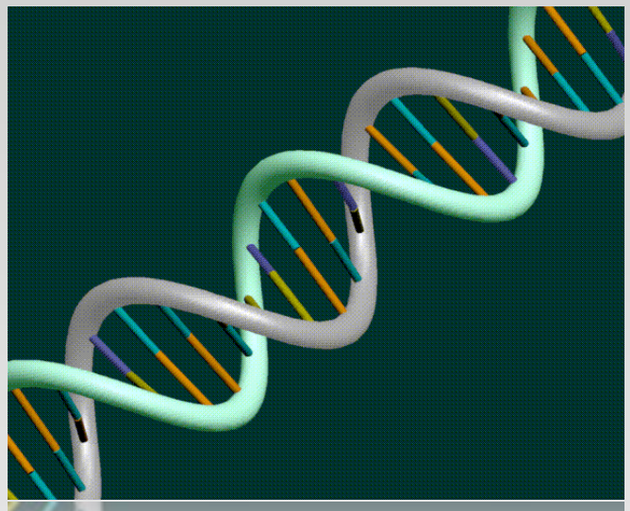
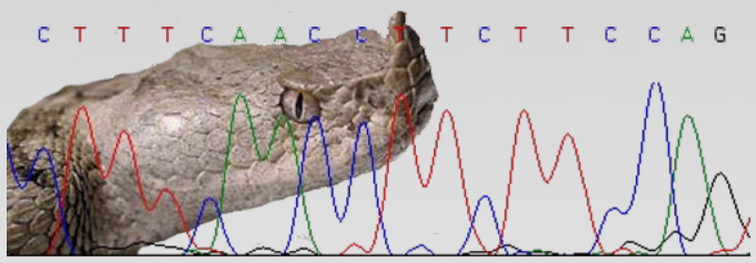
Conservation genetics

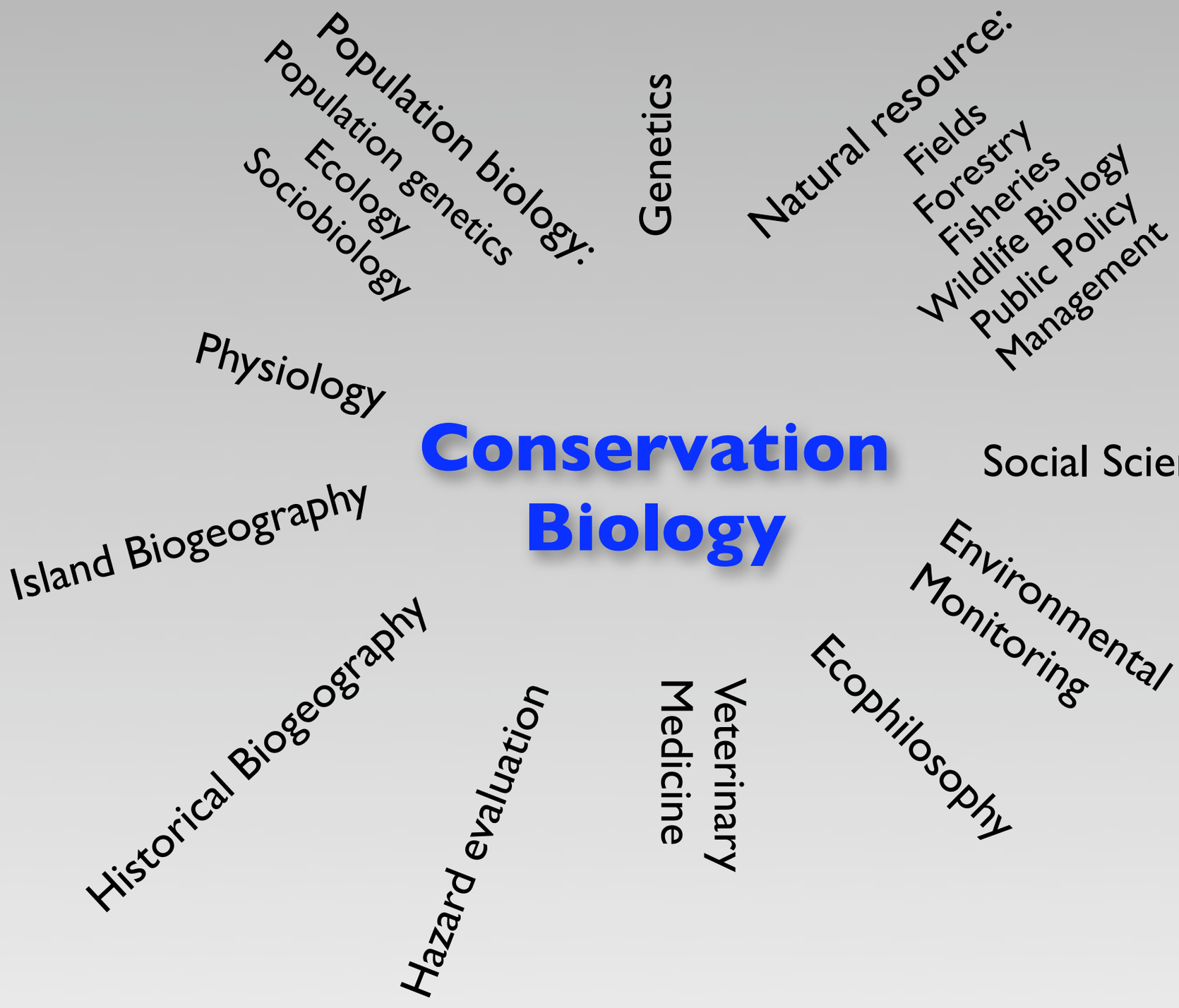




Sylvain Ursenbacher
NLU, room 36

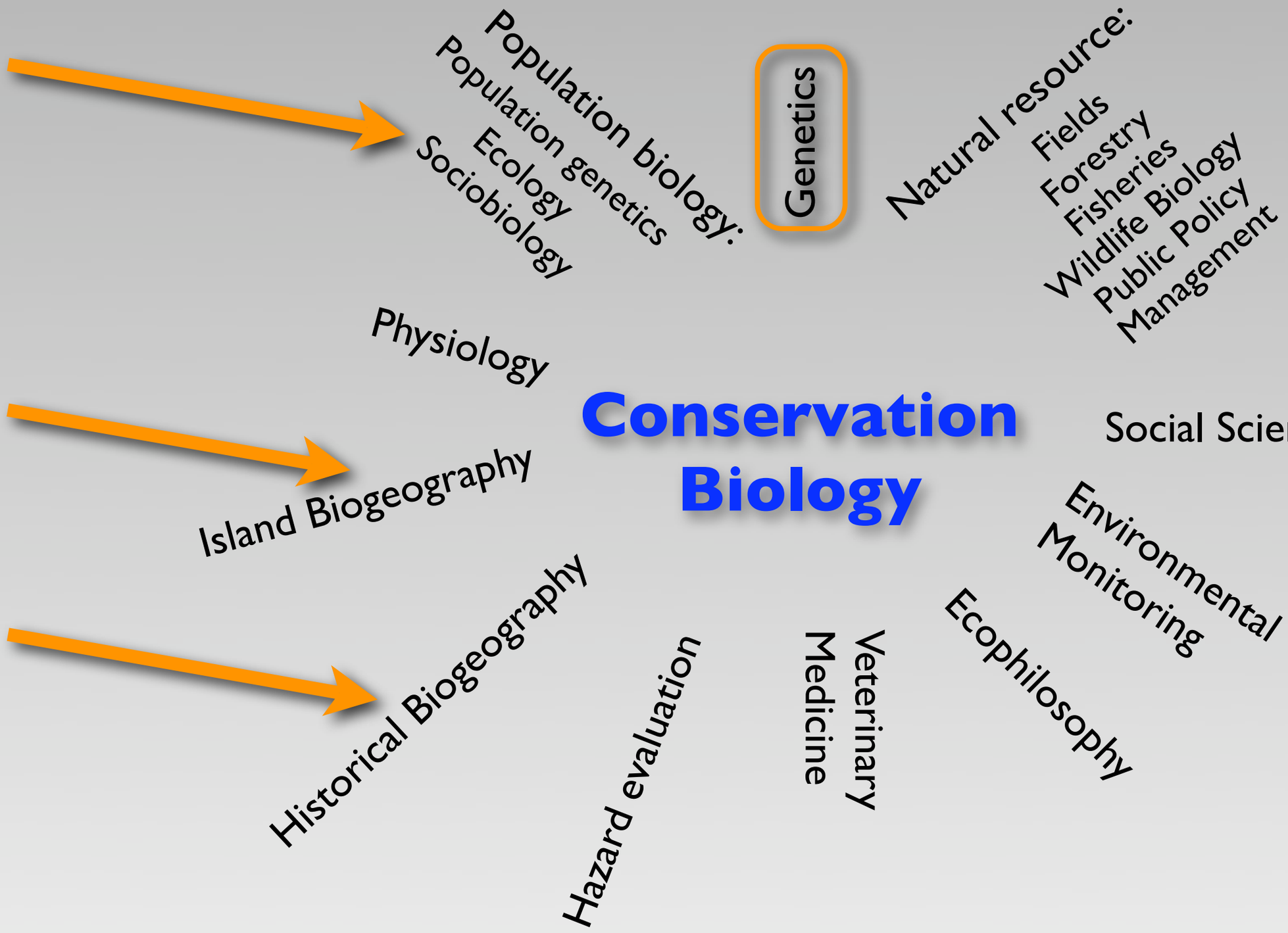
UNI BASEL

Introduction: *Conservation biology*



adapted from Soulé, 1985

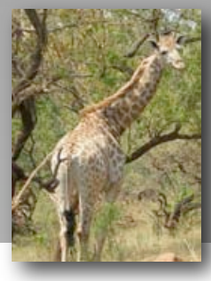
Introduction: *Conservation biology*

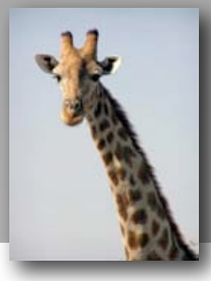


adapted from Soulé, 1985

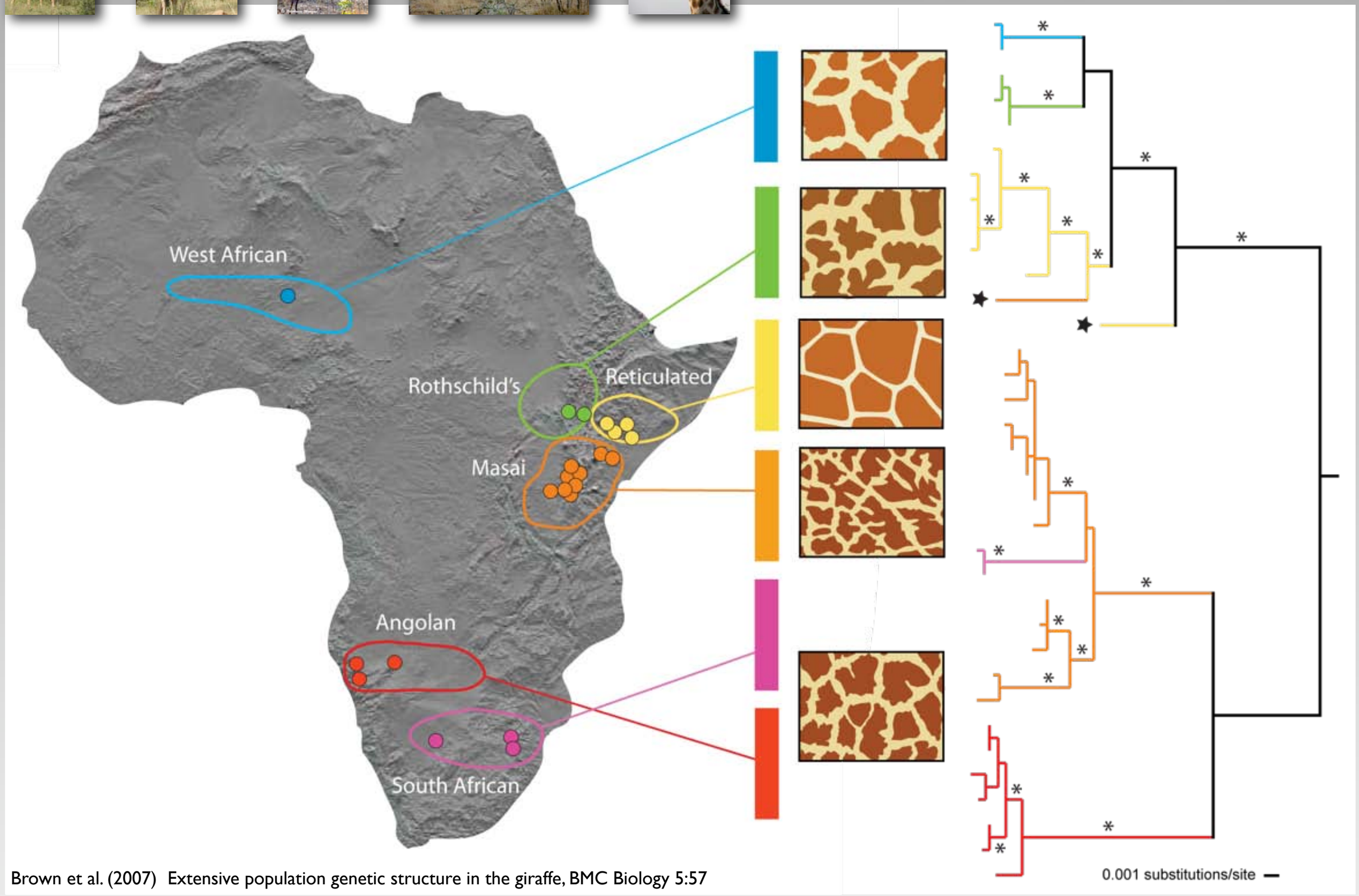
Introduction: *Conservation genetics*

- how genetic analyses can help threatened species: some examples...

