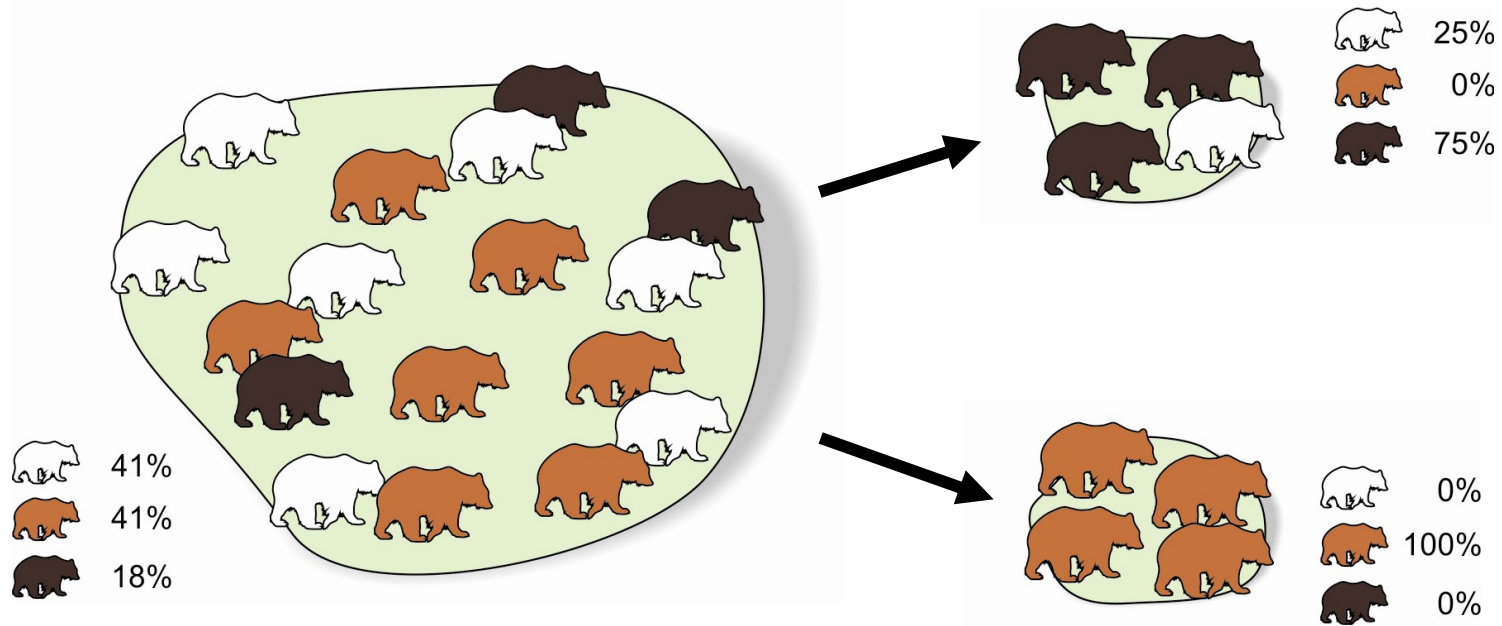
Populations experience a bottleneck in size during the period indicated by the shaded region and return to the original size of 1000 individuals afterward

Allele frequencies fluctuate much more during the bottleneck than before or after



The bottleneck causes divergence between populations. Before the bottleneck, allele frequencies are similar in all populations. After the bottleneck, allele frequencies differ greatly from one population to the next

Founder effect:



colonization of a novel territory (eg. island)

because of a small number of founders (even a single pregnant female)

→ random change of allele frequencies

→ reduction of variation

different environmental conditions → speciation

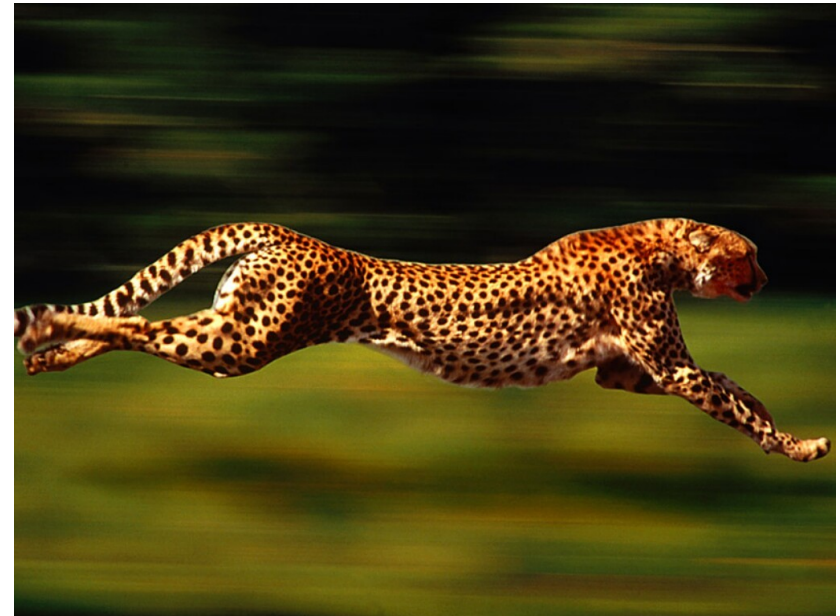
Examples of founder effect and bottleneck cheetah

30 individuals of *Acinonyx jubatus reineyi* from E Africa, 49 protein loci:
only 2 loci polymorphic ($P = 0,04$), mean heterozygosity $H_o = 0,01$

98 individuals of *A. j. jubatus* from S Africa: $P = 0,02$, $H_o = 0,0004$!

south-African individuals accept skin grafts of the east-African subspecies
without problems \Rightarrow monomorphism of MHC genes

assumed strong bottleneck
in the past



golden hamster

1930: [Israel Aharoni](#) (Hebrew Univ., Jerusalem) – female with offspring

escape of several individuals from captivity

1931: transport of several individuals to Britain 1937: private breeders

Recent genetic analyses including mtDNA → all golden hamsters currently kept in breeding colonies are descendants of a single female, probably that of 1930

mostly presented as an example of bottleneck but it is rather an example of founder effect



