ORIGIN OF GENETIC VARIATION



Consequence of the H-W principle:

if the assumptions of the H-W population hold true, polymorphism can be maintained solely by random mating and Mendelian inheritance





BUT!

real populations usually differ from the model:

population size finite
mating may be nonrandom
migration
selection
emergence of new alleles by mutation

MAIN MICROEVOLUTIONARY MECHANISMS:

mutation (incl. transposition)

recombination

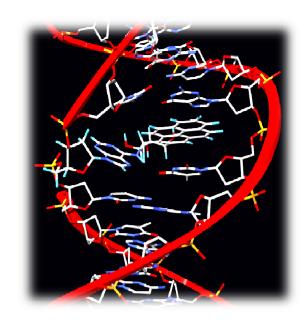
migration (gene flow)

nonrandom mating

natural selection

random genetic drift (incl. bottleneck, founder effect)

(molecular drive)



MUTATION



spontaneous × induced

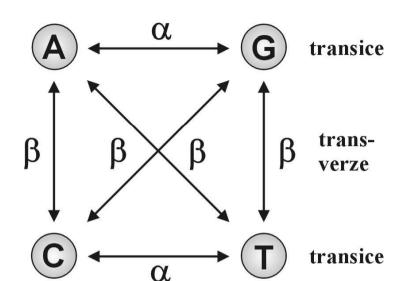
in germ cells × somatic

according to their deleterious/beneficial effect:

beneficial (positive) deleterious (lethal, negative) neutral

According to effect

point (gene) chromosomal genome



Point mutations:

substitutions (transitions, transversions)



synonymous

nonsynonymous

$$\begin{array}{ccc} \mathsf{GTC} \to \mathsf{GTA} & \mathsf{GTC} \to \mathsf{TTC} \\ \mathsf{Val} & \to \mathsf{Val} & \mathsf{Val} \to \mathsf{Phe} \\ & \mathsf{AAG} \to \mathsf{TAG} \\ & \mathsf{Lys} & \to \mathit{ochre} \ (\mathsf{stop}) \end{array}$$

missense nonsense insertions ACGGT → ACAGGT delections ACGGT → AGGT } indels → shift of reading frame

back mutations: generally 10-times lower frequencies

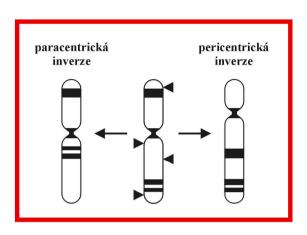
recurrent mutations → mutation pressure:

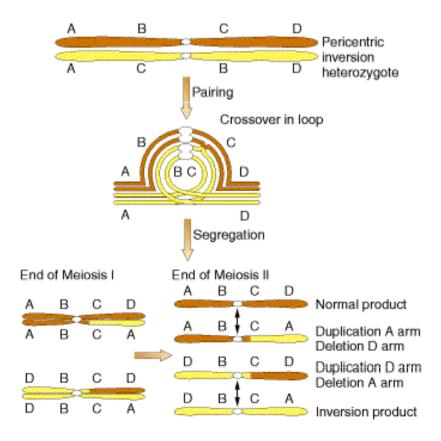
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eg. when allele frequency A = 0.5; 2N = 2000: after 1st generation \rightarrow N = 1001 \Rightarrow increase to 0,5005 after 100 generations \rightarrow 0.55 \dots
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⇒ change of allele frequencies by mutations very slow

Chromosomal mutations (chr. rearrangements)

inversions pericentric paracentric





translocations

fusions and dissociations

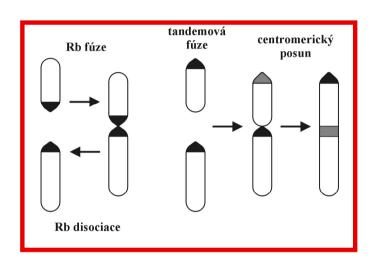
(Robertsonian translocations)



house mouse

MA AA AA

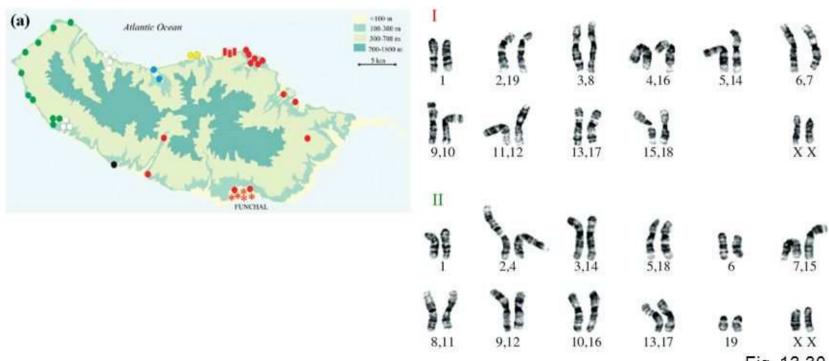
AX XX XX XX XX XX XX XX XX AX AA



Rapid chromosomal evolution in house mice on the island of Madeira

One population of mice introduced to island in 1400s

Two populations evolved different sets of Robertsonian translocations, hybrid offspring are sterile