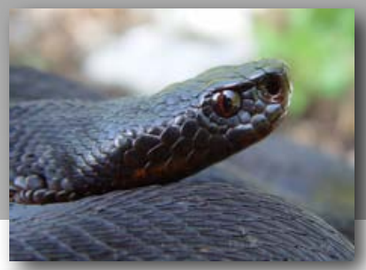



Giraffa camelopardalis



Brown et al. (2007) Extensive population genetic structure in the giraffe, BMC Biology 5:57



Vipera berus

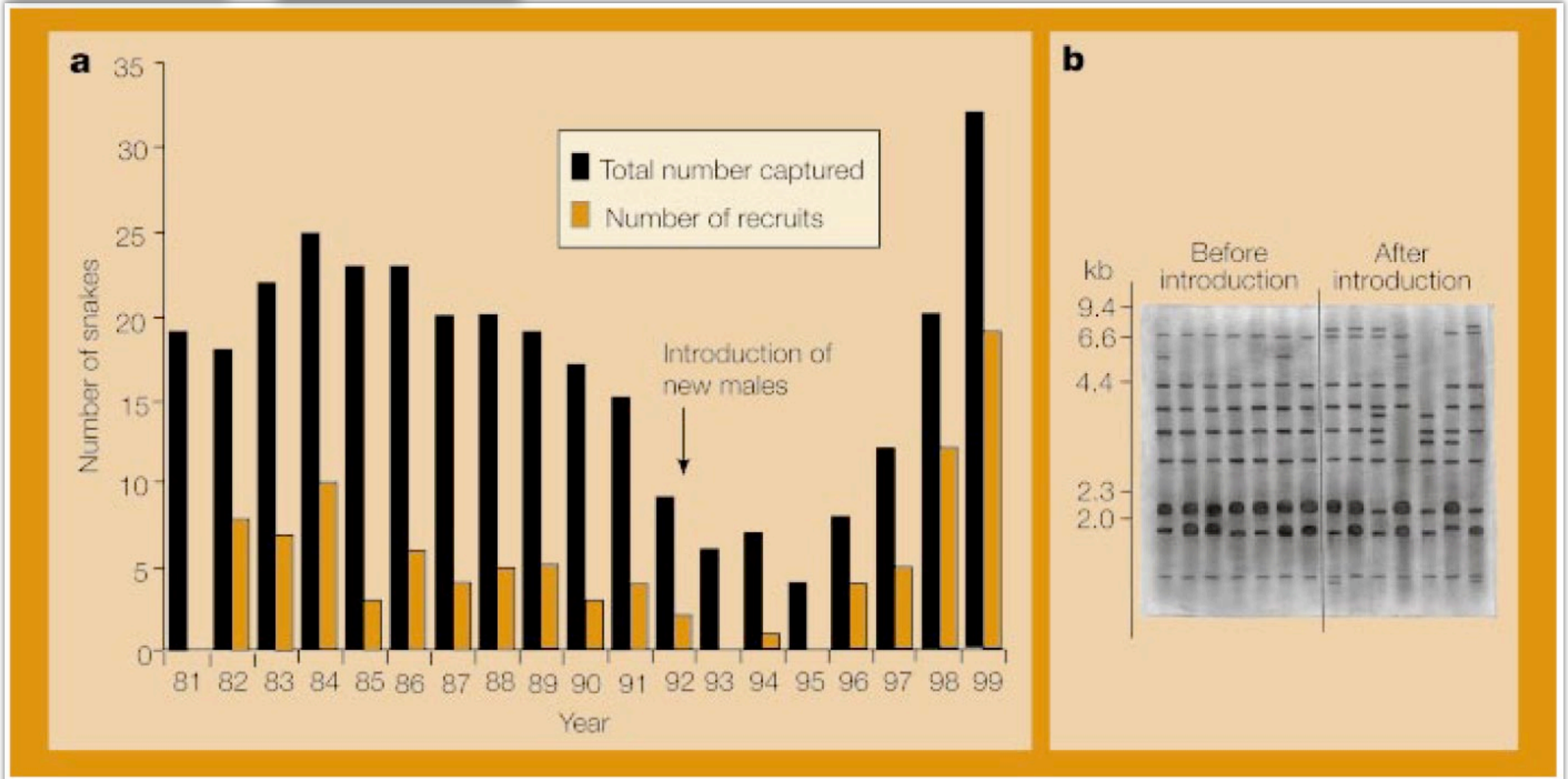


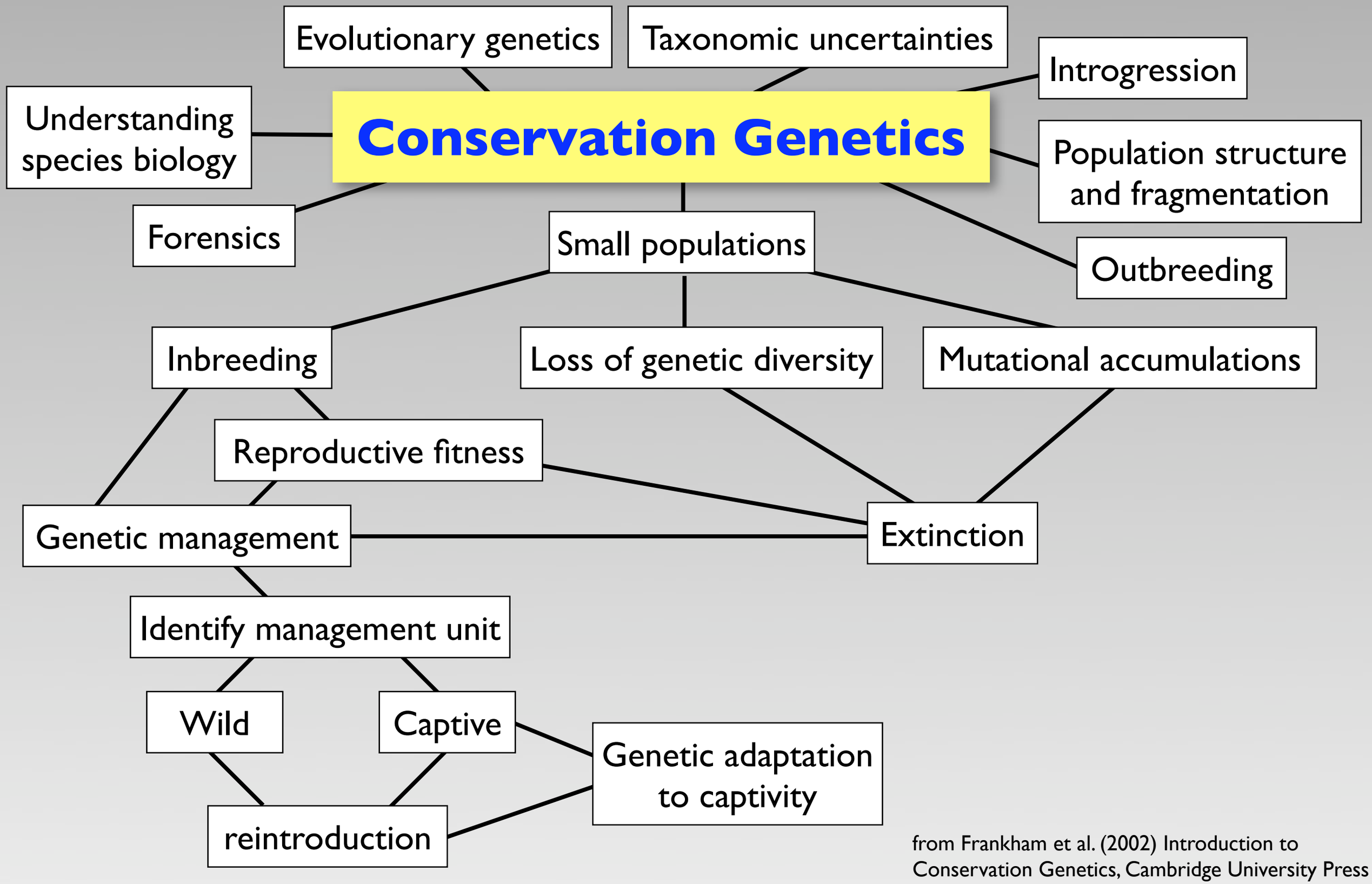
Figure 1 Introducing new males increases the genetic diversity and enables the adder population to recover. **a,** Total number and number of recruited male adders captured in Smygehuk from 1981 to 1999. **b,** Southern-blot analysis of major histocompatability complex (MHC) class I genes in seven males sampled before the introduction of new males (left) and in seven recruited males sampled in 1999 (right).

Madsen et al. (1999) Restoration of an inbred adder population, Nature 402, 34-35

Introduction: *Conservation genetics*

- how genetic analyses can help threatened species: some examples...
  - measure inbreeding / outbreeding depression
  - loss of genetic diversity
  - fragmentation of population and reduction of gene flow
  - genetic drift
  - define management unit
  - understand aspects of species biology important for their conservation

Introduction: *Conservation genetics*



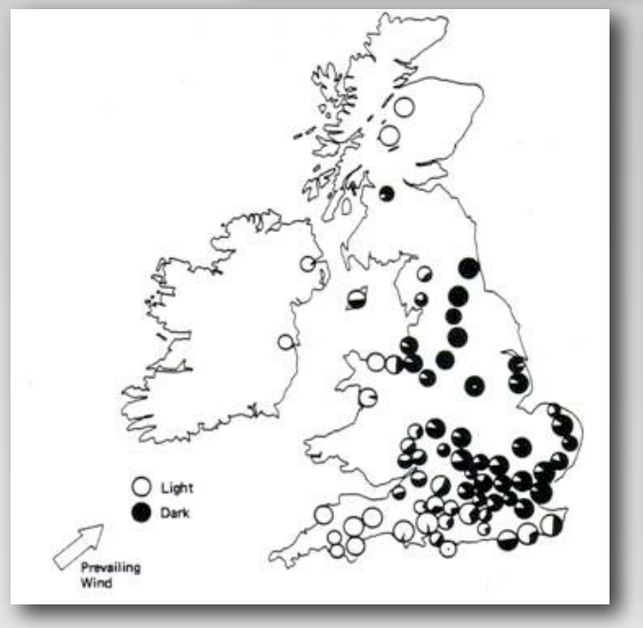
from Frankham et al. (2002) Introduction to Conservation Genetics, Cambridge University Press

Introduction: *why genetic diversity is important in populations...*

- genetic diversity reflects evolutionary potential
  - genetic diversity required to evolve or to adapt to new environment or environmental modifications.
  - ↗ genetic difference between individual ⇒ ↗ fitness of the most adapted

Introduction: *Why genetic diversity is important in populations...*

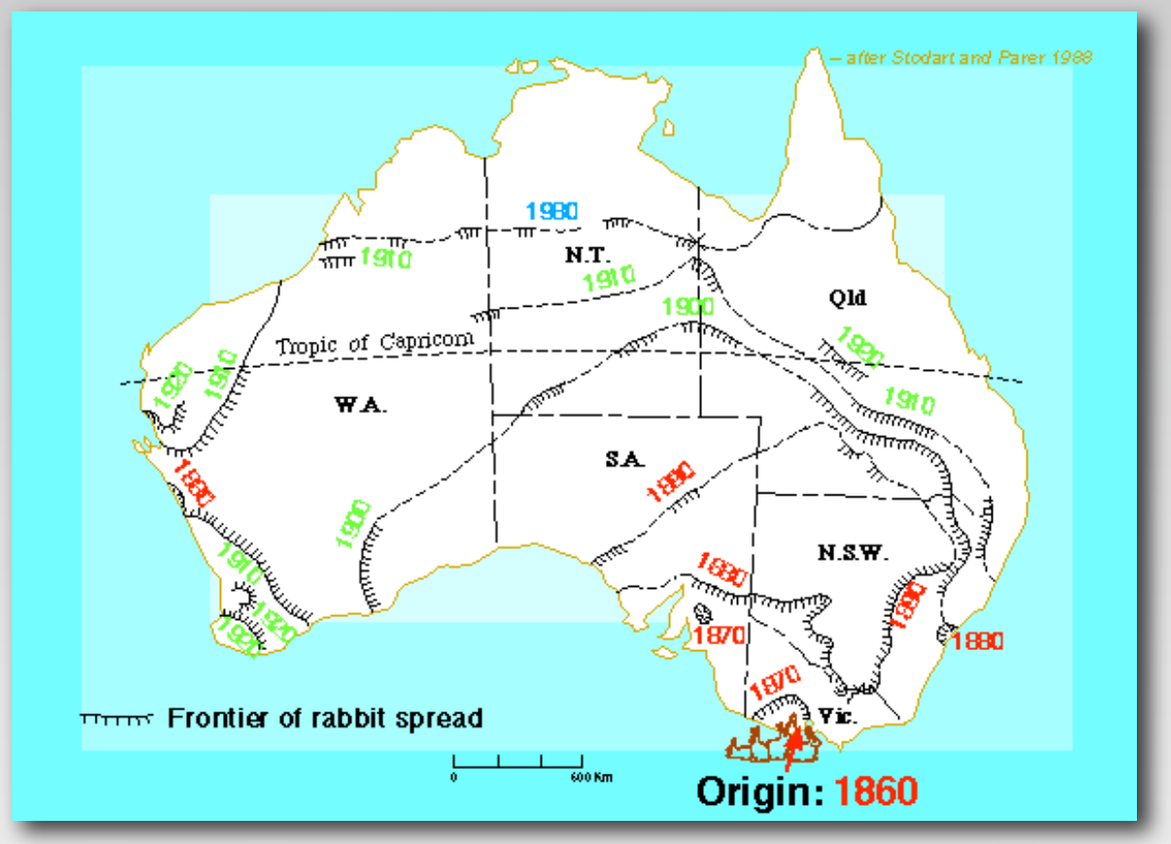
- genetic diversity reflect evolutionary potential
  - example 1 - habitat selection: peppered moth (*Biston betularia*) in UK
    - dark and light forms
    - night: active / day: resting on trees
    - ➡ camouflage critical for survival
    - light form: camouflaged on lichen-covered tree trunks
    - Industrialisation: kill lichen by sulphur pollution
    - ➡ light form: visible / dark form: camouflaged

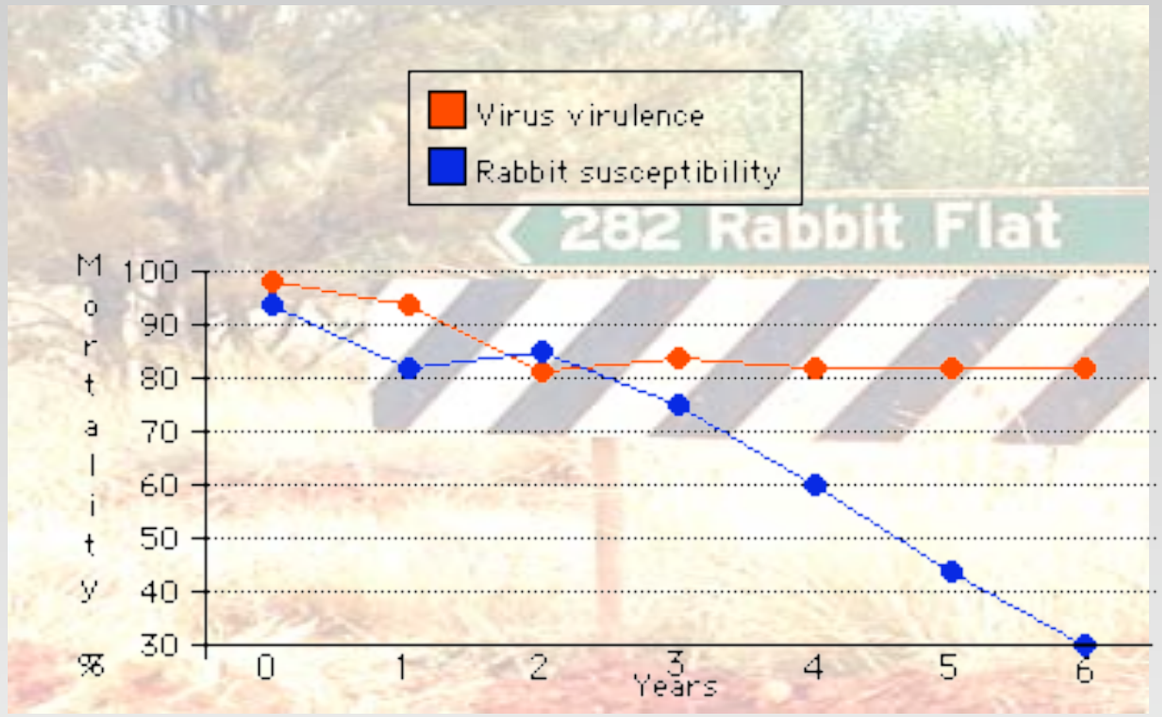


Grant (1999) Fine tuning the peppered moth paradigm, Evolution 53, 980-984
Kettlewell (1973) The Evolution of Melanism, Clarendon Press, Oxford, UK
Majerus (1998) Melanism: Evolution in Action. Oxford University Press
Kettlewell (1958) A survey of the frequencies of *Biston betularia* (L.) (Lep.) and its melanic forms in Great Britain, Heredity 12, 551-572
but see also: Rudge (2006) Myths about moths: a study in contrasts, Endeavour 30, 19-23