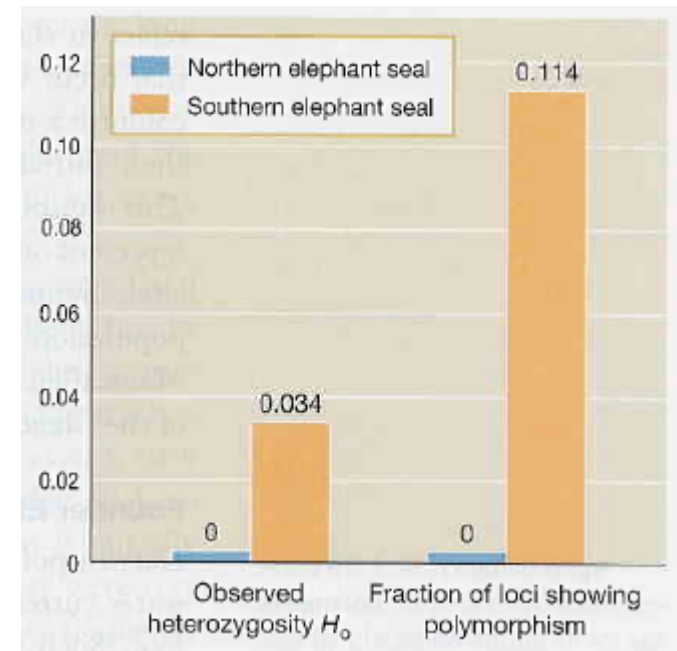
northern elephant seal

Mirounga angustirostris: in 19th century almost eradicated

→ 1892 last 8 individuals on the island of Guadelupe killed for museum collections

fortunately 10-20 individuals passed unnoticed → today > 100 000 inds.

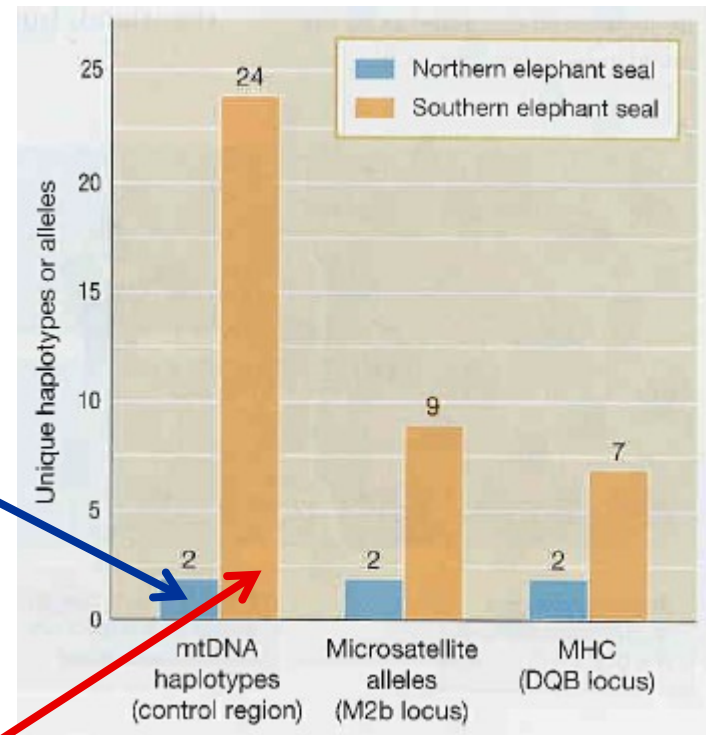
M. Bonnell a R.K. Selander (1974): blood samples of 159 individuals electrophoresis at 21 loci → no variation
likewise Hoelzel et al. (1993), 62 loci



Hoelzel et al. (1999): DNA markers



northern elephant seal
(*Mirounga angustirostris*)



southern elephant seal
(*Mirounga leonina*)

humans

a) Las Salinas (Dominican Republic):

Altagracia Carrasco:

several children with at least 4 men

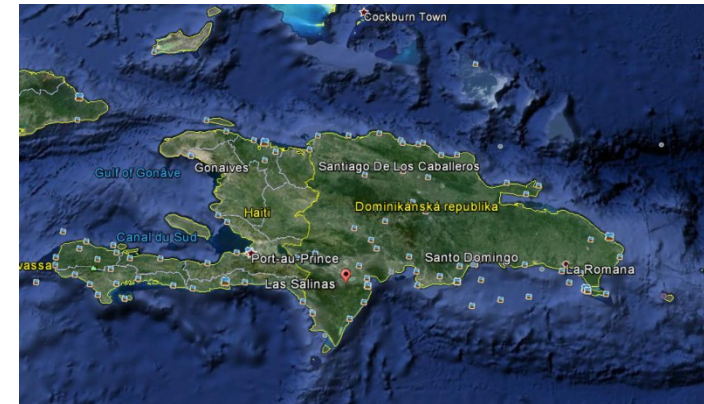
Carrasco heterozygous for substitution $T \rightarrow C$ in 5th exon of the 5- α -reductase 2 gene \Rightarrow TGG (Trp) \rightarrow CGG (Arg) at the 246th position of the protein

the enzyme catalyzes transformation of testosterone to DHT (dihydrotestosterone)

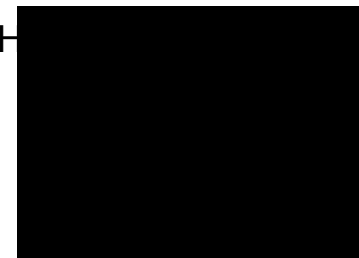
low activity of the mutant enzyme in homozygotes \Rightarrow boys have testes but other traits are female

in puberty testosterone production increases \Rightarrow transformation to men

in Salinas high frequency of the mutation \Rightarrow the word *guevedoces* (= „penis in 12“)



DH



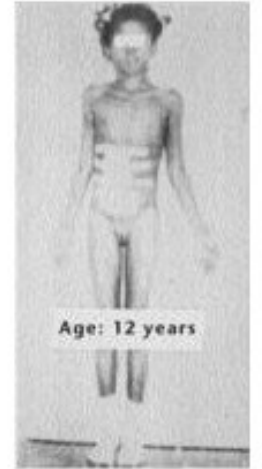




Age: 4 years



Age: 8 years



Age: 12 years



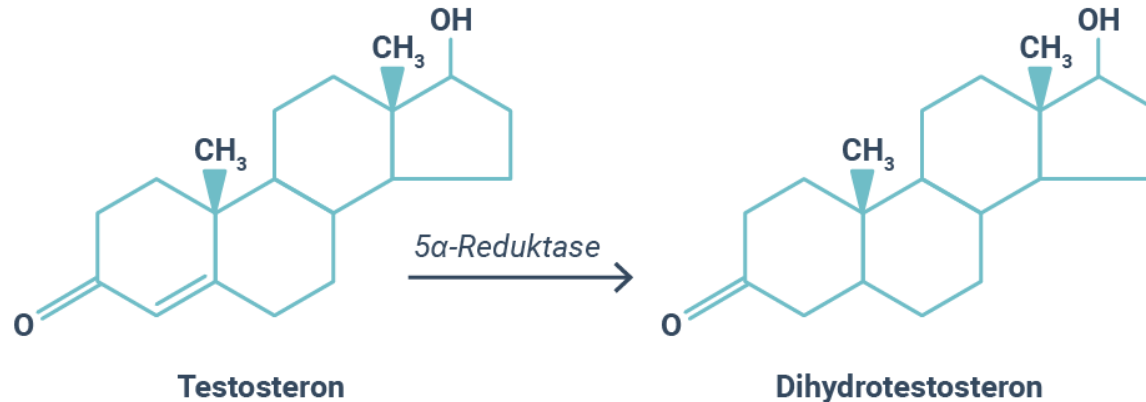
Age: 18 months



Age: 19 years



heterozygote for mutation of 5- α -reductase 2, catalyzing change of testosterone to dihydrotestosterone (DHT)