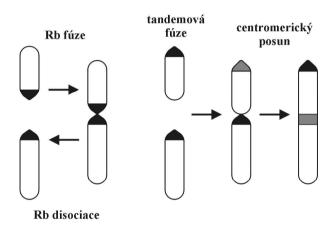
translocations

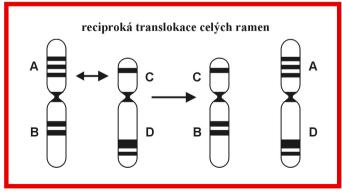
fusions and dissociations

whole-arm reciprocal translocations (WART)

house mouse



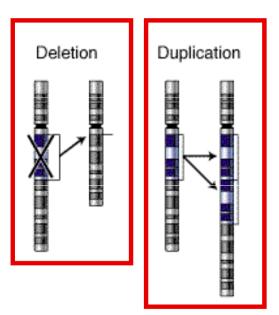


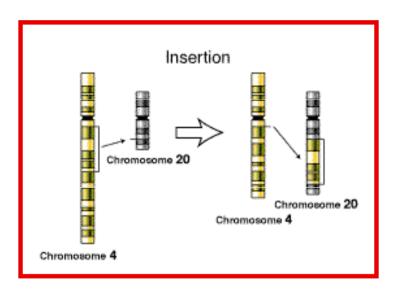


deletions

duplications

insertions





Genome mutations

-somies (monosomies, trisomies)

mostly incompatible with life

monosomies: the only viable = X0 (Turner syndrom)

trisomies: imbalance in gene dosage (increased expression of the trisomic pair)

viable trisomies: XXY, XXX, XYY, Patau syndrom (chr. 13), Edwards s. (chr. 18), Down s. (chr. 21)

-ploidies (polyploidy)

especially plants

in animals less frequent (invertebrates, fishes, amphibians)

during the vertebrate evolution 2 rounds of whole genome duplications (2R-hypothesis)

polyploid individuals usually bigger (increased cell volume)

odd multiples of the genome \rightarrow problems in meiosis \Rightarrow reproductive barrier (not always – eg. triploid frogs)

autopolyploidy: combination of two identical genomes

fusion of cells endoreplication abortive cell cycle

allopolyploidy: combination of two different genomes

fusion of diploid gametes polyspermy

Randomness and mutation rate (µ)

mutation effects random, position and rate nonrandom

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transitions > transversions
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mutation "hotspots": CpG in animals (methylated $C \rightarrow T$); TpT in Procaryota "SOS reactions" in Bacteria, minisatellites (VNTR), microsatellites (STR)

mtDNA > nuclear DNA

sex chromosomes > autosomes

influence of proximity of the replication start, centromeres, telomeres, repetitive sequences, intensity of transcriptions

cold-blooded animals: > temperature \Rightarrow > μ

RNA viruses (HIV)

parasites

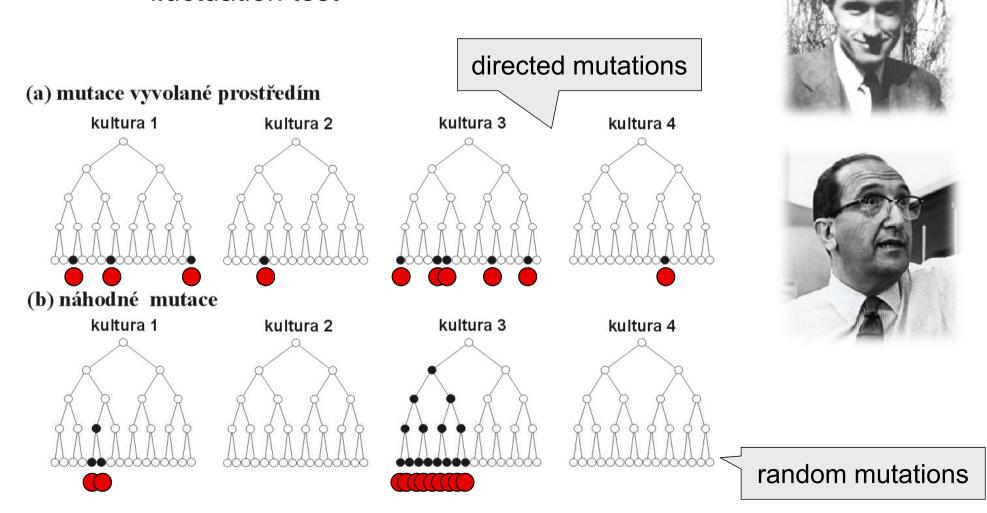
antigens, immunoglobulins

> μ of somatic mutations

males > females: humans 6x, rodents, fox: 2x ... more cell divisions in germ cells

Adaptive (directed) mutations?

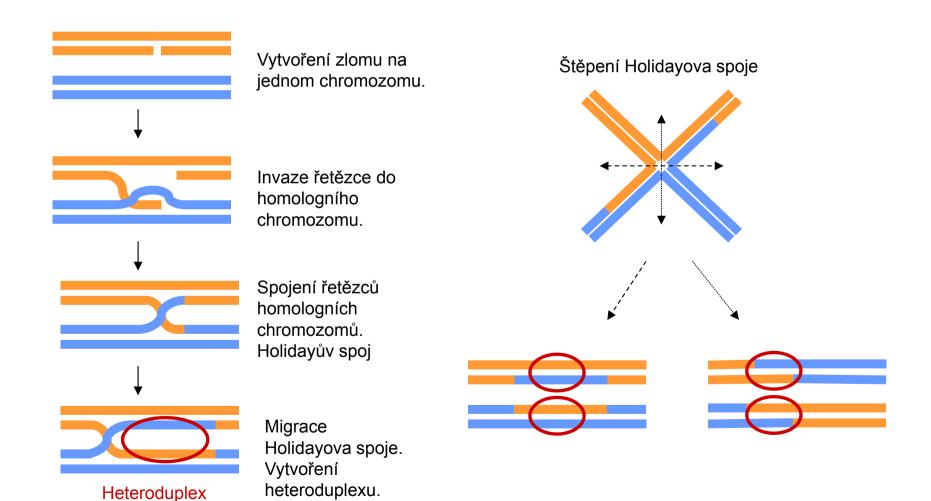
Max Delbrück, Salvador Luria (1943): fluctuation test