Introduction: *why genetic diversity is important in populations...*

- genetic diversity reflect evolutionary potential
  - example 2 - disease resistance: resistance to myxoma virus in Australian rabbits
    - introduction of rabbits in Australia: 1860
    - control measure: introduction of myxoma in 50'
    - ➡ high mortality rate first years
    - high selection for resistance



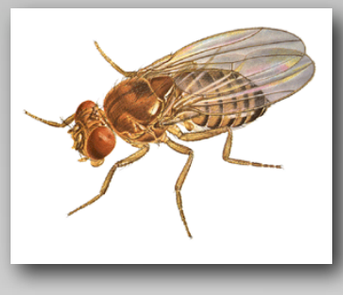


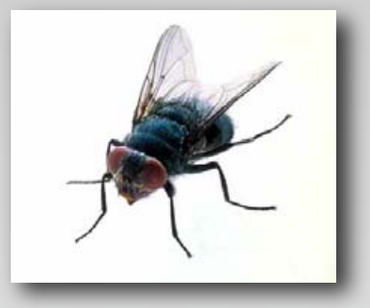
Introduction: *why genetic diversity is important in populations...*

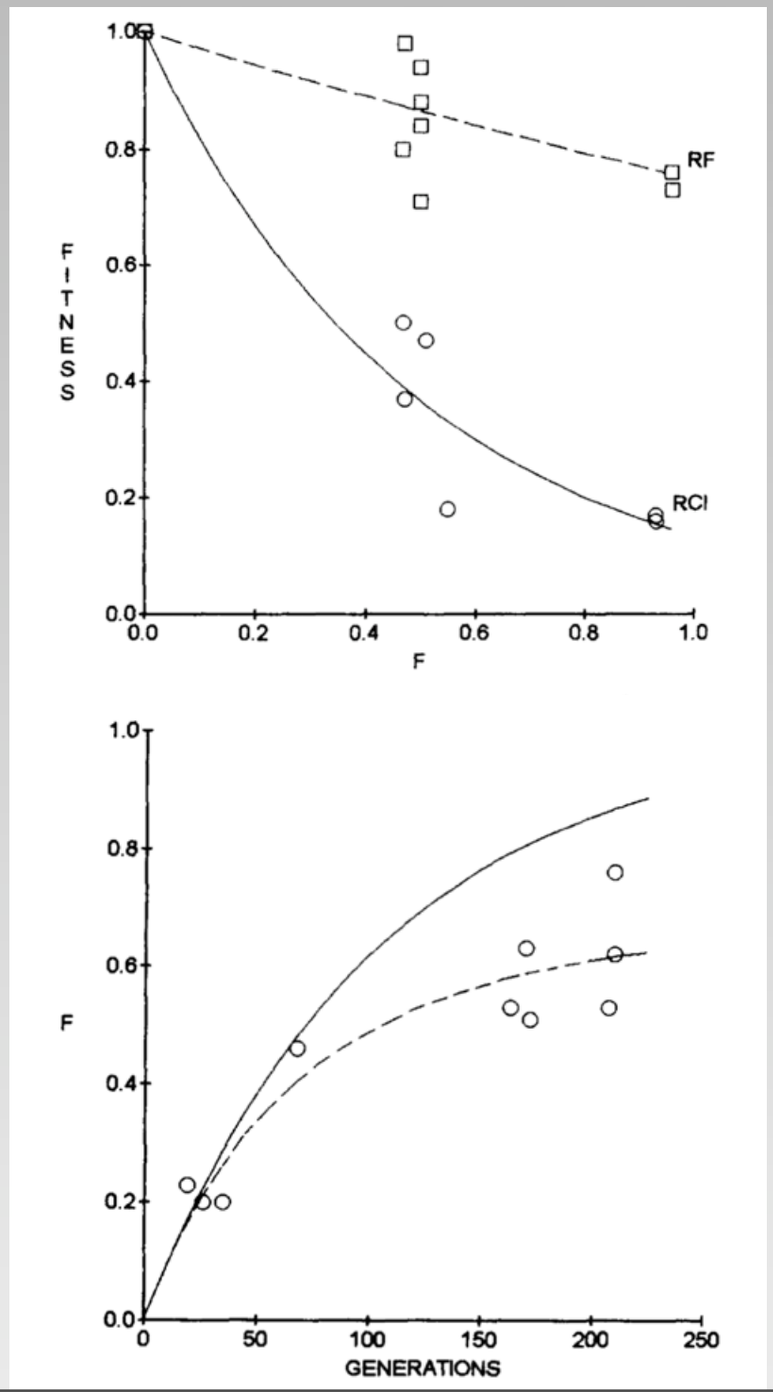
- genetic diversity reflects evolutionary potential
  - genetic diversity required to evolve or to adapt to new environment or environmental modification.
  - ↗ genetic difference between individual ⇒ ↗ fitness of the most adapted
- loss of genetic diversity often associated with inbreeding, reduction of reproductive fitness and extinction risk

Introduction: *Why genetic diversity is important in populations...*

- loss of genetic diversity often associated with inbreeding, reduction of reproductive fitness and extinction risk
  - example 3 - captive fruit fly/housefly populations
    - 60 captive fruit fly (*Drosophila melanogaster*) populations, maintained during 210 generations
      mean population size: 67 individuals
    - ➡ 15/60 populations extinct after 210 generations
    - 6 captive housefly (*Musca domestica*) populations, maintained during 64 generations
      population size: 50 individuals
    - ➡ 5/6 populations extinct after 64 generations







Latter & Mulley (1995) Genetic Adaptation to Captivity and Inbreeding Depression in Small Laboratory Populations of *Drosophila melanogaster*, Genetics 139, 255-266
Reed & Bryant (2000) Experimental tests of minimum viable population size, Animal Conservation 3, 7-14

Introduction: *Why genetic diversity is important in populations...*

- loss of genetic diversity often associated with inbreeding, reduction of reproductive fitness and extinction risk
  - example 4 - large metapopulation (Finland) of the Glanville fritillary butterfly (*Melitaea cinxia*)
    - 42 butterfly populations genotyped in 1995
    - survival and extinction recorded in 1996
    - ➡ 36 populations survived
    - extinction rate high for populations with lower heterozygosity even corrected for demographic and environmental variables (pop. size, area, ...)

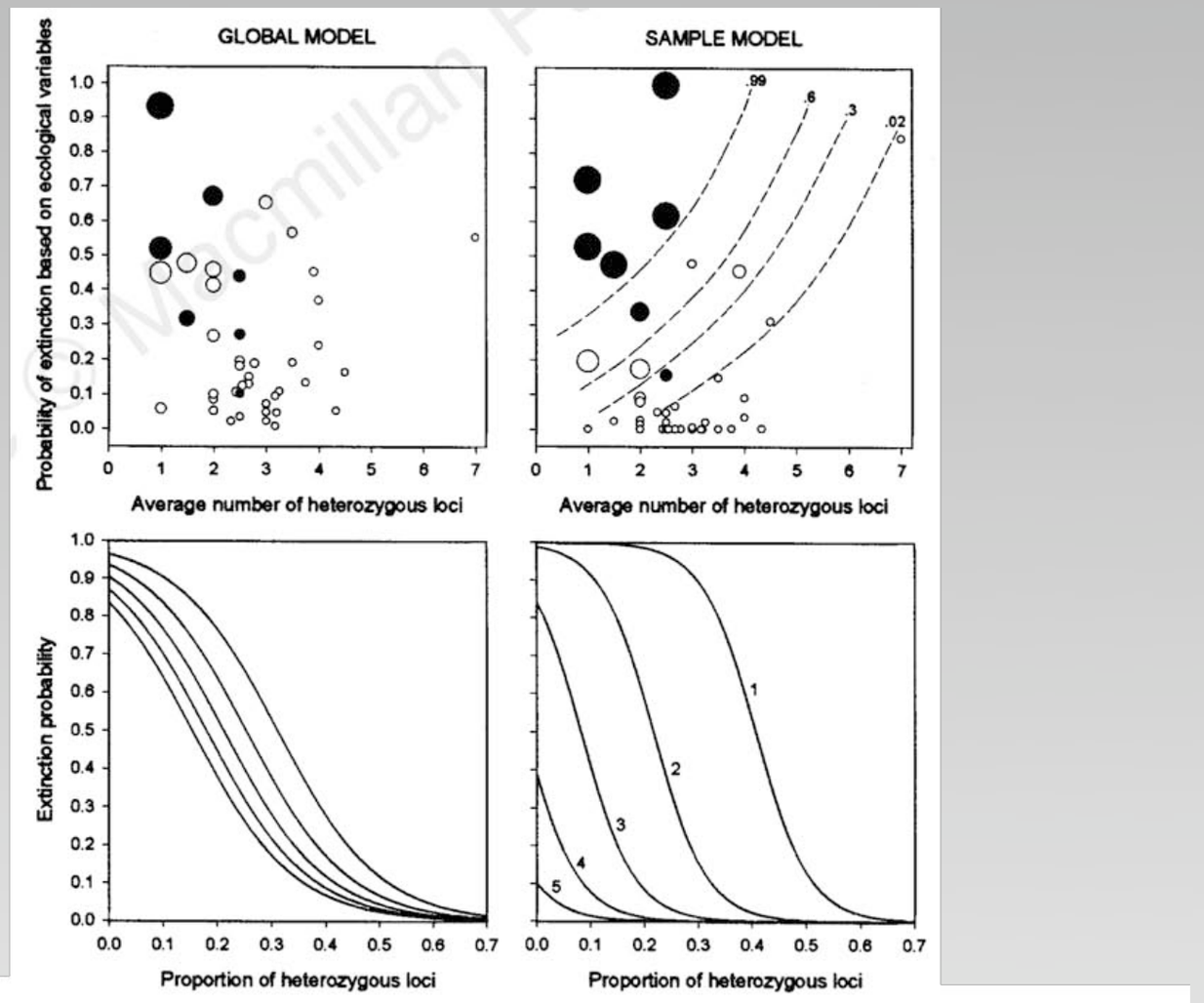


Figure 2 For both global and sample models (Table 1), the upper panels show: (1) the observed average number of heterozygous loci in extinct (black) and surviving (white) populations; (2) the probability of extinction predicted by the models without heterozygosity compared with the observed heterozygosity; (3) the probability of extinction predicted by the full model, including heterozygosity (proportional to circle size). For the sample model, we have drawn appropriate isoclines for the extinction risk predicted by the model, including ecological factors and heterozygosity. These figures illustrate that both the ecological factors and heterozygosity influence the extinction risk (for statistical analysis, see Table 1). Lower panels show the relationship between the risk of local extinction and heterozygosity predicted by the global and sample models (Table 1). Model predictions are shown for local population sizes of 1–5 larval groups, fixed at the lower quartile value of change in regional density (N_{trend}) and the lower quartile value of meadow area in the global model; and fixed at the lower quartile value of regional density (N_{neigh}) and median flower abundance in the sample model.

Saccheri et al. (1998) Inbreeding and extinction in a butterfly metapopulation, Nature 392, 491-494

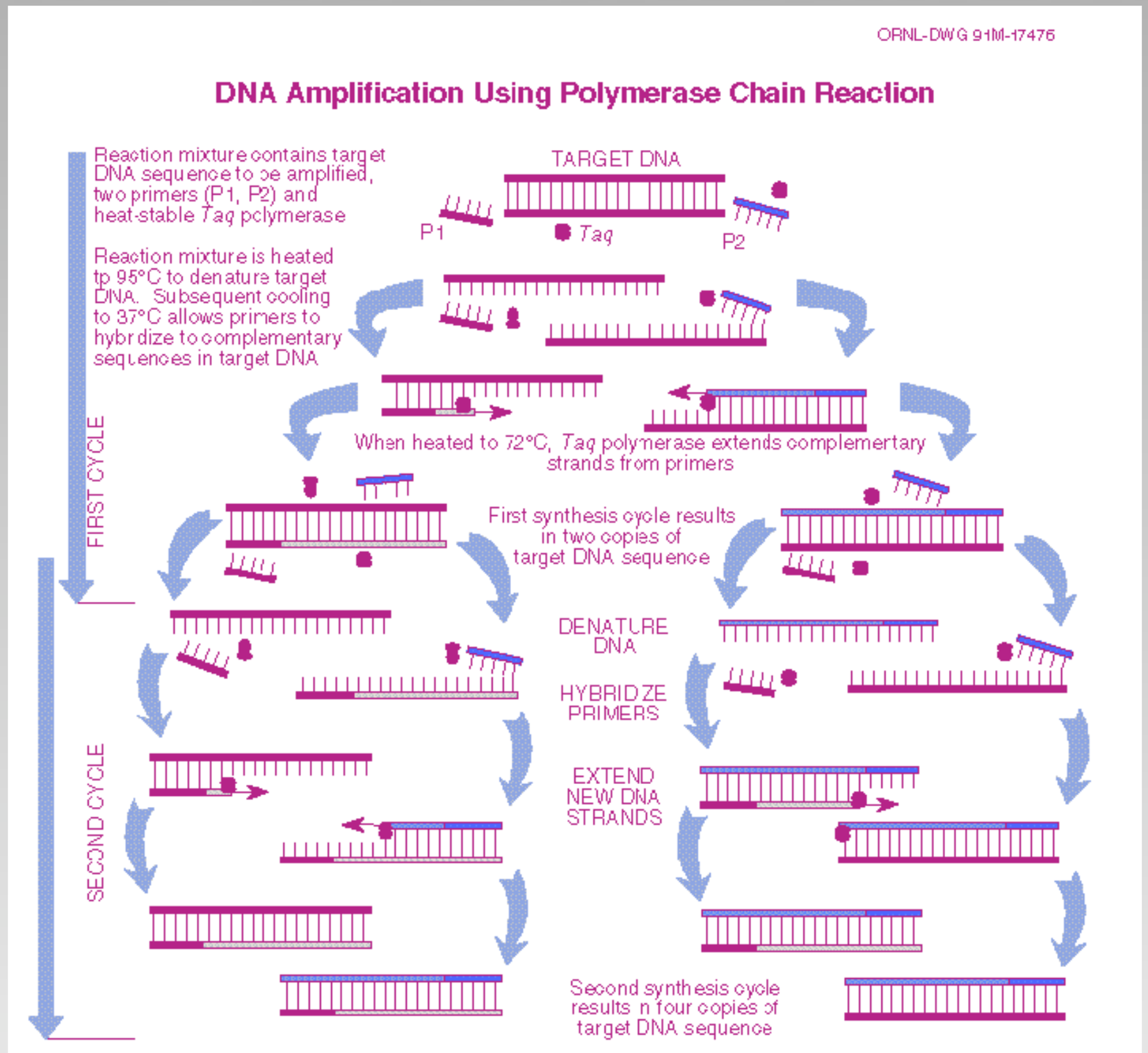
Genetic tools: *DNA sampling*

- invasive methods (dead animals)
  - entire animal/plants (e.g. insects)
  - internal tissue: liver, heart, ...
- non-invasive methods
  - blood sample
  - part of the body: hairs, feathers, scales, sloughed skin, ... leafs, flowers, ...
  - buccal swab
  - faeces
  - ...