low activity of mutant enzyme \rightarrow development of female traits in homozygotes, hidden testes

increased production of testosterone during puberty \Rightarrow transformation into male

founder effect: Altagracia Carrasco \rightarrow multiple offspring with min. 4 men in small population \Rightarrow now high incidence of the phenomenon

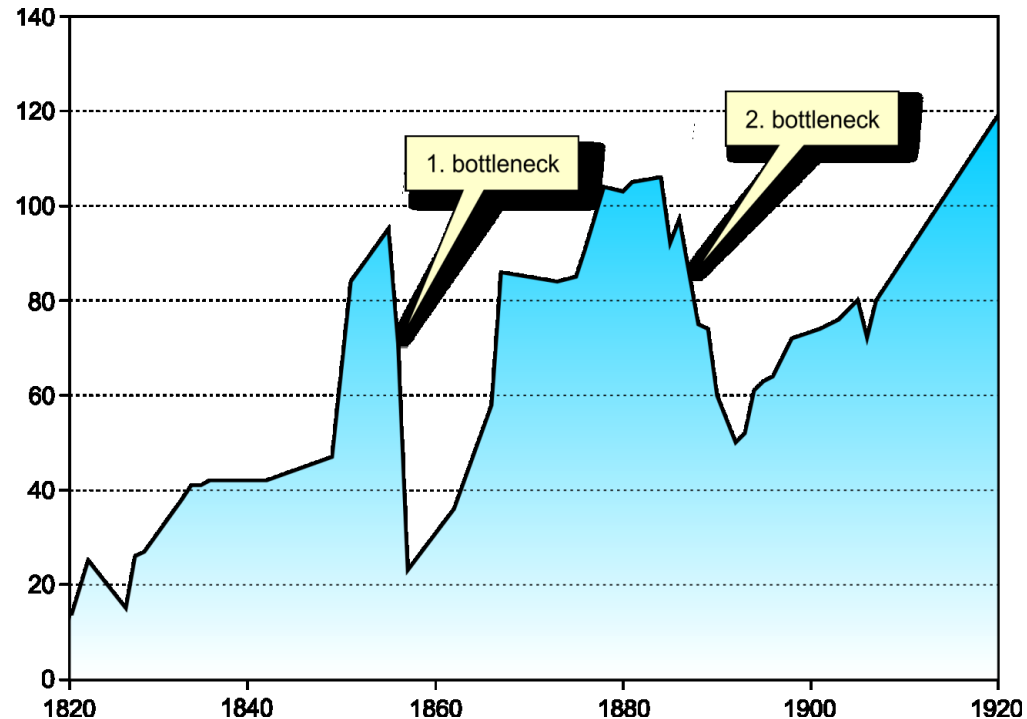
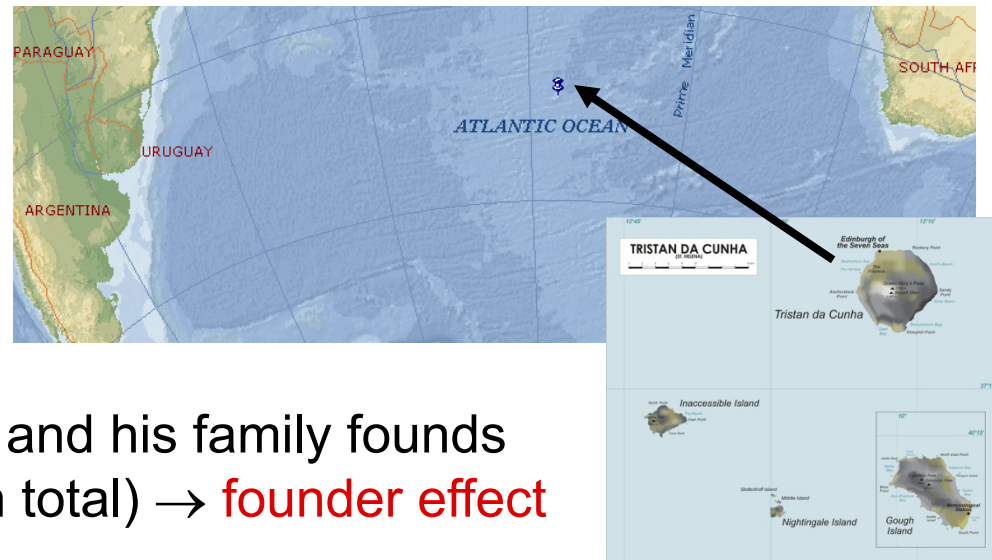
Tristan da Cunha:

1816 military garrison

1817 garrison withdrawn;

Scottish corporal **William Glass** and his family founds a small colony (20 individuals in total) → **founder effect**