RECOMBINATIONS

mutations → new alleles

recombinations → new genotypes (exception = intragenomic recombination)



in many organisms crossing-over important for right meiosis (at least 1 c-o per chromosome, otherwise aneuploidies)

women with $> c-o \rightarrow > children$

children of older women \rightarrow > recombinations

differences in various parts of chromosome (near centromeres and telomeres etc., differences among organisms)

small chromosomes > recombination frequencies

recombination "hotspots":

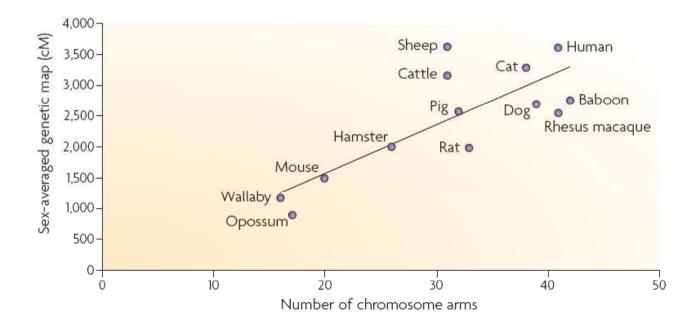
humans ~25 000 absent in *Drosophila* and *Caenorhabditis elegans* frequent appearance and disappearance demise of 1 point often compensated by increased aktivity of a neighbour point

differences in recombination rate between sexes:

- Haldane-Huxley rule: if one sex doesn't recombine, it is the heterogametic sex
- if both sexes recombine, mostly in females > recombinations (man 1,7x, mouse 1,3x)

differences between species:

- species with more small chromosomes \rightarrow more recombinations than species with less large chromosomes
- correlation with the number of arms: more recombinations in karyotypes with large numbers of chrom. arms (at least 1 c-o/arm to avoid aneuploidies?)

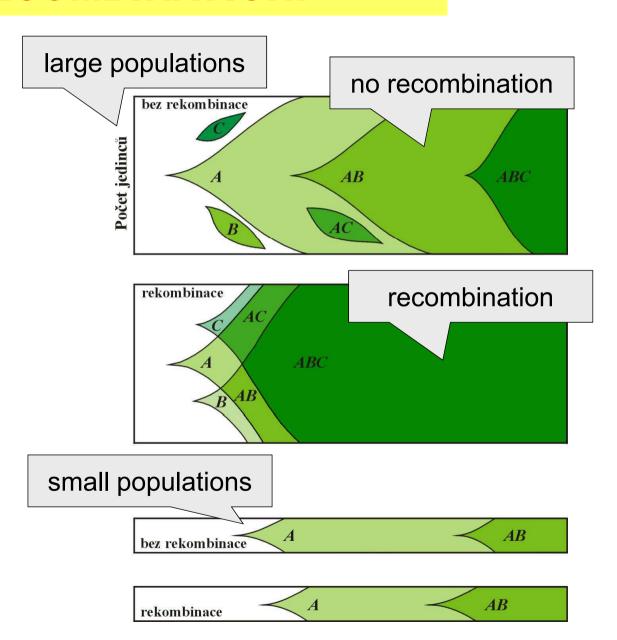


EVOLUTIONARY CONSEQUENCES OF RECOMBINATION:

Recombination and polymorphism:

absence of recombination

⇒ linkage disequilibrium



EVOLUTIONARY CONSEQUENCES OF RECOMBINATION:

Recombination and polymorphism:

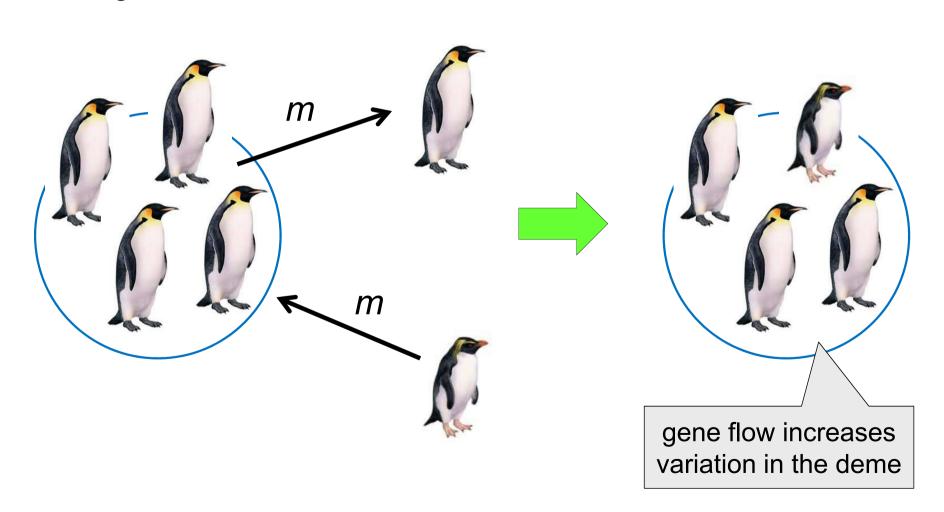
positive selection: selective sweep hitchhiking (draft)
more frequent appearance of rare alleles