Genetic tools: *DNA extraction*

- first: lysis of the tissue/sample using proteinase
- numerous protocols
  - standard Phenol/Chloroform (Sambrock et al. 1989)
    ⊕ low cost / ⊖ high toxicity
  - CTAB: more adapted to plants (or amphibians)
  - CHELEX:
    ⊕ quick / ⊖ not for long storage
  - columns: several companies, e. g. Qiagen, Promega, Sigma,...
    ⊖ expensive / ⊕ high purity DNA

Genetic tools: *DNA amplification (PCR)*



Source: *DNA Science*, see Fig. 13.

Measuring genetic diversity

- different markers (regions)
  - under selection or not
  - lineage: maternal, paternal or both
  - easy/difficult to develop, use or analyse
  - cheap/expensive

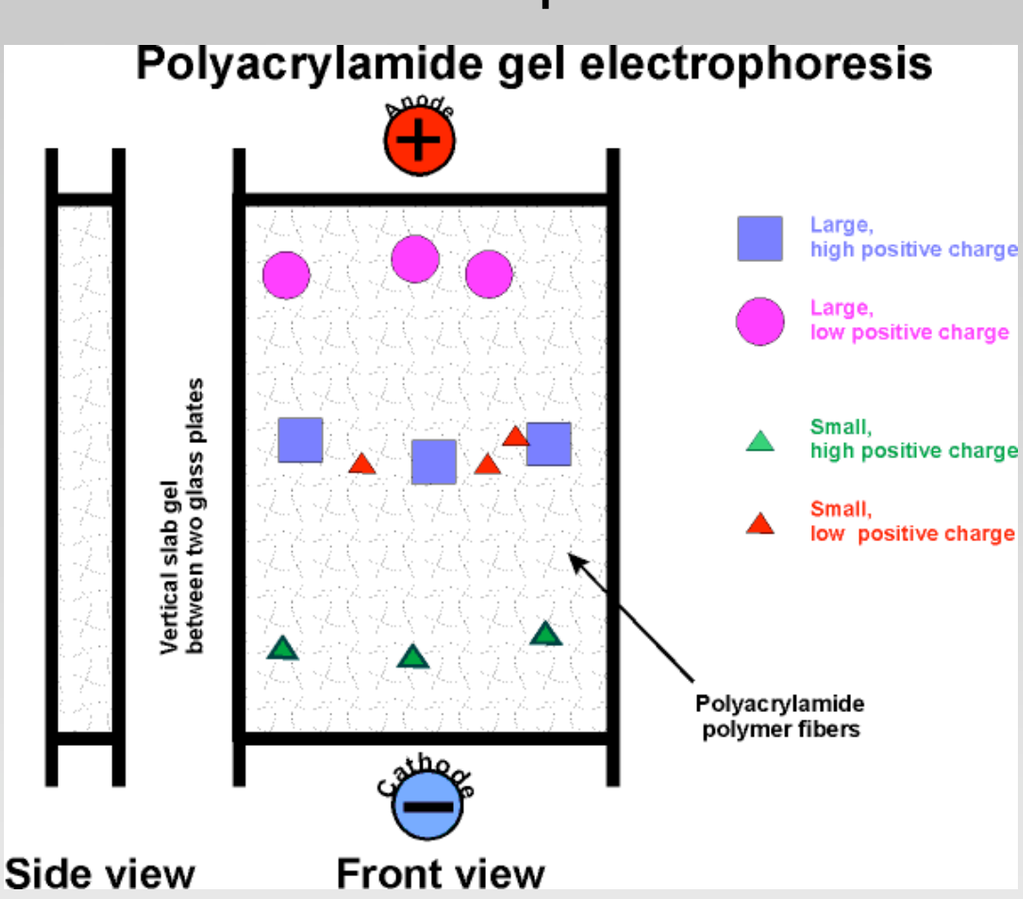
  - Proteins / Allozymes
  - sequencing
  - Restriction Fragment Length Polymorphism (RFLP)
    Amplified Fragment Length Polymorphism (AFLP)
    Random Amplified Polymorphic DNA (RAPD)
    DNA fingerprints (minisatellites)
  - microsatellites
    Single Nucleotide Polymorphism (SNP)
    Single Strand Conformational Polymorphism (SSCP)

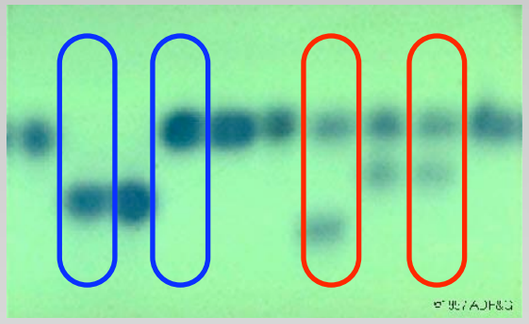
Genetic markers: *Proteins / Allozymes*

- separate proteins according to their electric charge and molecular weights

DNA coding for a protein	...ATG CTT GAC GTT ...	...ATG CTT G**G**C GTT ...
mRNA	...AUG CUU GAC GUU ...	...AUG CUU G**G**C GUU ...
amino acid composition	... -met - leu - asp - val - ...	... - met - leu - **gly** - val - ...

- electrophoresis



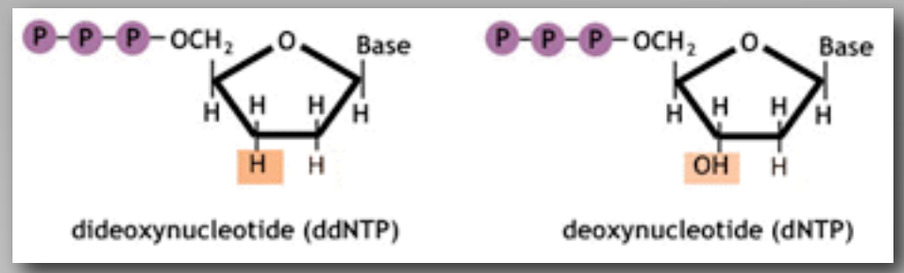


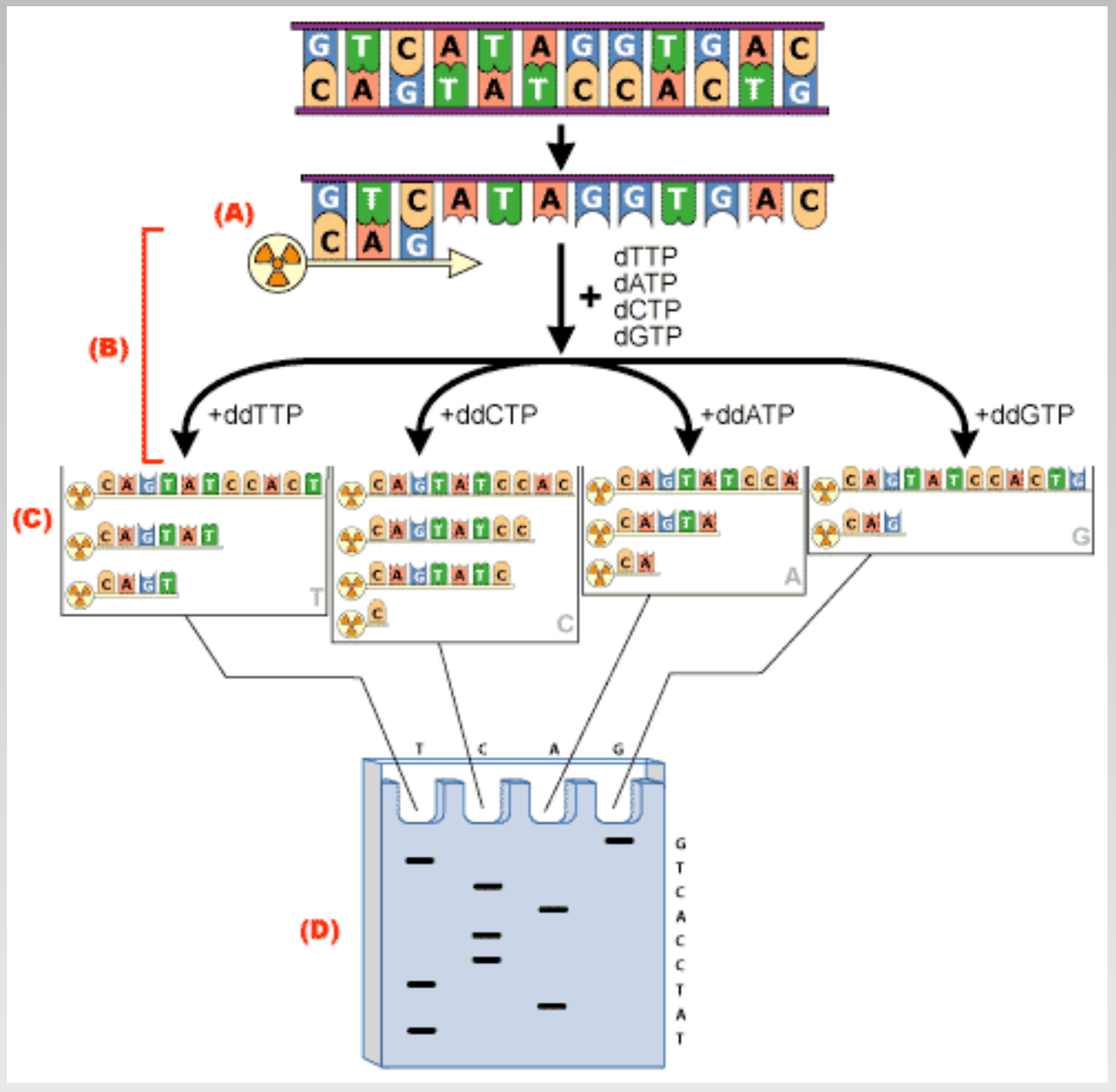
⊖ only 30% of DNA base changes result in charge changes: underestimation of genetic diversity

⊖ need high quantity of material (blood, kidney, liver) not really useful for endangered species

Genetic markers: *Sequencing*

- "reading" DNA sequences





DNA Polymerase reads the template strand and synthesizes a new second strand to match:

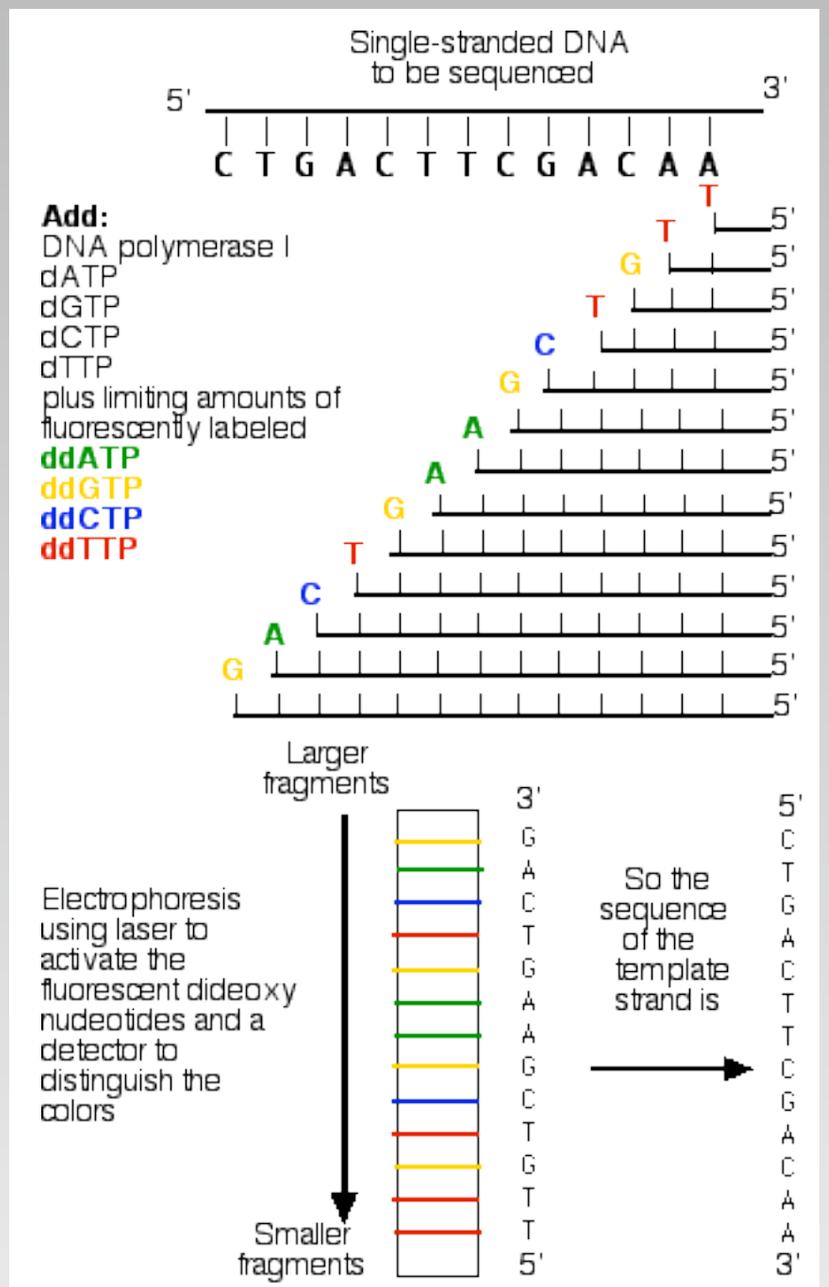
```
5' - TACGCGGTAACGGTATGTTCGACCGTTTAGCTACCGAT
3' - ATGCGCCATTGCCATACAAGCTGGCAAATCGATGGCTAGAGATCCAA - 5'
```