during 80 years 2 strong bottlenecks

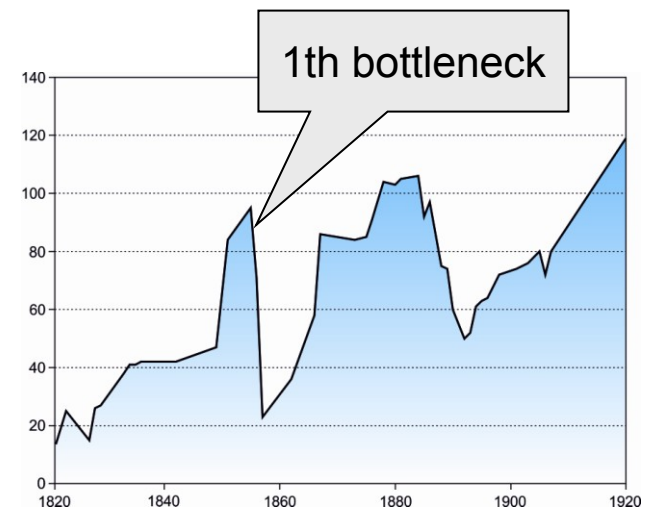
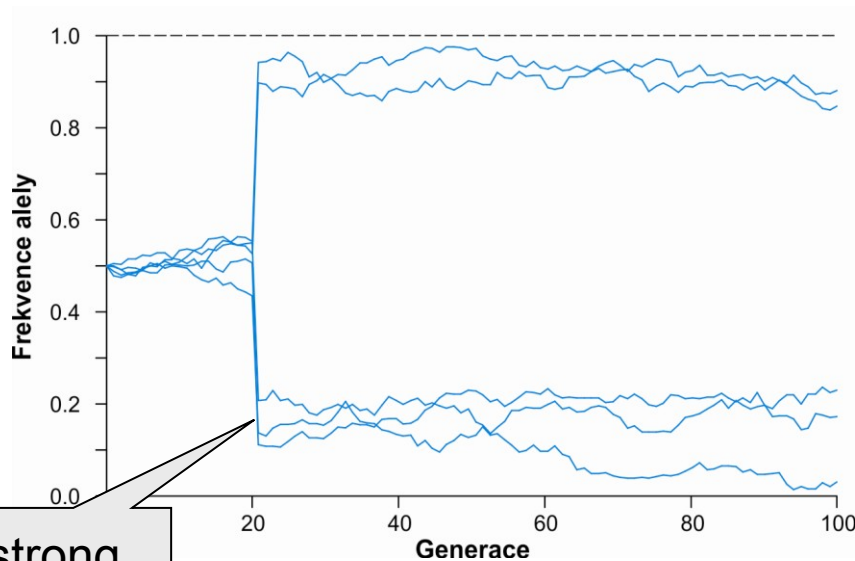


1851: a missionary arrival

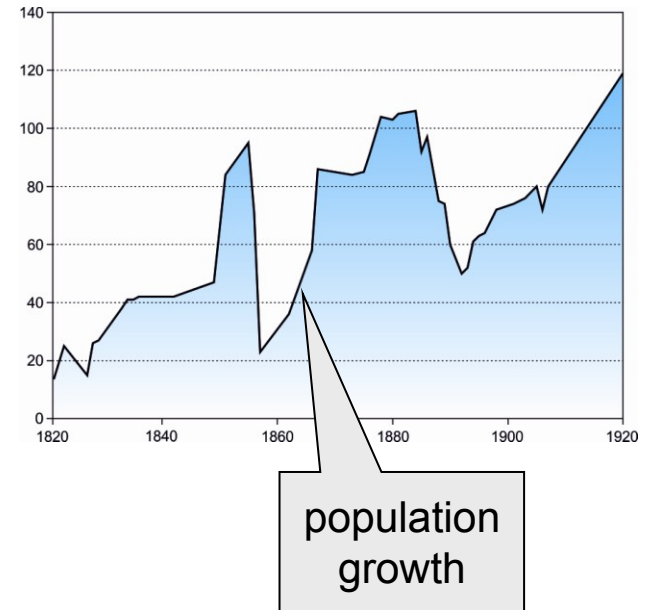
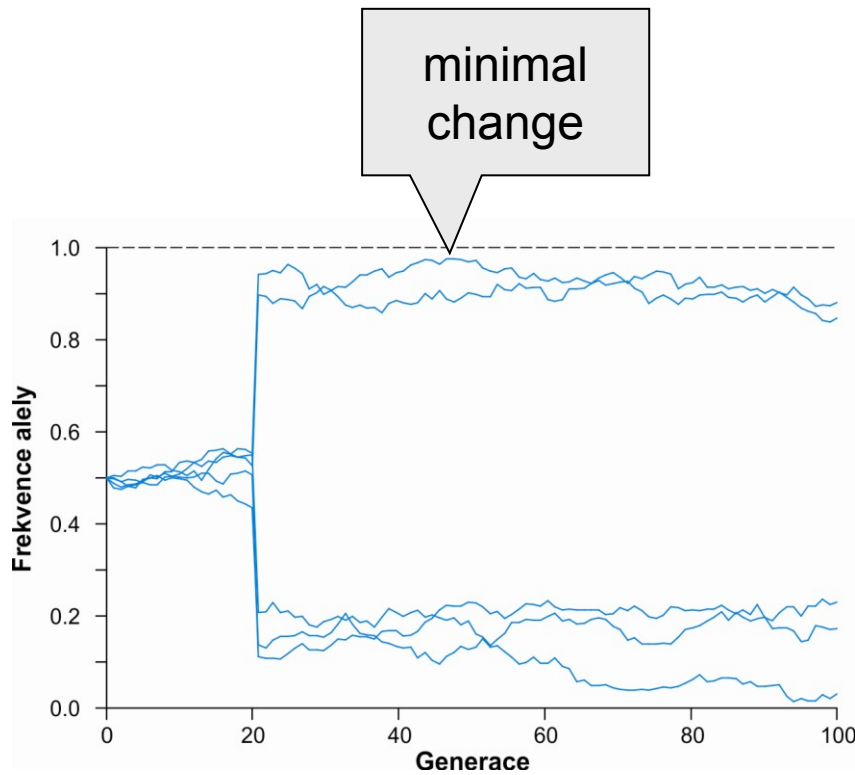
1853: death of W. Glass

1856: departure of 25 Glass's descendants to America, departure of other 45 people with the missionary

⇒ 103 inds. (1855) → 33 (1857) ... **1st bottleneck**

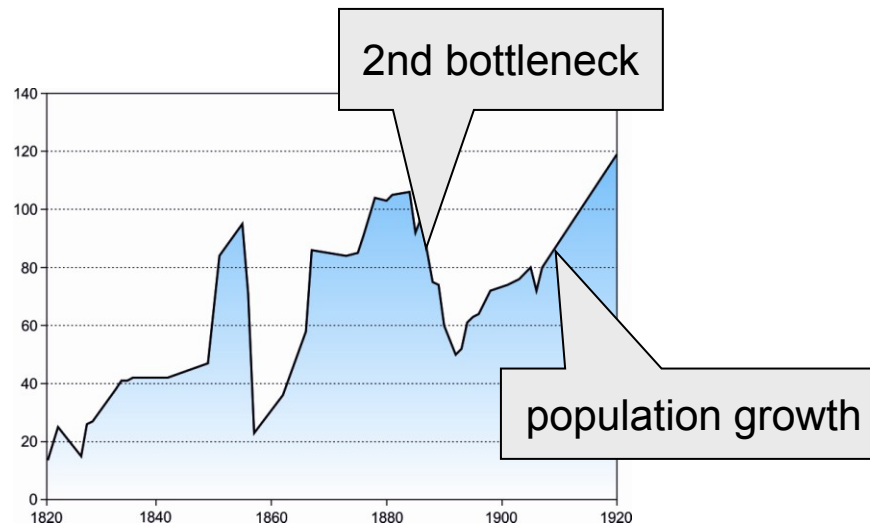


1857–1884: population growth \Rightarrow conservation of changes caused by previous bottleneck \rightarrow less changes during 27 years than during 2 years 1855–1857