negative selection: background selection

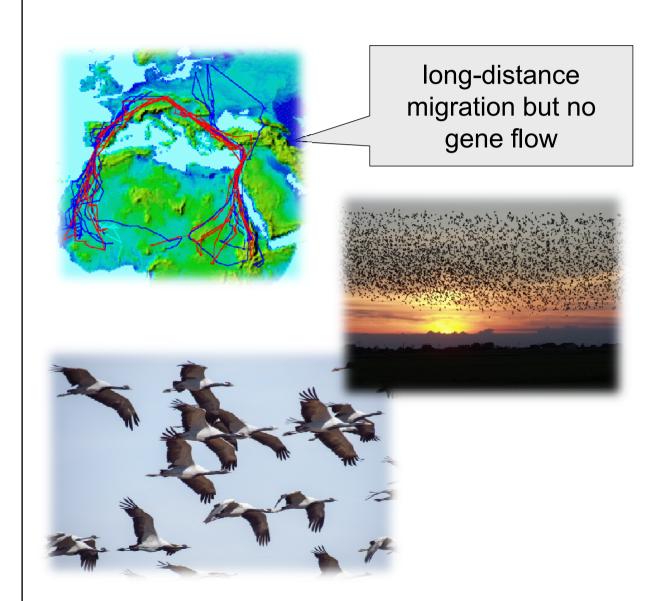
 \rightarrow loss of polymorphism

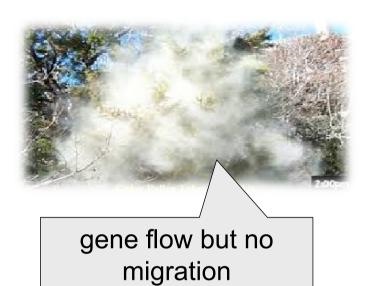
MIGRATION (GENE FLOW)

Migration rate, *m* = proportion of gene copies appearing in the population by immigration from other populations in the given generation



MIGRATION (GENE FLOW)







METHODS OF GENE FLOW ESTIMATION:

1. direct

capture-mark-recapture (CMR)

finger clipping, special dyes, tattooing, tags, rings, collars,

genetic marking













1. Direct methods

remote tracking – telemetry

transmitters, anntenas; GPS systems

... more expensive, time consuming









Risk of underestimation of gene flow!!

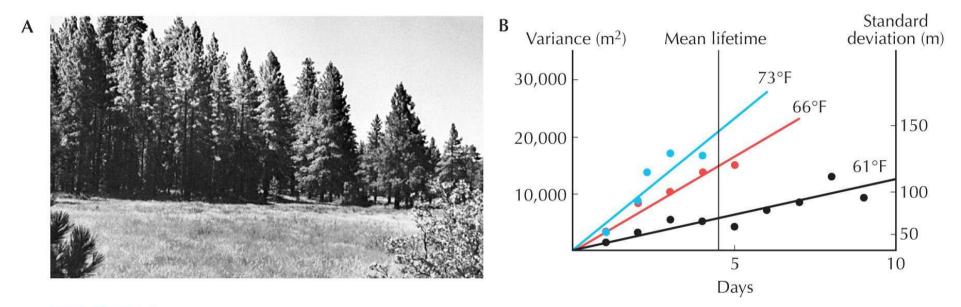


FIGURE 16.4. Dobzhansky and Wright (1943) measured the rate of dispersal of *Drosophila pseudoobscura* by releasing marked flies at sites in the Sierra Nevada, California (*A*). Over the following days, flies were caught in a series of traps. The graphs (*B*) show how the variance of the distribution of marked flies increased over time. The three sets of points show results from experiments at different times during the summer: Rates of movement increase strongly with temperature. The rate of diffusion of genes is estimated by assuming a mean lifetime of 4.5 days (*vertical line*).

2. Indirect methods

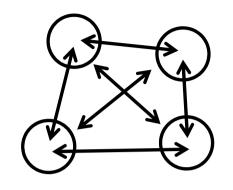
molecular markers

gene flow models

maximum likelihood and Bayesian programs

dispersal: distance between parents and offspring

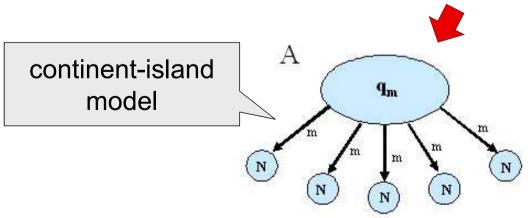
A) Island model

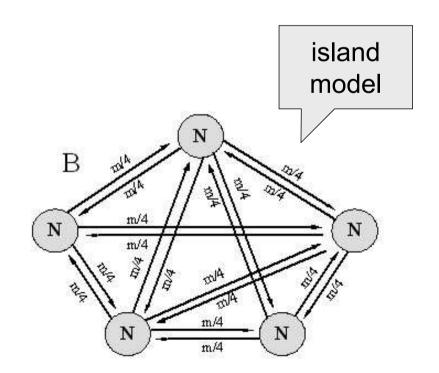




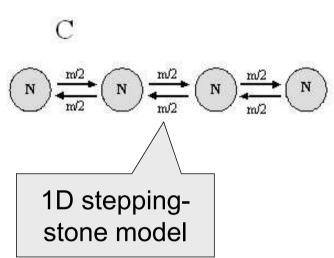
S. Wright (F-statistics): $F_{ST} = 1/(4Nm + 1) \Rightarrow Nm = (1/F_{ST} - 1)/4$... Nm = number of migrants per generation