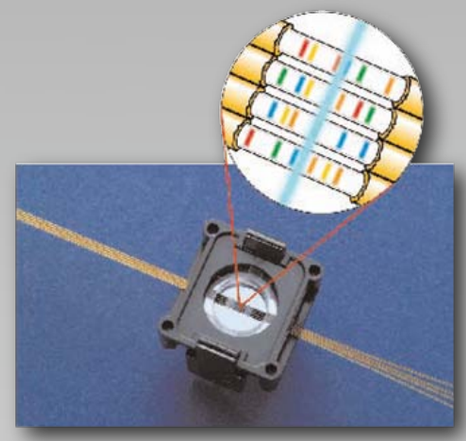
IF 5% of the T nucleotides are actually dideoxy T, then each strand will terminate when it gets a ddT on its growing end:

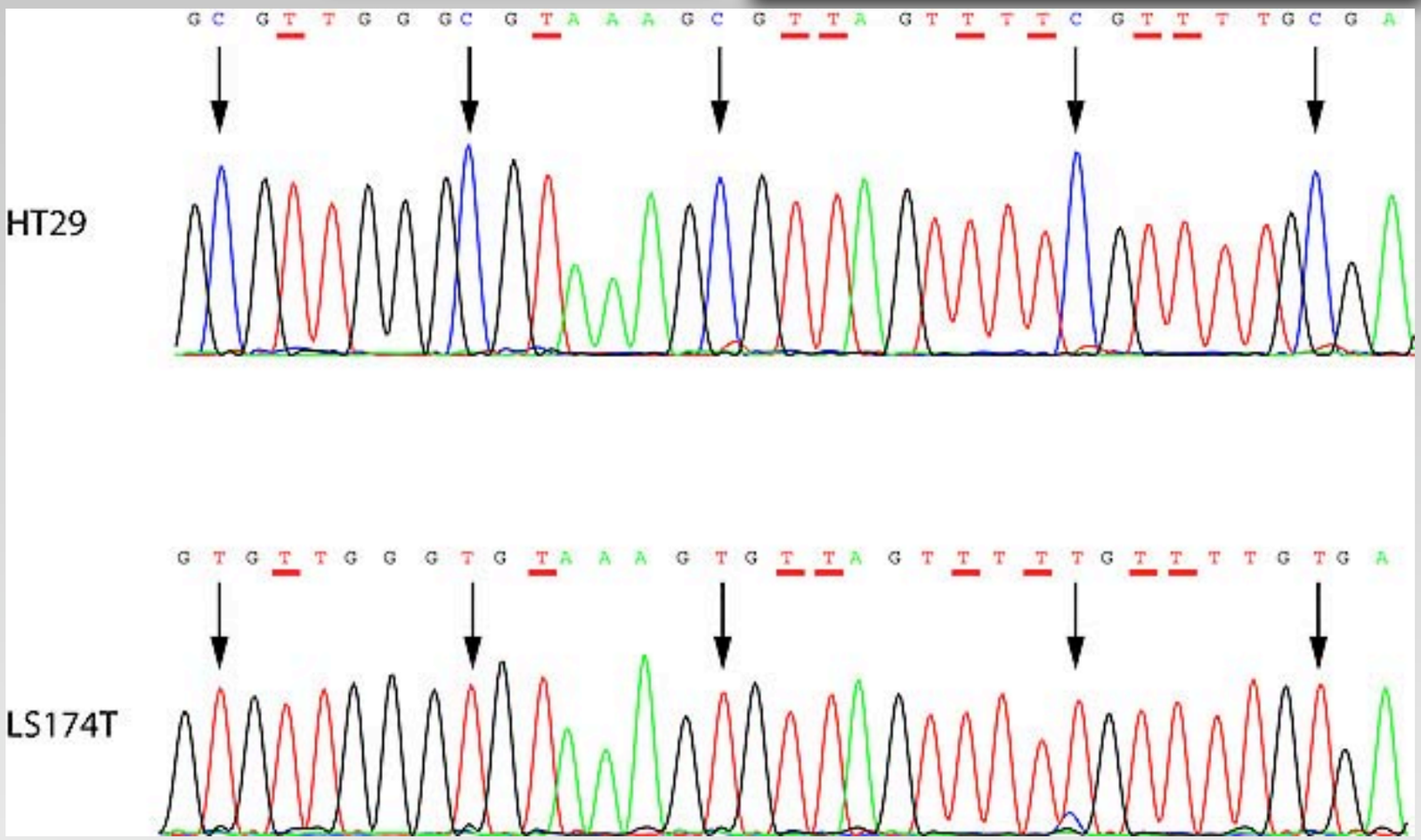
```
5' - TACGCGGTAACGGTATGTTCGACCGTTTAGCTACCGAT•
5' - TACGCGGTAACGGTATGTTCGACCGTTTAGCT•
5' - TACGCGGTAACGGTATGTTCGACCGTTT•
5' - TACGCGGTAACGGTATGTTCGACCGTT•
5' - TACGCGGTAACGGTATGTTCGACCGT•
5' - TACGCGGTAACGGTATGTT•
5' - TACGCGGTAACGGTATGT•
5' - TACGCGGTAACGGTAT•
5' - TACGCGGTAACGGT•
5' - TACGCGGT•
```

Genetic markers: *Sequencing*

- "reading" DNA sequences





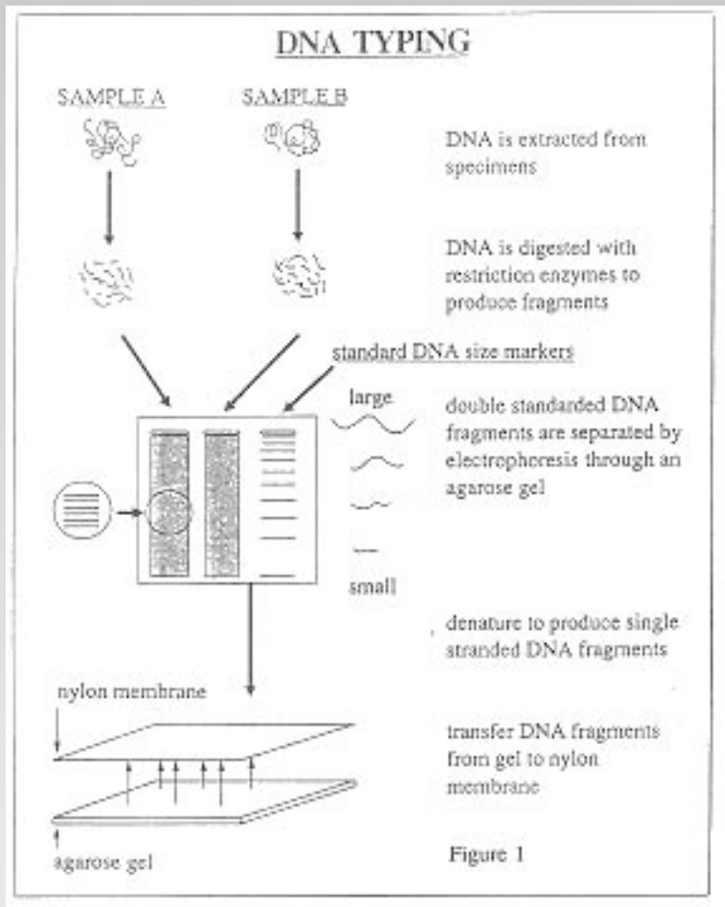


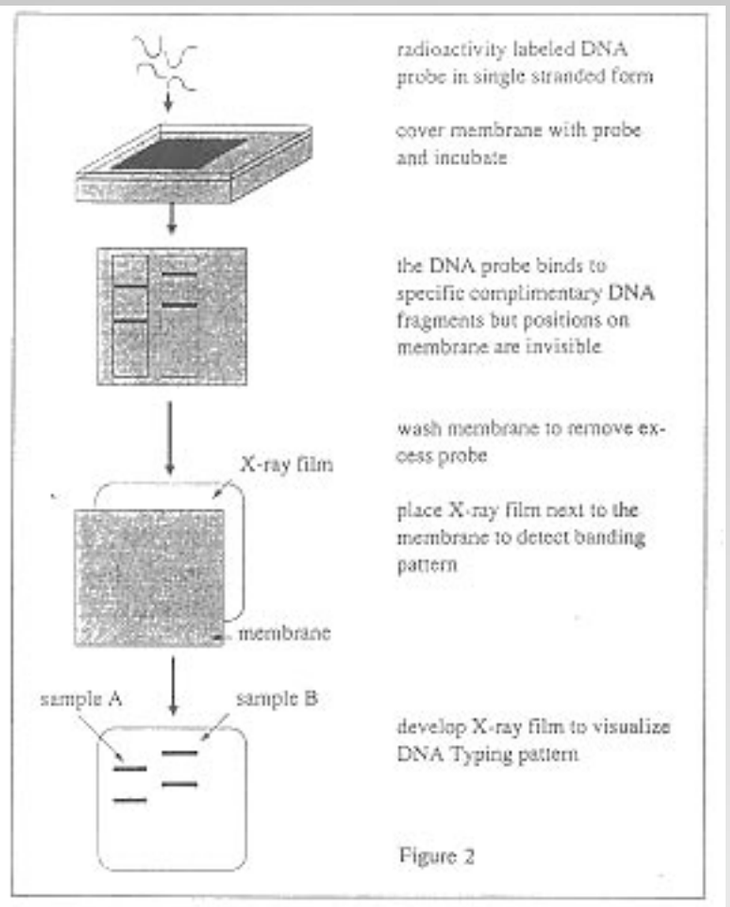
Genetic markers: *Sequencing*

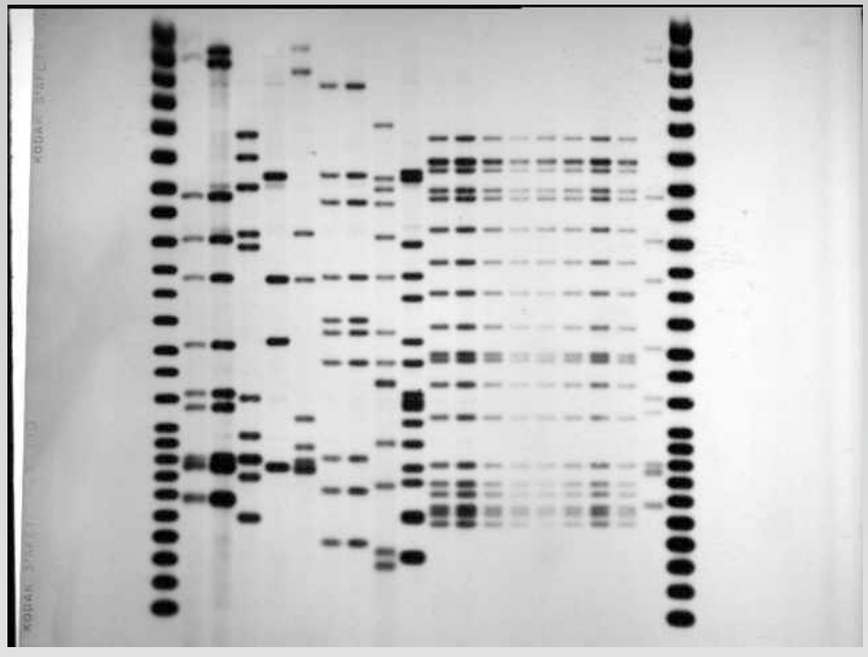
- "reading" DNA sequences
  - ⊖ high cost
  - ⊖ problems with heterozygosity
  - ⊖ primers sequences must be known

Genetic markers: *Restriction Fragment Length Polymorphism (RFLP)*

- use define restriction enzymes to cut randomly in the whole genome → numerous DNA fragments with diff. sizes
  - differences at the restriction enzyme cutting site
- electrophoresis (agarose or other)
  - ⊖ need large amount of DNA; not for non-invasive methods

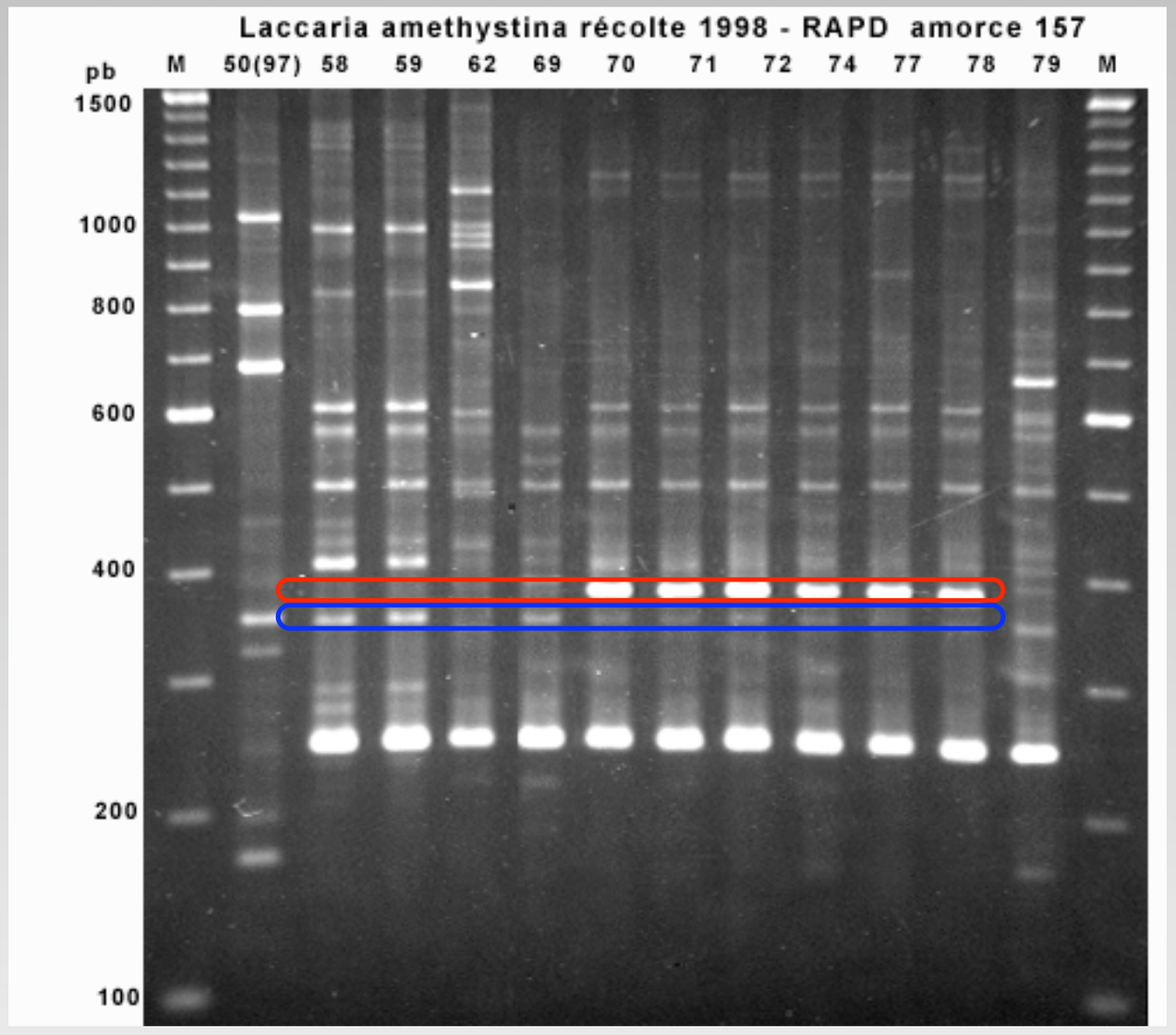






Genetic markers: *Random Amplified Polymorphic DNA (RAPD)*

- PCR reaction using random primers (10-20 bp), producing several fragments with different length
- electrophoresis to see the different fragments



⊖ repeatability of the results not always good...