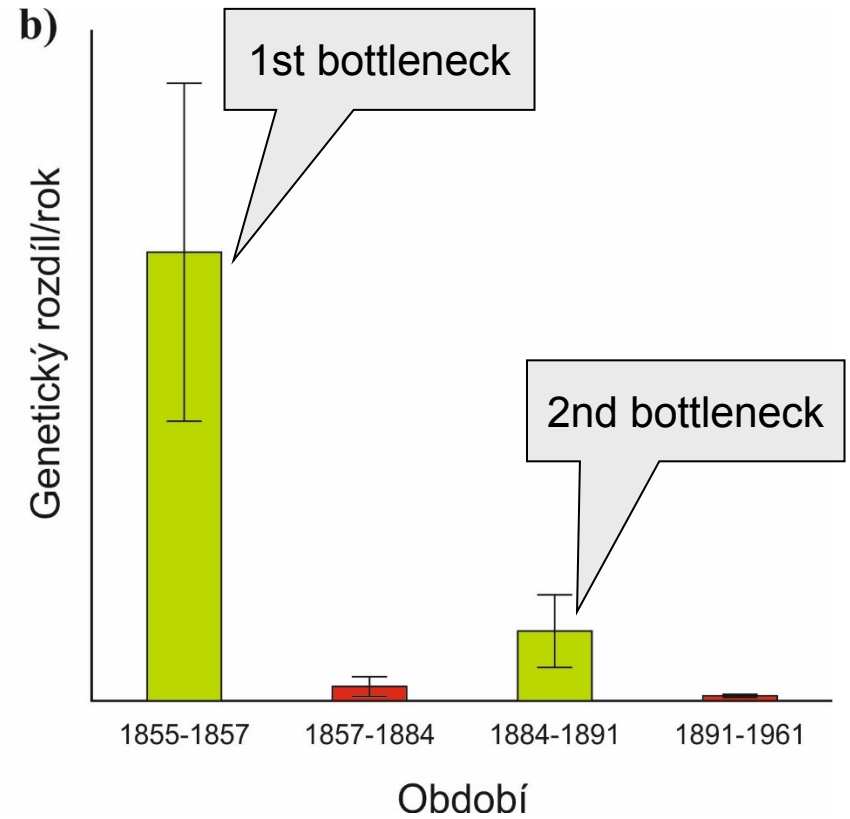
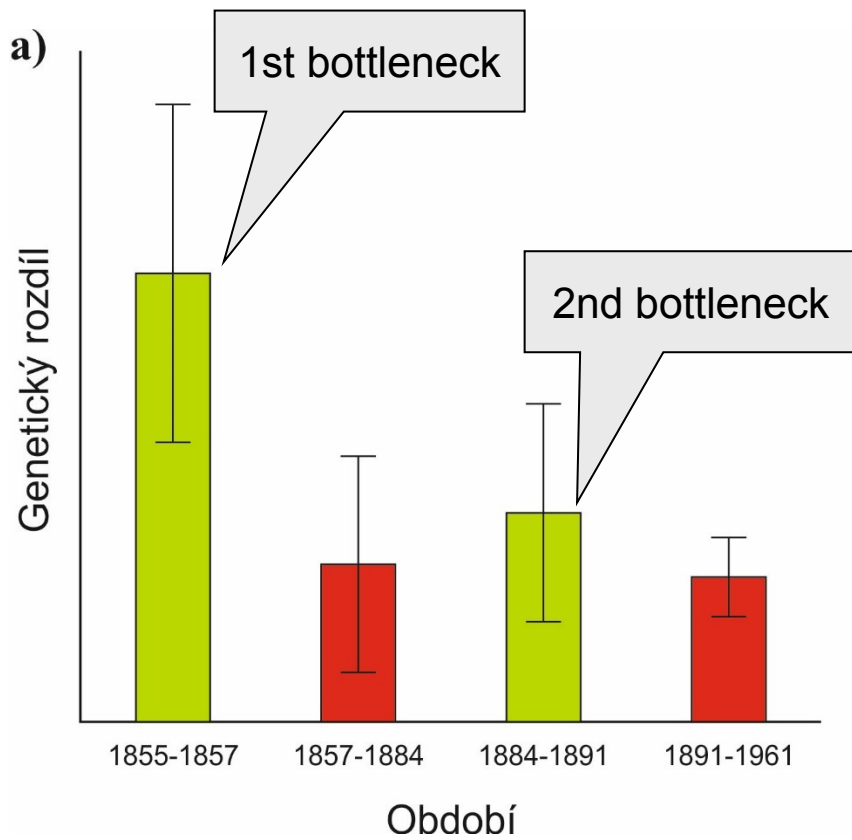


1884–1891: drowning of 15 men, only 4 adult remains, of which 2 very old („Island of Widows“) → departure of many widows with their children