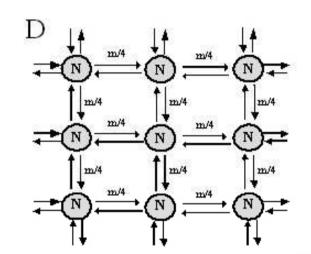
Island model can be also asymmetric:





B) Isolation by distance models discontinuous = stepping stone model





2D steppingstone model



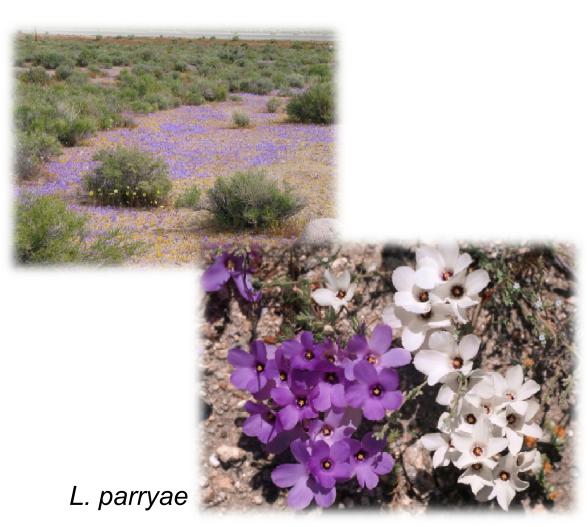


B) Isolation by distance models continuous

Linanthus parryae (Polemoniaceae), Mojave Desert (California)
T. Dobzhansky, Sewall Wright







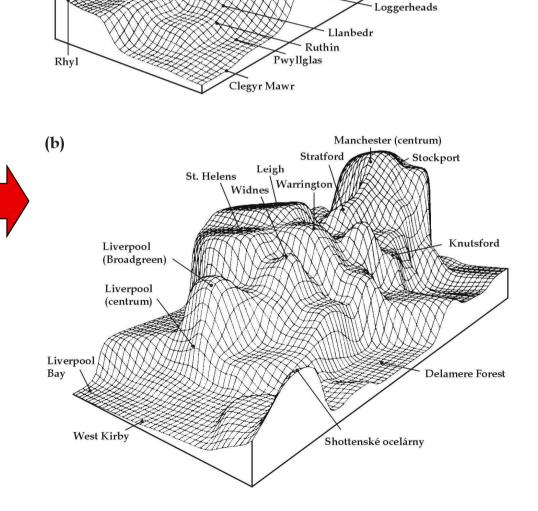
Gene flow consequences:

(a)

genetic homogenization of subpopulations, preventing their genetic divergence

in many species migration severely reduced

Eg.: melanic forms of moths in England



Manchester

(centrum) 7 Stockport

Wrexham Hawarden

Mold

Liverpool

(centrum)