⊖ dominant markers

Genetic markers: *dominance / co-dominance principle*

AA	Aa	aa
---P___P----	---P___P----	---P----------
---P___P----	---P----------	---P----------
DNA fragment on a gel		
___	___	no band

A: dominant / a: recessive
P: primer similar to the DNA seq.
_____ PCR product

- dominance: when heterozygotes are not distinguishable from homozygotes
  - AA with PCR product, aa without PCR product, Aa with PCR product
- co-dominance: when heterozygotes are distinguishable from homozygotes
  - AA with low mobility, aa with high mobility, Aa with a medium mobility

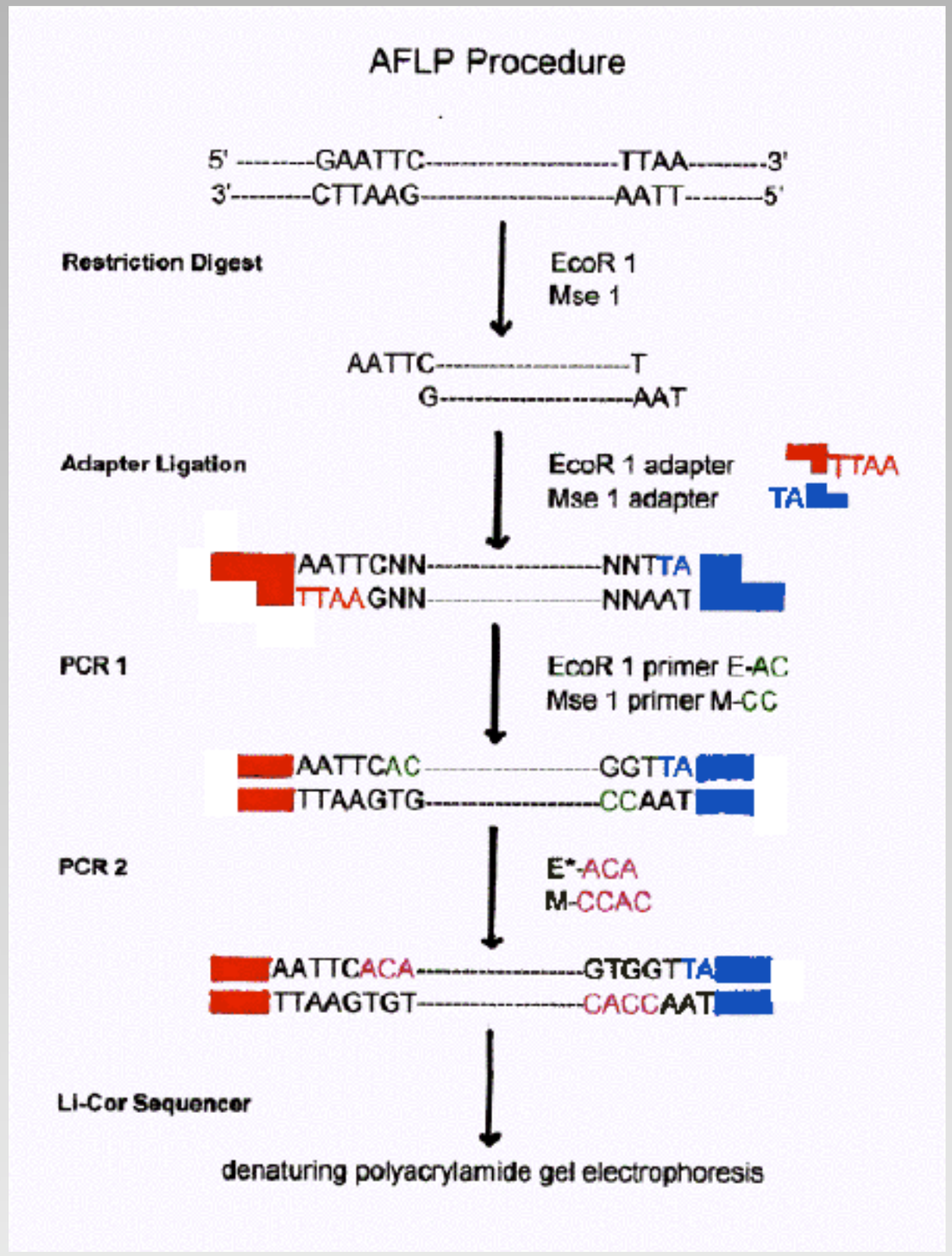
→ difficulties in the analyses

Genetic markers: *Amplified Fragment Length Polymorphism (AFLP)*

- method close to RAPD
- DNA cut with a restriction enzyme, and short DNA fragments of known sequence are attached to the cut ends

  ⊕ more accurate than RFLP
  no repeatable problems

  ⊖ dominant markers



Genetic markers: *microsatellites*

- also named STR (short tandem repeats) or
  SSR (simple sequence repeats)
- tandem repeats of a short DNA segment (1-5 bp)

maternal origin ATATATATATATATATATAT $(AT)_9$

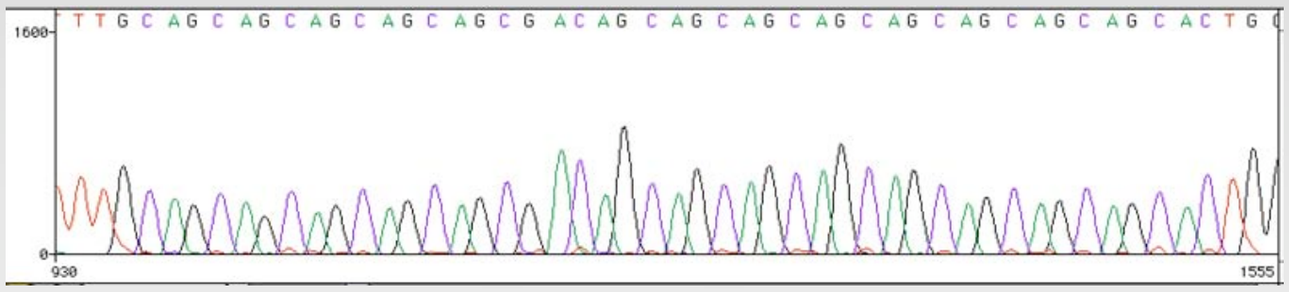
paternal origin ATATATATATATATATATATATAT $(AT)_{11}$

- between two conserved regions flanking the microsatellites

stable ATATATATATATATATATAT stable

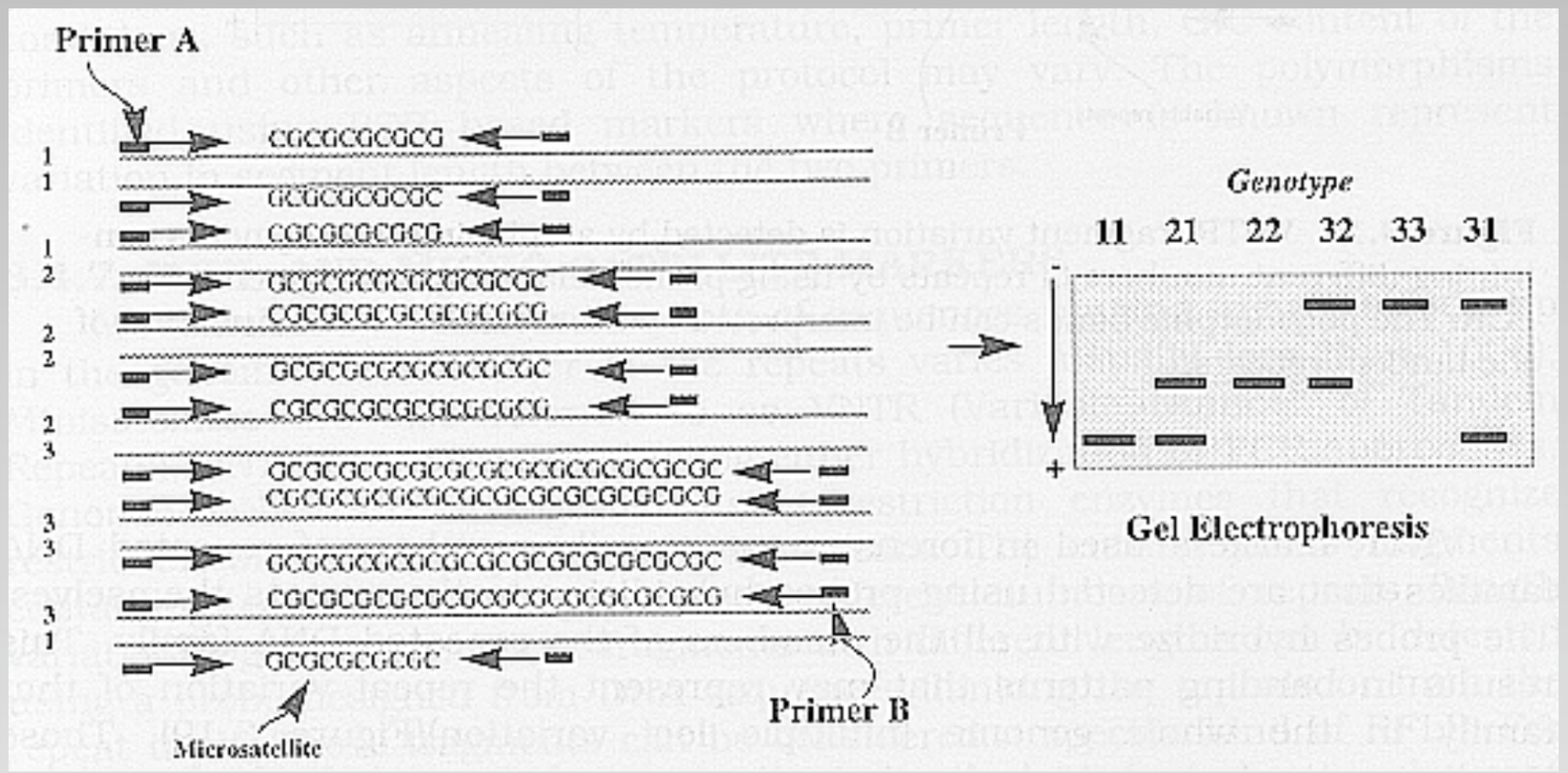
stable ATATATATATATATATATATATAT stable

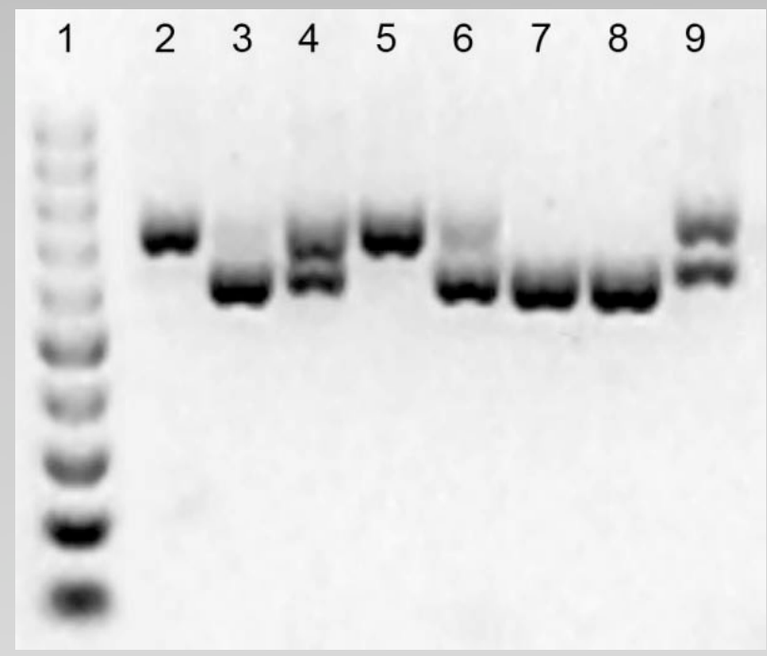
- reason of the polymorphism: polymerase "slippage" or "stuttering"

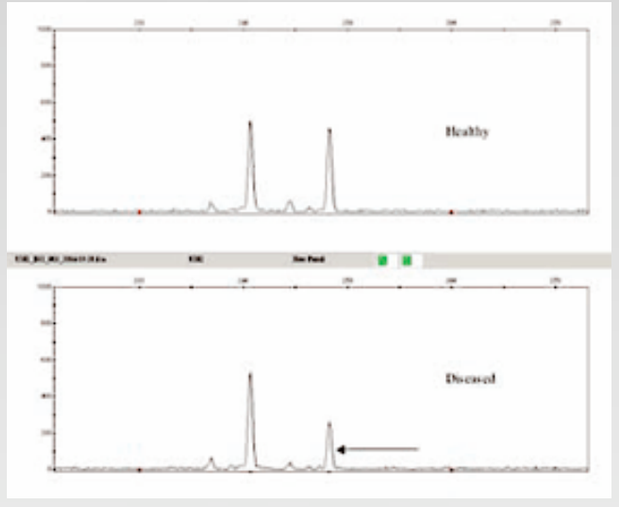


Genetic markers: *microsatellites*

- must found the conserved regions flanking the microsatellites
- separation using electrophoresis (agarose gel or sequencer)