⇒ 106 inds. (1884) → 59 (1891) ... **2nd bottleneck**

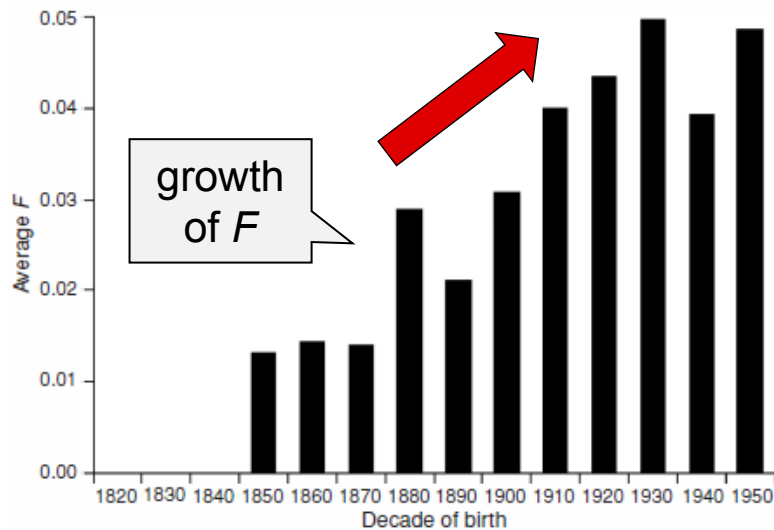


again, the following growth has „frozen“ the changes



Genetic changes during population growth lower than during bottlenecks

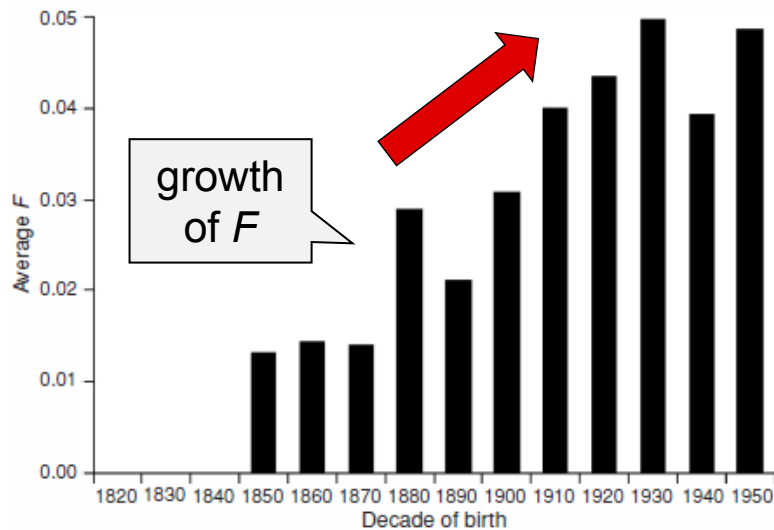
Inbreeding on Tristan da Cunha:



Despite the outbreeding strategy (choice of the least related partner), ie. $F_{IS} < 0$, the level of autozygosity increased



Inbreeding on Tristan da Cunha:



Despite the outbreeding strategy (choice of the least related partner), ie. $F_{IS} < 0$, the level of autozygosity increased

Table 3.3. First Eight Marriages between Biological Relatives on Tristan da Cunha Showing Date of Marriage, Number of Available Women of Marriageable Age,^a and Number of Available Women Not Related to Groom

Marriage between Relatives	Date of marriage	Number of available women	Number of non relatives
1	1854	7	3
2	1856	9	2
3	1871	1	0
4	1876	1	0
5	1884	7	1
6	1888	8	0
7	1893	3	0
8	1898	1	0

no unrelated woman available!

^a Sixteen years and over, single, and not a sister of the groom.

RELATION BETWEEN DRIFT AND GENE FLOW

Generace 0

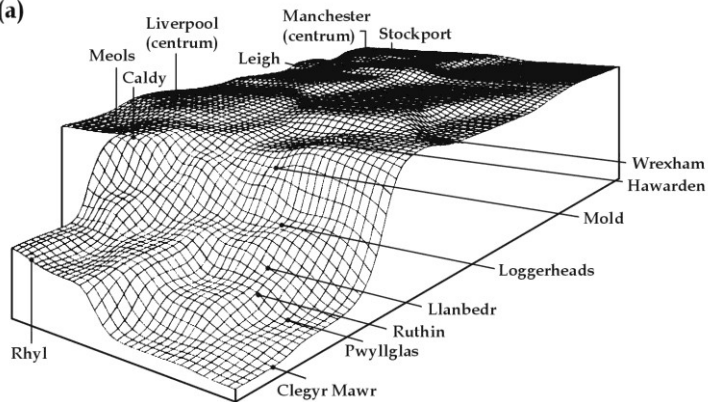
50

110

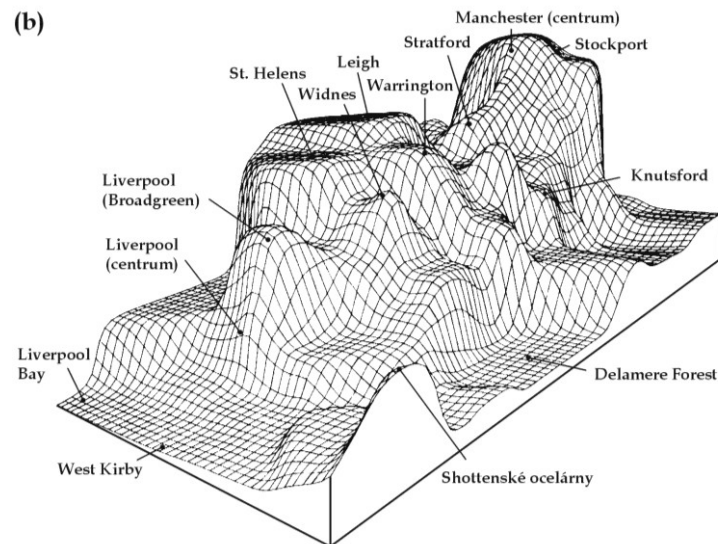
higher
gene flow

lower
gene flow

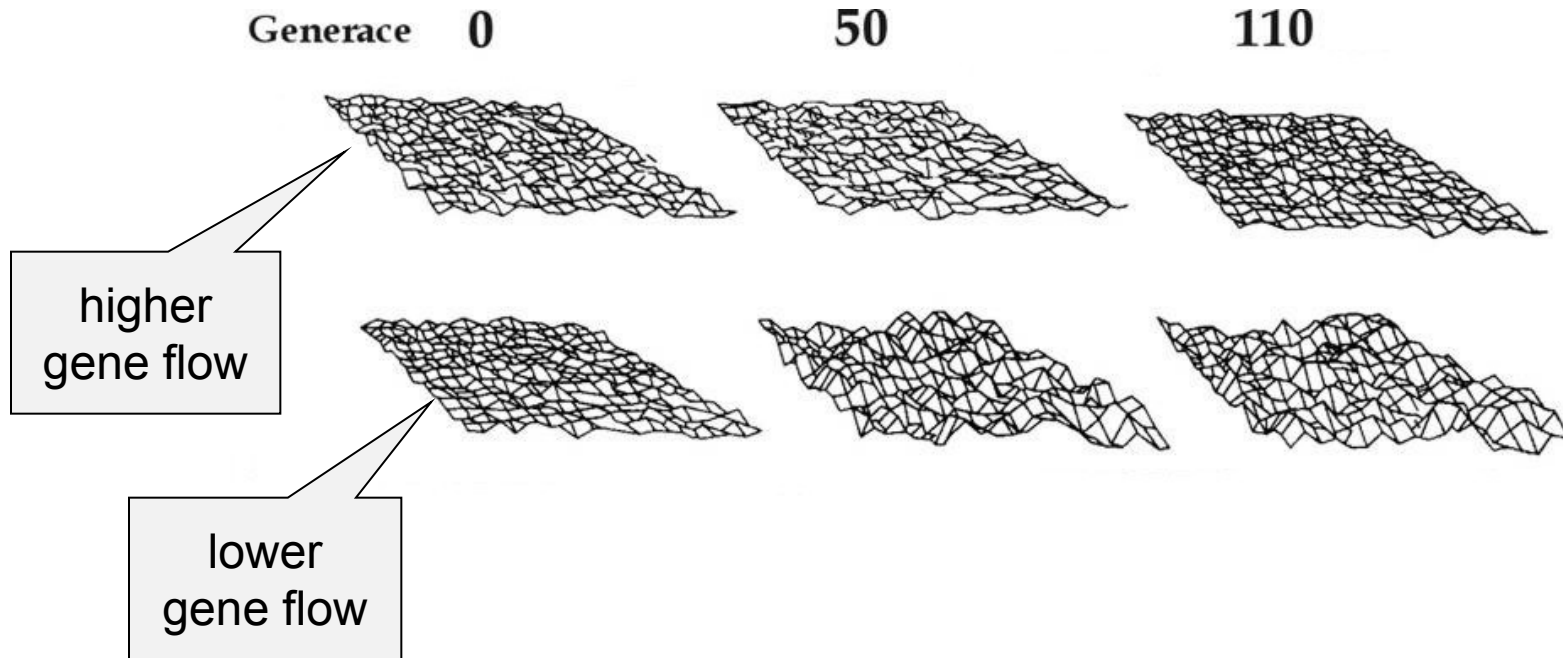
(a)