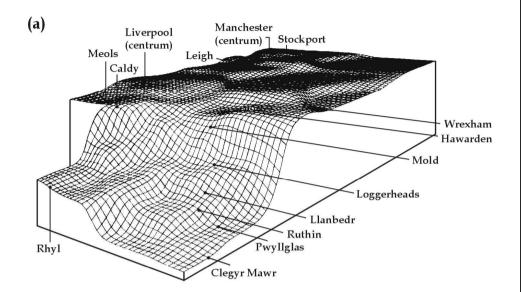


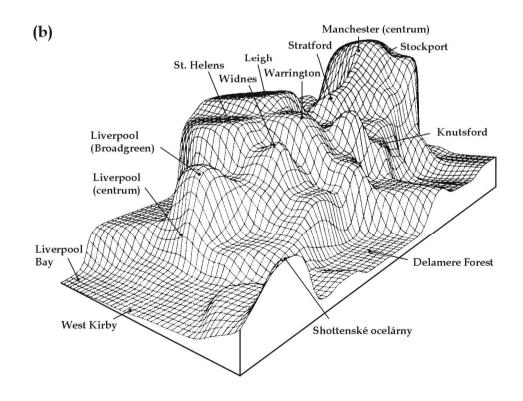


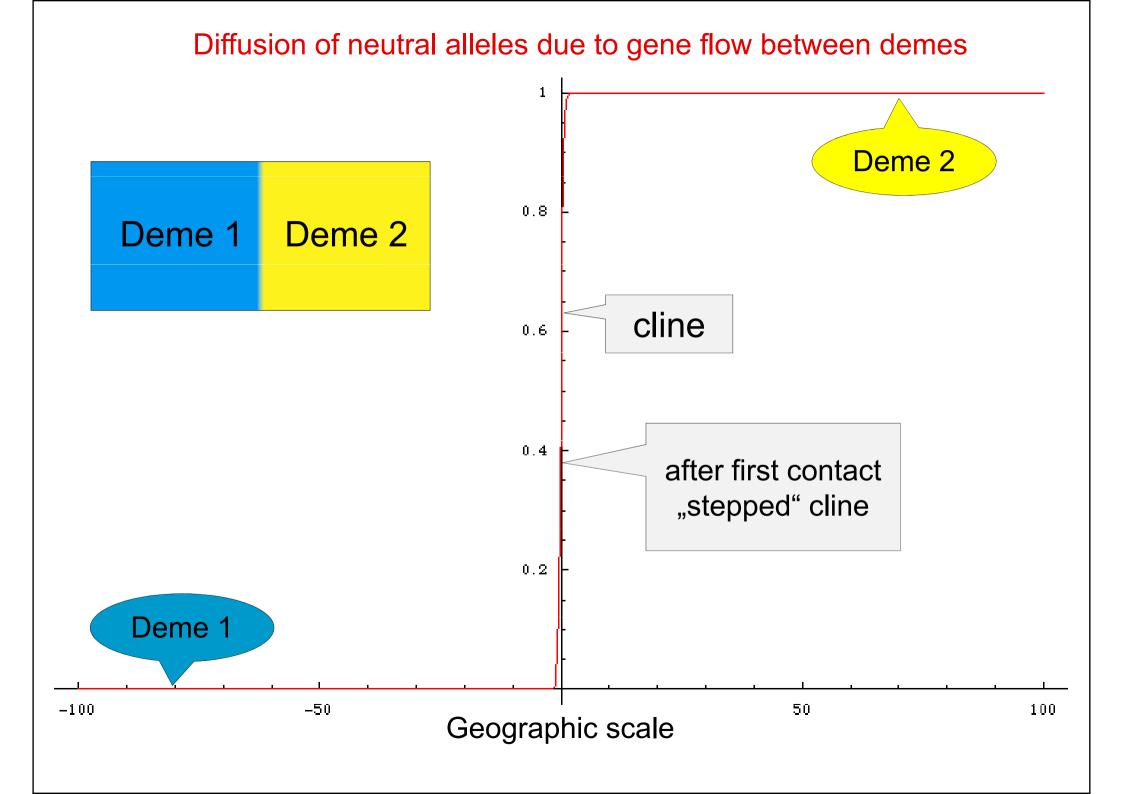
peppered moth (Biston betularia)