⊖ difficult to identify the conserved regions flanking the microsatellites

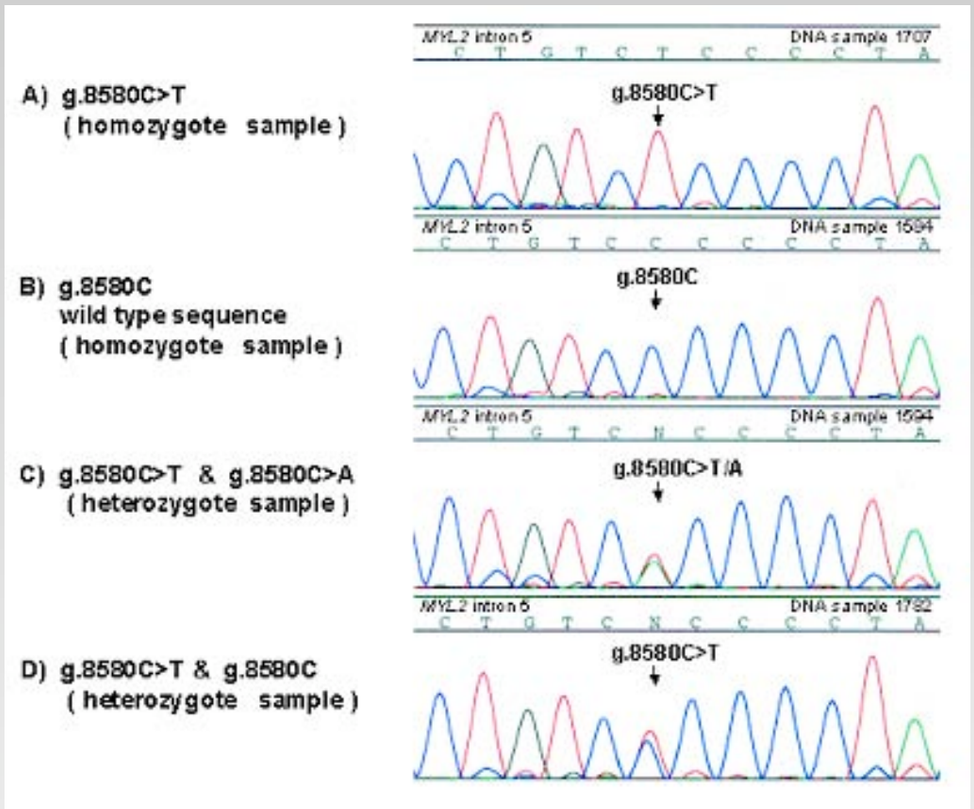
Genetic markers: *other markers*

- *DNA fingerprints (minisatellites)*
  - core repeat sequences of 10-100 bp

    ⊕ highly variable / ⊖ high quantity of DNA, difficult to set-up / old method

- *Single Nucleotide Polymorphism (SNP)*
  - punctual mutation in a gene, present in >1% of the population

    ⊕ possible difference in the protein expression / ⊖ need sequencing



Genetic markers: *other markers*

- *DNA fingerprints (minisatellites)*
  - core repeat sequences of 10-100 bp

    ⊕ highly variable / ⊖ high quantity of DNA, difficult to set-up / old method

- *Single Nucleotide Polymorphism (SNP)*
  - punctual mutation in a gene, present in >1% of the population

    ⊕ possible difference in the protein expression / ⊖ need sequencing

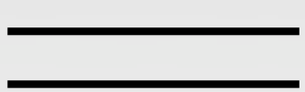
- *Single Strand Conformational Polymorphism (SSCP)*
  - *using difference of mobility for slightly different DNA fragments*

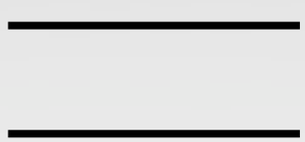
    ⊕ differentiation without sequencing / ⊖ difficult to set-up

GATTGCGTA**G**CGTACTAGCG**A**CAGCTAG
GATTGCGTA**C**CGTACTAGCG**T**CAGCTAG

GATTGCGTA**C**CGTACTAGCG**A**CAGCTAG
GATTGCGTA**C**CGTACTAGCG**T**CAGCTAG

gel





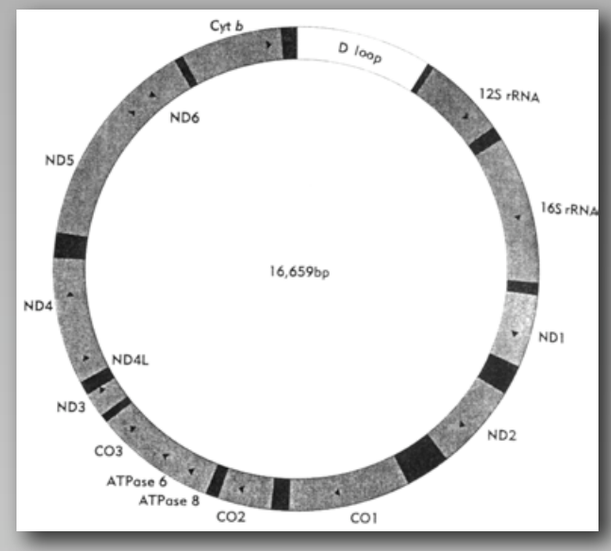
Genetic markers: *summary*

	first use	Basis	Polymorphism	Level of polymorphism	Dominance / co-dominance	selection	development	cost	non-invasive sampling
Allozymes	1966	amino-acid polymorphism	change in amino-acid	low	co-dominant	under	none	low	no
sequencing	1975	sequencing of PCR product of a defined gene/ region	nucleotide polymorphism, inserts, deletion	low/high	co-dominant	no or under	none	high	yes
RFLP	1970's	Randomly fragmented DNA	length of the fragments	medium	co-dominant	no (rarely under)	limited	moderate	no
RAPD	1990	Random amplified DNA fragments	amplifiable or not amplifiable fragment	medium	dominant	no (rarely under)	limited	low/ moderate	yes
AFLP	1995	Random amplified DNA fragments	amplifiable or not amplifiable fragment	medium	dominant	no (rarely under)	limited	moderate / high	yes
microsatellites	end of 1980's	PCR amplification of a unique loci, harbouring simple sequences repeats	variation in the number of repeats	high	co-dominant	no	long time, high cost	moderate	yes

Genetic markers: *summary*

	first use	Basis	Polymorphism	Level of polymorphism	Dominance / co-dominance	selection	development	cost	non-invasive sampling
Allozymes	1966	amino-acid polymorphism	change in amino-acid	low	co-dominant	under	none	low	no
sequencing	1975	sequencing of PCR product of a defined gene/ region	nucleotide polymorphism, inserts, deletion	low/high	co-dominant	no or under	none	high	yes
RFLP	1970's	Randomly fragmented DNA	length of the fragments	medium	co-dominant	no (rarely under)	limited	moderate	no
RAPD	1990	Random amplified DNA fragments	amplifiable or not amplifiable fragment	medium	dominant	no (rarely under)	limited	low/ moderate	yes
AFLP	1995	Random amplified DNA fragments	amplifiable or not amplifiable fragment	medium	dominant	no (rarely under)	limited	moderate / high	yes
microsatellites	end of 1980's	PCR amplification of a unique loci, harbouring simple sequences repeats	variation in the number of repeats	high	co-dominant	no	long time, high cost	moderate	yes

Mitochondrial markers



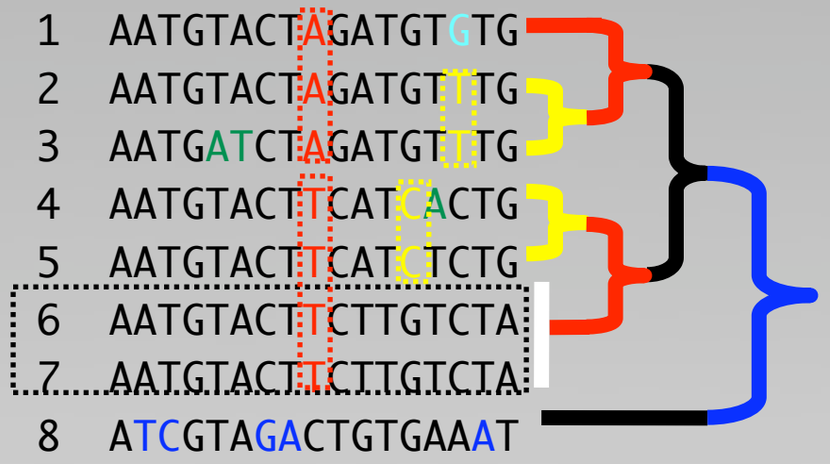
- numerous copies in a cell
- only maternal lineage / no (limited) heterozygosity
- animals: about 15-17k bp
  - well known: sequencing from defined primers
  - most interesting regions:
    - Control region (highly variable non-coding region: intra population → species)
    - cytochrome b (subspecies → genus)
    - NADH dehydrogenase 1-6 (subspecies → genus)
    - CO1 (species → order)
    - 12S / 16S (species → order)
- plants: 200k bp to >2400k bp
  - sequencing of some parts
  - presence of microsatellites in the mtDNA

Mitochondrial analyses

- only maternal lineage / no (limited) heterozygosity
- limited mutation rate: 1-10% / million of years
- methods used: SEQUENCING
- reconstruction of lineage, relationship between genus, species: PHYLOGENY
- relationship within a species, with implication of the geography e.g. PHYLOGEOGRAPHY: geographical distribution of genealogical lineages

Mitochondrial analyses: *phylogenetic trees*

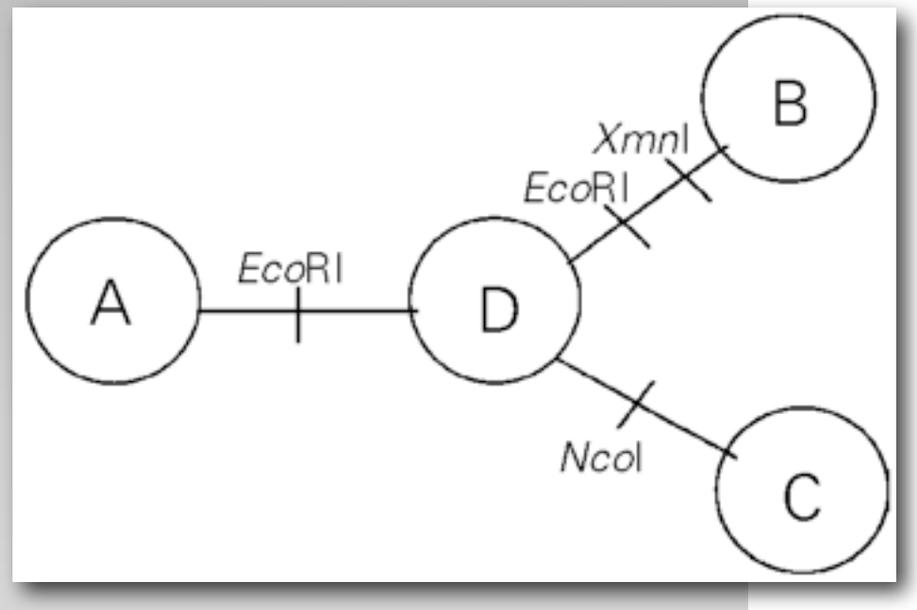
- regrouping most similar haplotypes

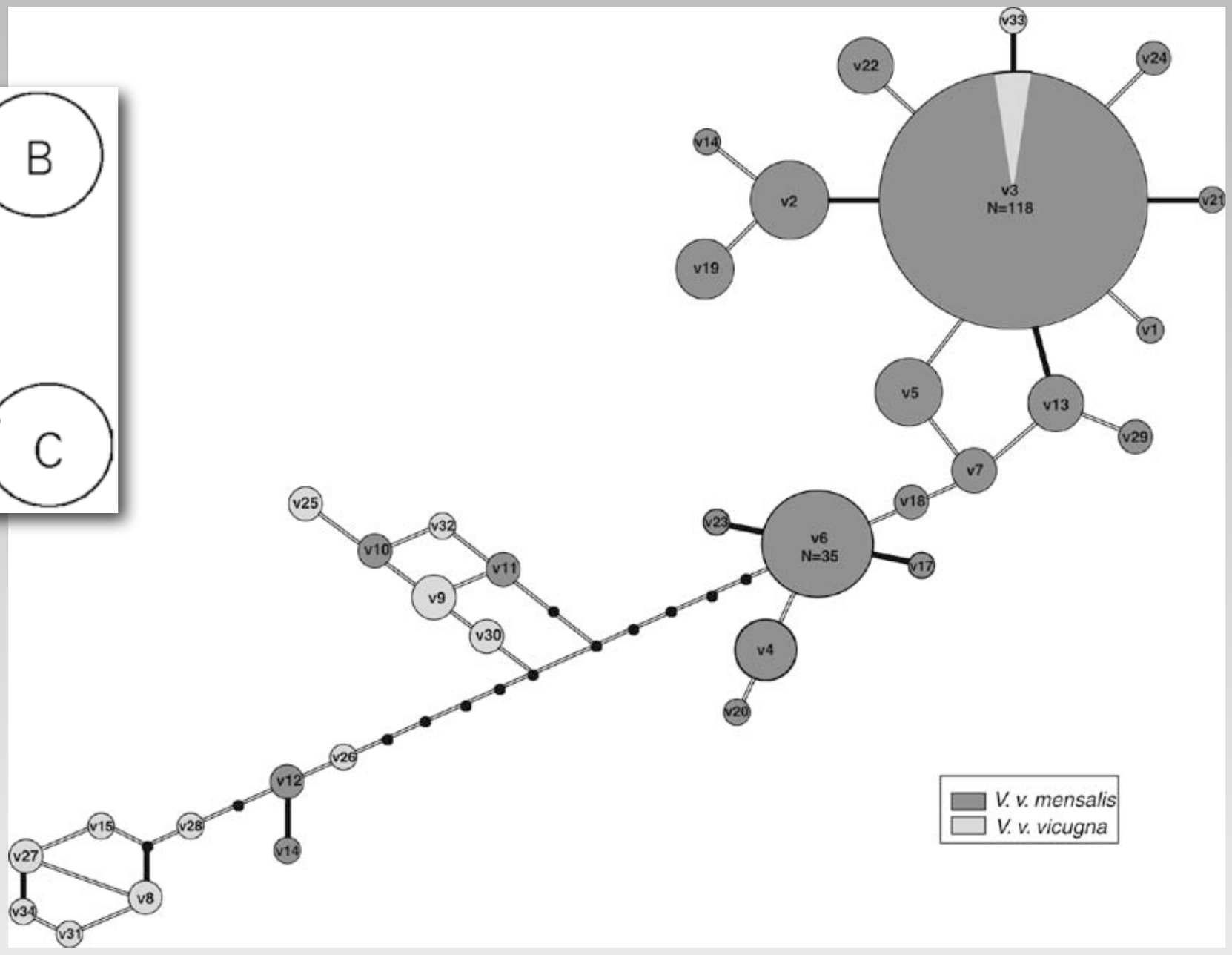


- different methods:
  - maximum likelihood: the tree with the lowest probability
  - maximum parsimony: the less number of steps (mutations)
  - genetic distance (Neighbour joining): regrouping most similar haplotypes
  - Bayesian method: posterior probability, after simulating and keeping the most probable trees

Mitochondrial analyses: network

- re-create all steps (mutation) between all haplotypes with a minimum steps





Nuclear markers

- paternal and maternal lineages: 2 copies ⇒ heterozygosity
- mutation rate:
  very low (e. g. coding region) to very high (e. g. microsatellites)
- use for
  - pedigree reconstruction (maternal-paternal lineages)
  - level of inbreeding
  - population differentiation
  - migration estimation
  - differentiated behaviour (migration, ...) between sexes
  - ...