(b)



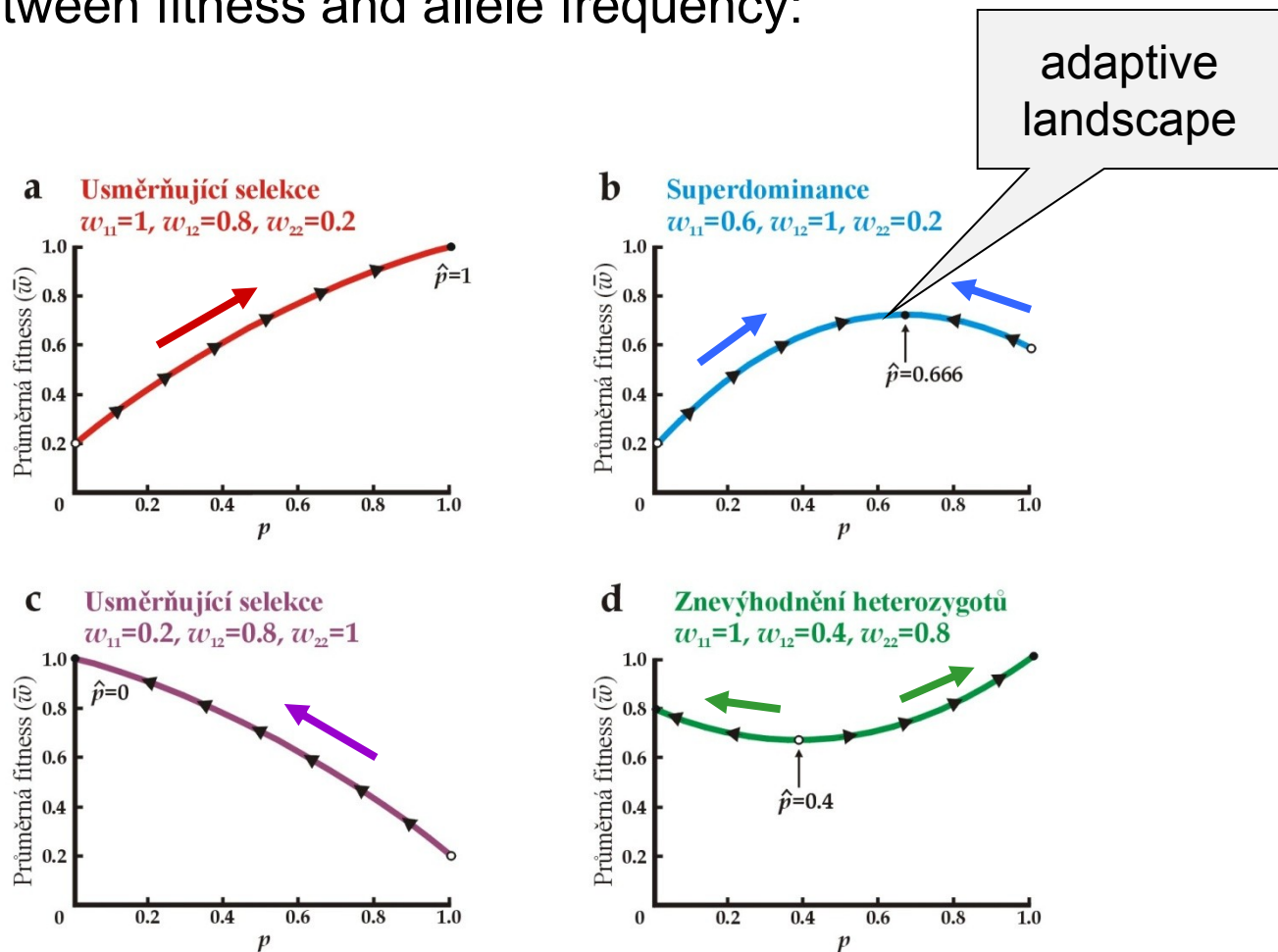
RELATION BETWEEN DRIFT AND GENE FLOW

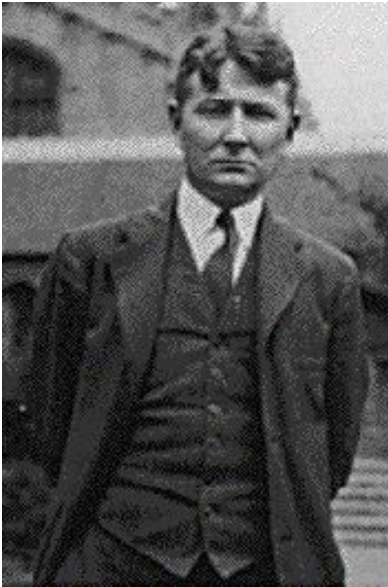


Gene flow and drift have opposite effects:
drift increases divergence among demes × migration „homogenizes“ demes