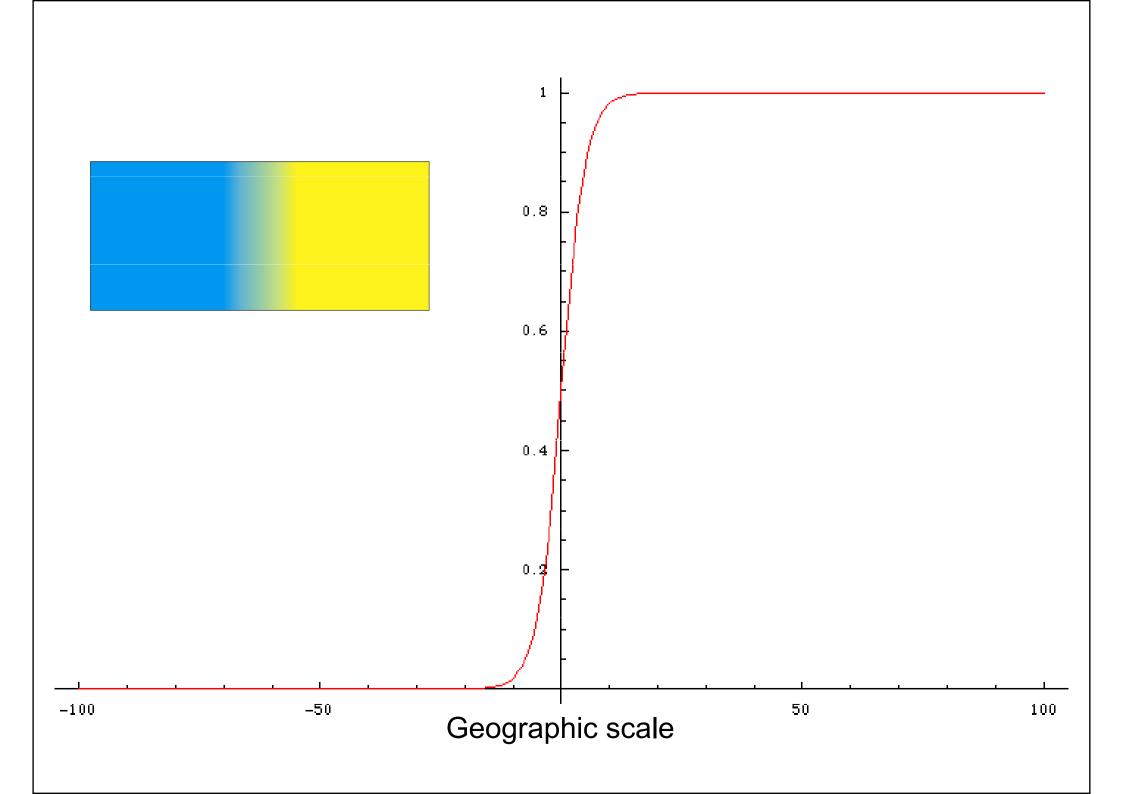


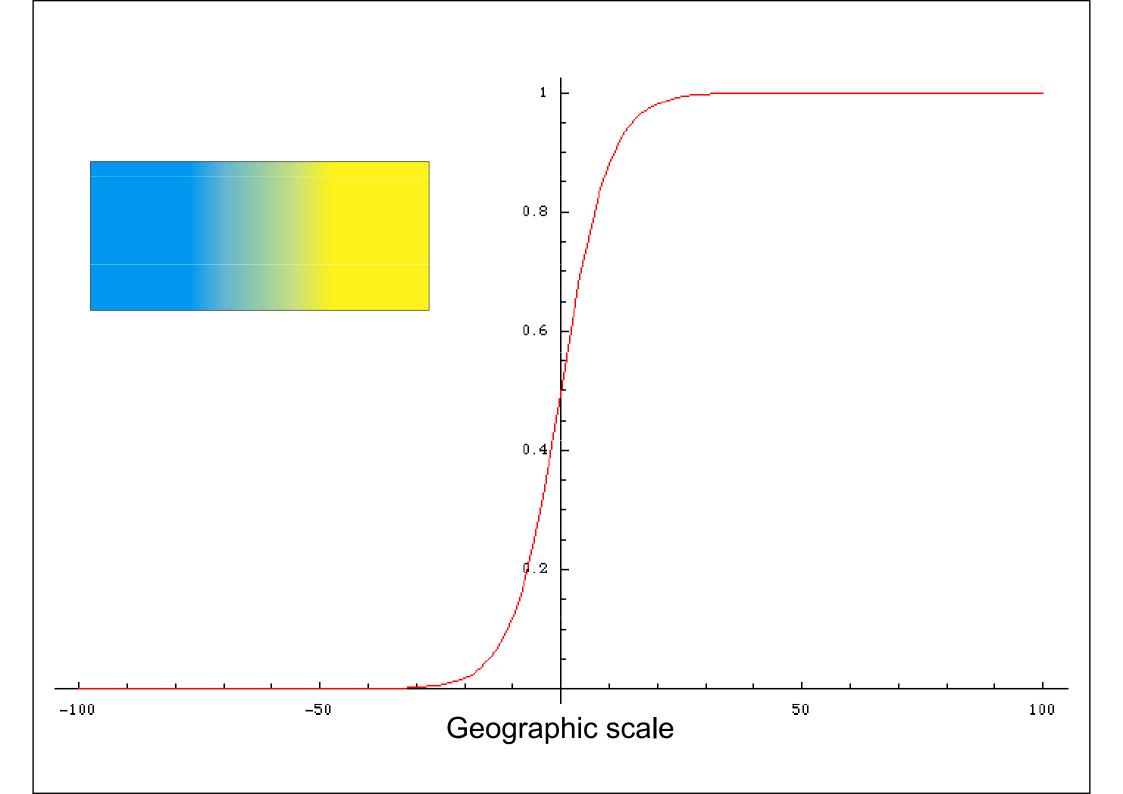
scalloped hazel (Odontoptera [Gonodontis] bidentata)

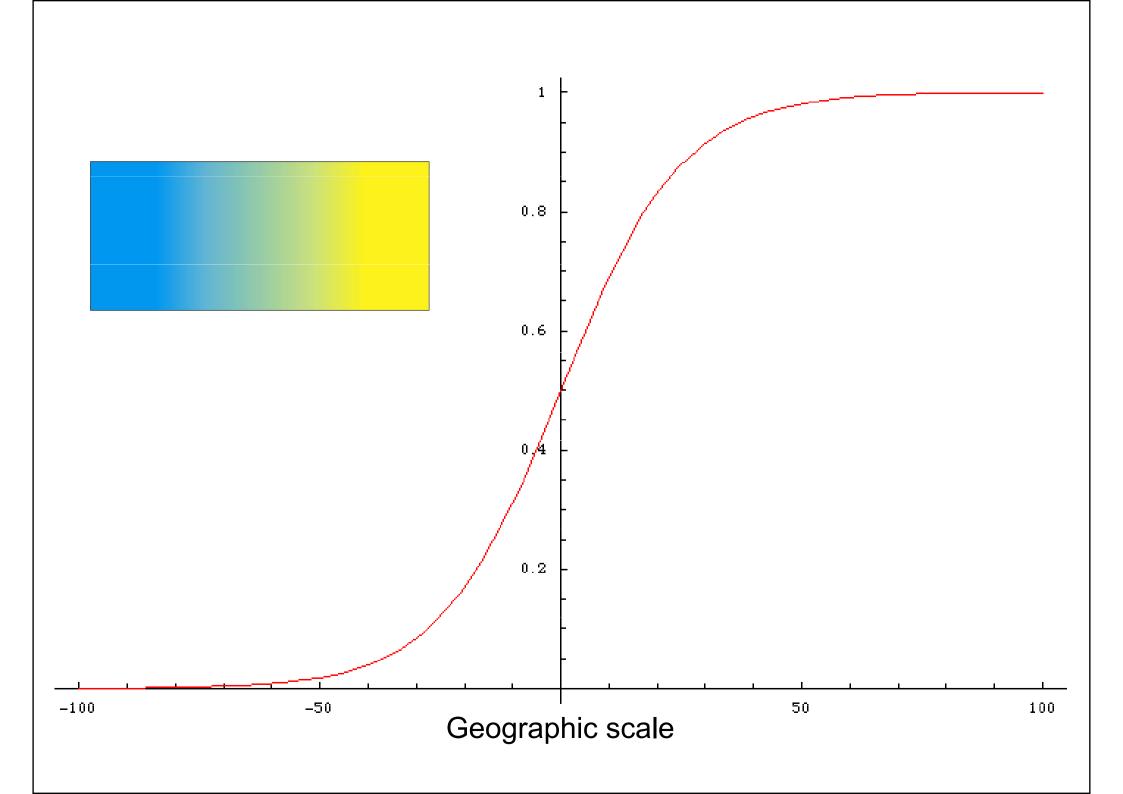


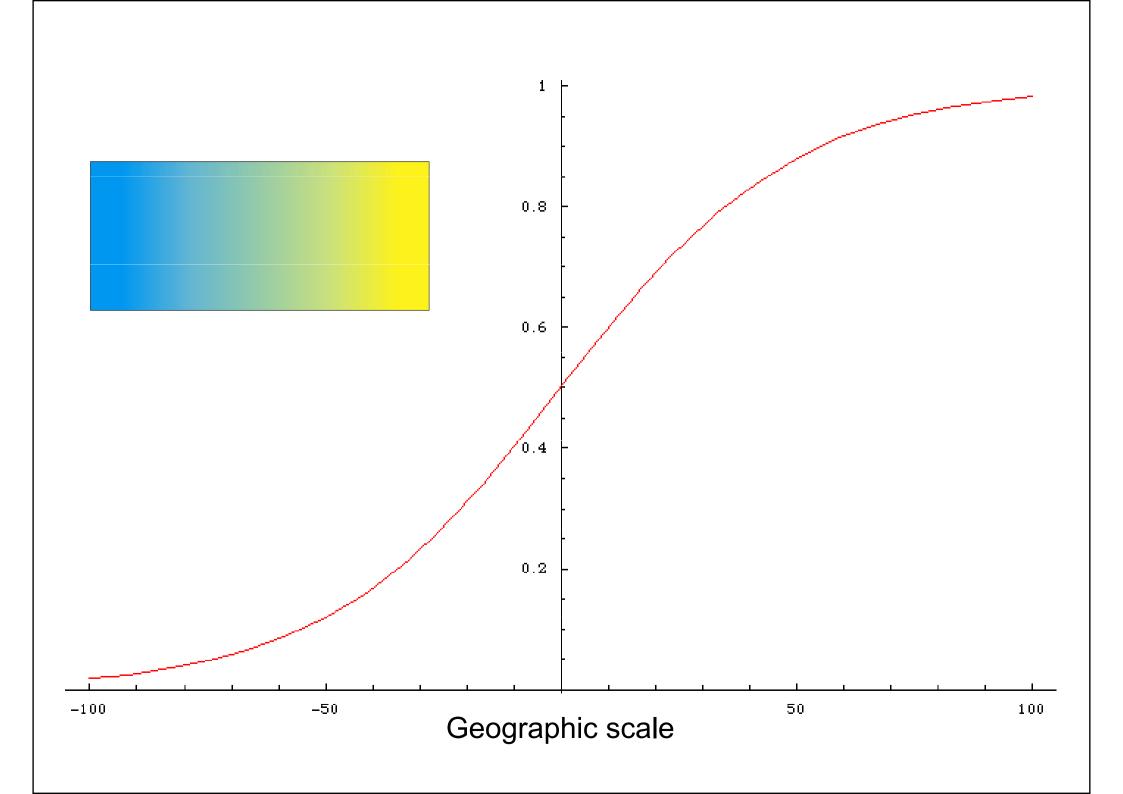




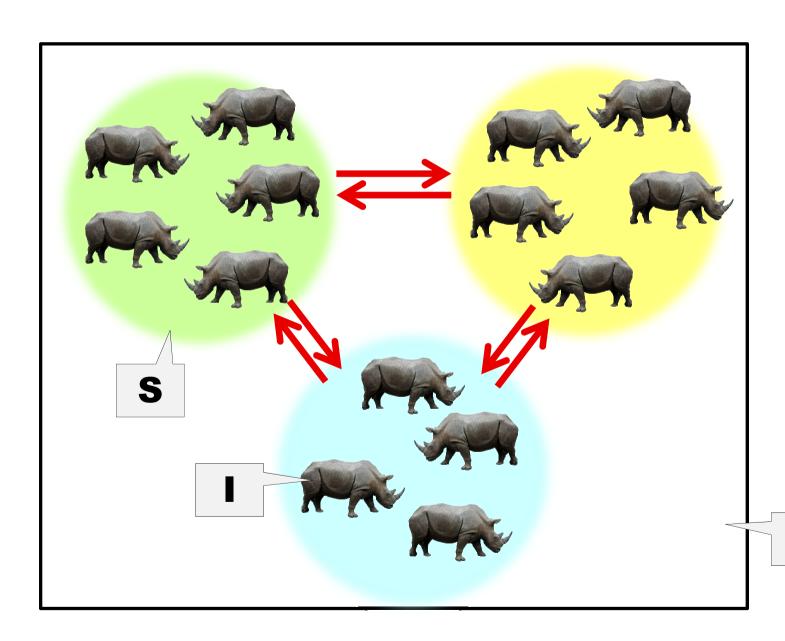








Sewall Wright - F-statistics:





T

F_{IS} (= inbreeding coefficient)

→ reduction of HZ in a subpopulation due to inbreeding

$$F_{IS} = (H_S - H_I)/H_S$$
 $-1 \le F_{IS} \le +1$



 F_{ST} (= fixation index) \rightarrow reduction of HZ due to population substructuring

$$F_{ST} = (H_T - H_S)/H_T$$
 $0 \le F_{ST} \le +1$

 $F_{\text{IT}} \rightarrow$ reduction of HZ both due to population substructuring and inbreeding

$$F_{\text{IT}} = (H_{\text{T}} - H_{\text{I}})/H_{\text{T}}$$
 $(1 - F_{\text{IS}}) (1 - F_{\text{ST}}) = 1 - F_{\text{IT}}$