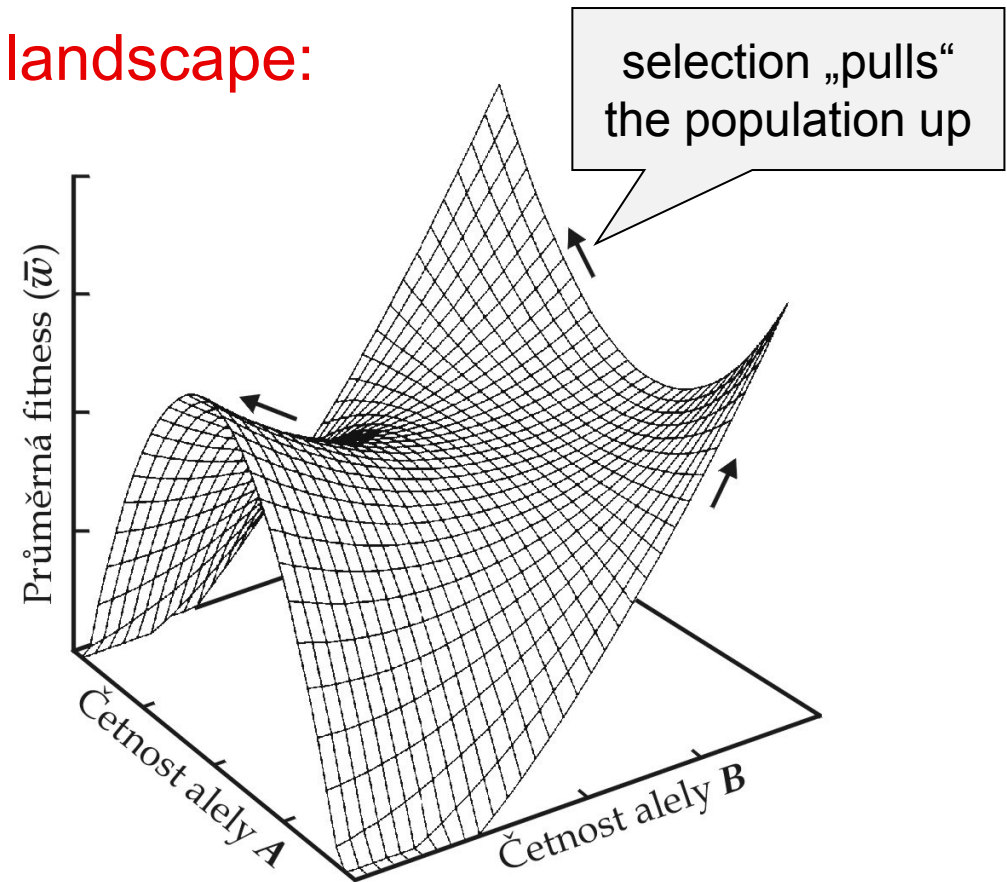
RELATION BETWEEN DRIFT AND SELECTION

relation between fitness and allele frequency:





Adaptive landscape:



Adaptive landscape has been defined in different ways, most often as a field of average allele frequencies

number of dimensions = number of loci + fitness

surface of such landscape is continuous

Shifting balance theory (SBT)

Assumptions:

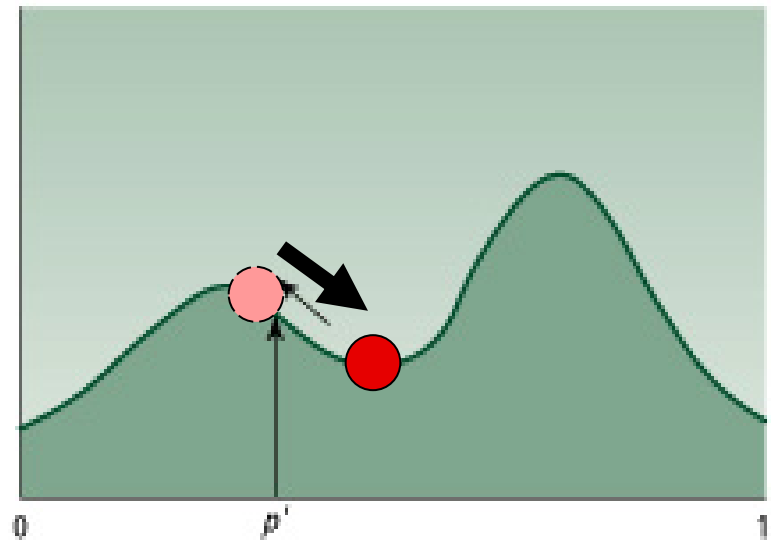
environment changes \Rightarrow populations in constant change

mutations \Rightarrow new dimensions, new ways upwards

small populations (drift) \Rightarrow possibility to move down to adaptive valleys

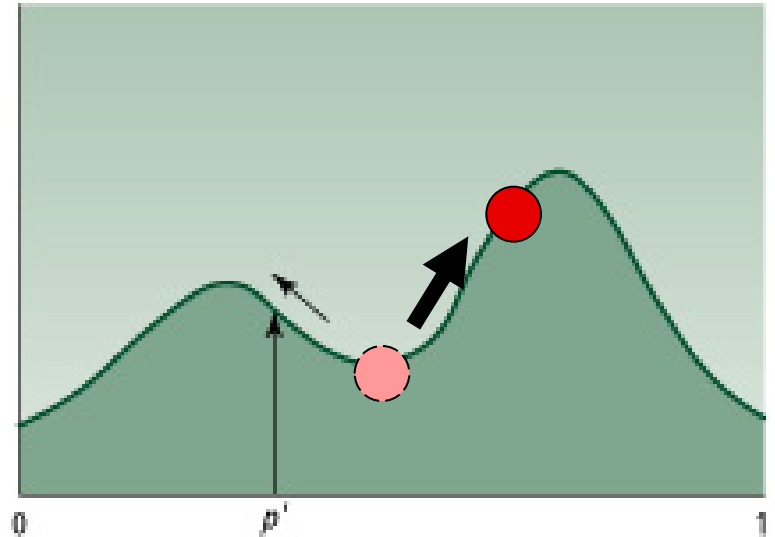
3 phases of SBT:

1. contemporary fitness reduction of a local population due to drift \rightarrow chance to approach the area of attraction of a higher peak



3 phases of SBT:

2. intrademic selection → „pulling“ of the population towards a new peak



3. interdemic selection → spread of the deme's members at the higher peak to surrounding demes

The whole process seen as shifting of the balance between drift, intrademic, and interdemic selection

2 views on evolution in populations:



S. Wright



R.A. Fisher

small local populations

combination of selection, drift and migration

epistasis, pleiotropy,
dependence of allele effects on context

speciation as a byproduct of local adaptations in epistatic systems

large panmictic populations

mutation and selection

additive effects of genes,
allele effects independent of context

disruptive or locally divergent selection