Nuclear markers: *some definitions*

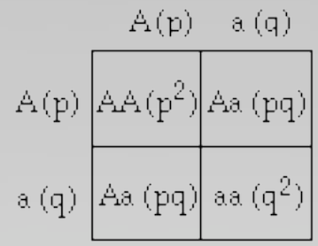
- **Locus**: a segment of DNA, e.g. a microsatellites, coding for a protein, ...
- **Alleles**: different forms of the same locus, e.g. different length of a microsatellite, different amino-acidic chain in a protein, ...
- **Heterozygote**: an individual with two different allele at a locus e.g. alleles A_1A_2 for the locus A
- **Average heterozygosity**: mean of the heterozygosity at all loci
- **Allelic diversity**: average number of alleles per locus

Nuclear markers

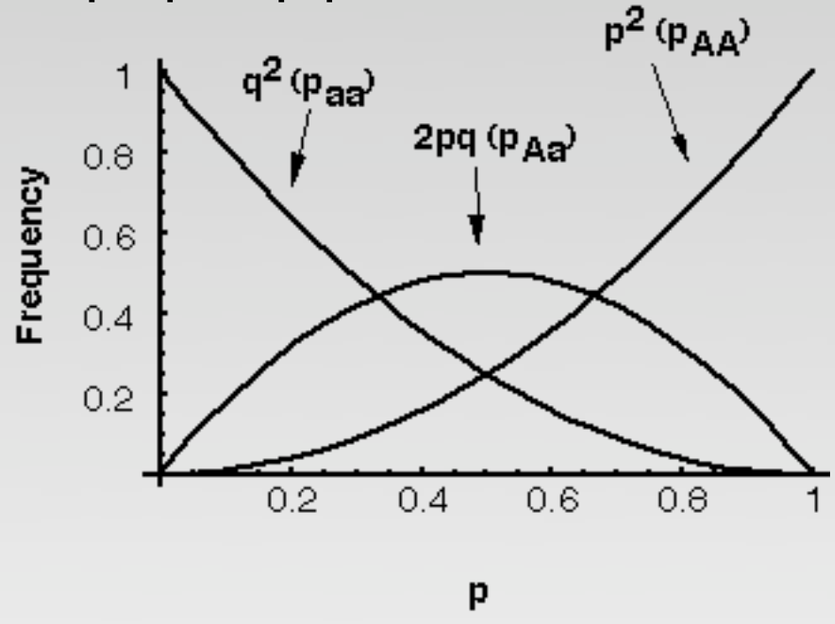
- markers used
  - microsatellites
    - when microsatellites already developed
    - no limitation by cost
    - more for animals (sometimes difficult to find in plants)
    - neutral markers
  - AFLP
    - when no microsatellites already exist and cannot been developed (time, cost)
    - plants
    - dominance is not a problem
  - RFLP
    - when no microsatellites already exist and cannot been developed (time, cost)
    - limit the cost
    - plants
    - dominance is not a problem
  - RAPD, enzymes, sequencing, SSCP, fingerprints, ...
    - particular cases

Nuclear marker analyses: *Hardy-Weinberg (HW) equilibrium*

- in large population, with random mating and no mutation, migration or selection
- allele and genotypes frequencies in equilibrium
- e.g. locus with alleles A_1 and A_2, relative frequency of p and q, where p+q=1



- proportion of A_1A_1: (♀p - ♂p) pxp = p^2
- proportion of A_2A_2: (♀q - ♂q) qxq = q^2
- proportion of A_1A2: (♀p - ♂q AND ♀q - ♂p) 2* pxq = 2pq



Nuclear marker analyses: *genetic diversity characteristics*

- expected heterozygosity (gene diversity): H_E
  - for p and q allele frequency: $H_E = 2pq$
  - for more alleles: $He = 1-\sum p_i^2$ for all alleles frequencies
- observed heterozygosity: H_O
  - proportion of heterozygotes at a locus
- allelic richness: A (or A_R)
  - average number of alleles per locus

Nuclear marker analyses: *genetic diversity characteristics*

- example 1

	AA	AB	BB	total
number	27	23	5	55
genotype frequency	0.49	0.42	0.09	1.0

- estimation of alleles frequency:

  p = [(2*27)+(1*23)] / [2*55] = 0.70

  q = [(2*5)+(1*23)] / [2*55] = 0.30

  p + q = 0.70 + 0.30 = 1

- expected heterozygosity: H_E

  $He = 1-\sum p_i^2 = 1 - [0.70^2 + 0.30^2] = 1-[0.49+0.09] = 1-0.58 = 0.42$

- observed heterozygosity: H_O

  no heterozygotes / total number = 23 / 55 = 0.42

- allelic richness: A (or A_R)

  average number of alleles per locus = 2

Nuclear marker analyses: *genetic diversity characteristics*

- example 2

	91/91	91/95	91/97	95/95	95/97	97/97	total
number	10	24	6	23	9	8	80
genotype frequency	0.125	0.30	0.075	0.2875	0.1125	0.10	1.0

- estimation of alleles frequency:
  - p = [(2*10)+(1*24)+(1*6)] / [2*80] = 0.312
  - q = [(2*23)+ (1*24)+(1*9)] / [2*80] = 0.494
  - r = [(2*8)+ (1*6)+(1*9)] / [2*80] = 0.194
  - p + q +r = 0.312 + 0.494 + 0.194 = 1
- expected heterozygosity: H_E

  $He = 1-\sum p_i^2 = 1 - [0.312^2 + 0.494^2 + 0.194^2] = 1-0.38 = 0.62$
- observed heterozygosity: H_O

  no heterozygotes / total number = 24 + 6 +9 / 80 = 0.49
- allelic richness: A (or A_R)

  average number of alleles per locus = 3

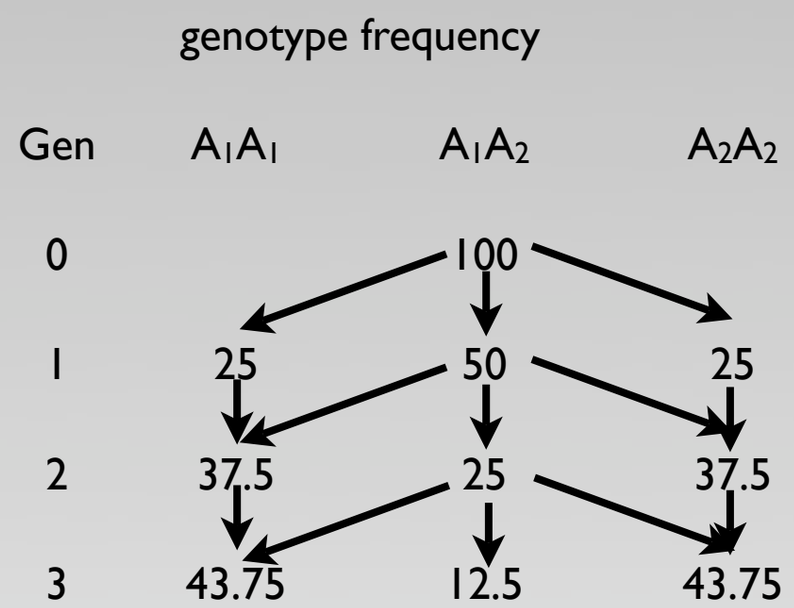
Nuclear marker analyses: *Deviations from Hardy-Weinberg (HW) equilibrium*

- causes
  - inbreeding
  - assortative and disassortative mating
  - fragmented populations

Nuclear marker analyses: *Deviations from Hardy-Weinberg (HW) equilibrium*

- causes
  - inbreeding
    - definition: mating with relatives
    - with inbreeding: decrease of heterozygotes (compare to HW equilibrium)

      e.g.: selfing



  - assortative and disassortative mating
  - fragmented populations

Nuclear marker analyses: *Deviations from Hardy-Weinberg (HW) equilibrium*

- causes
  - inbreeding
  - assortative and disassortative mating
    - preferential selection of mate with similar (assortative) or different (disassortative) genotype

      e.g.: human female selection:

      disassortative odour preferences in human (Wedekind et al., 1995; Wedekind & Furi 1997; Thornhill et al. 2003) ➡ disassortative

      MHC-disassortative mating observed between partners (Ober et al., 1997)

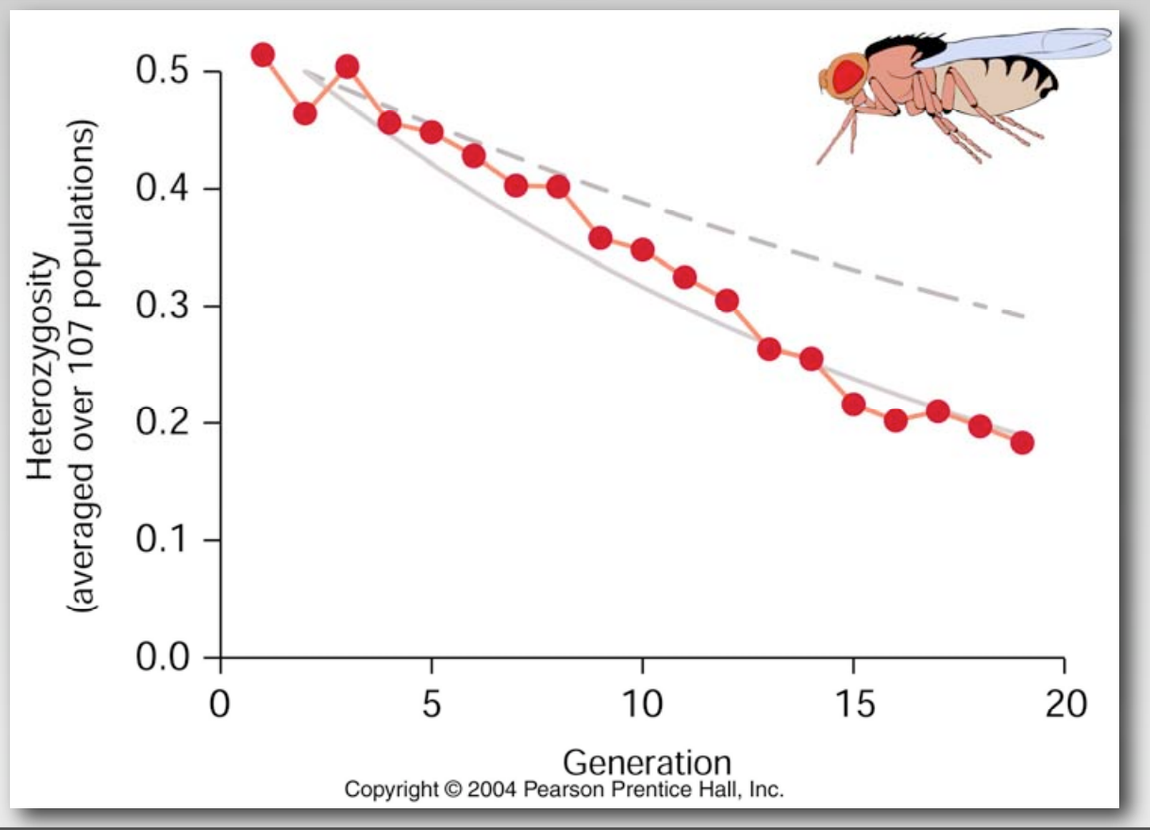
      BUT: MHC-similar facial preferences
  - fragmented populations

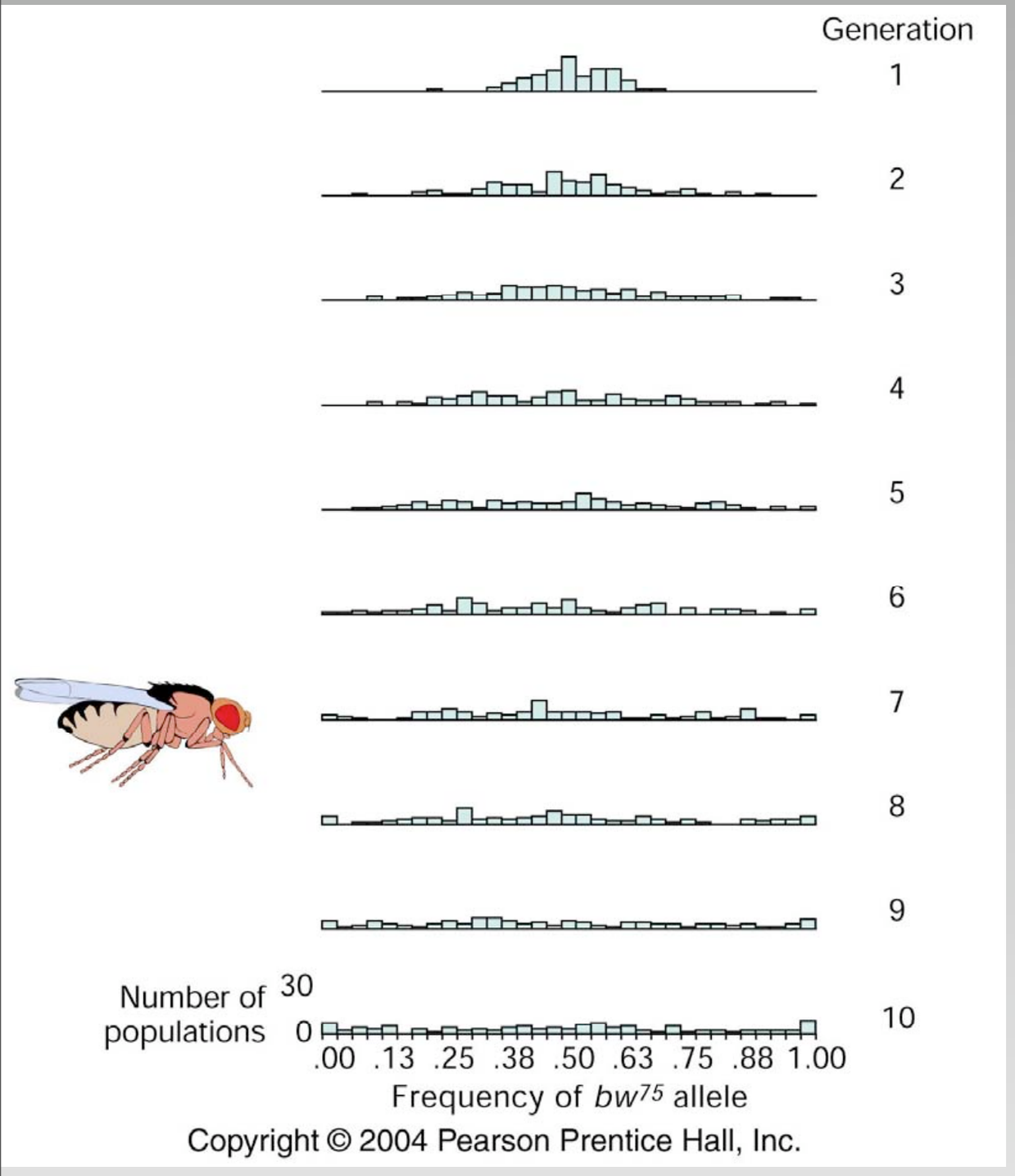
MHC (Major histocompatibility Complex): is a large genomic region or gene family found in most vertebrates. It plays an important role in the immune system, autoimmunity, and reproductive success.

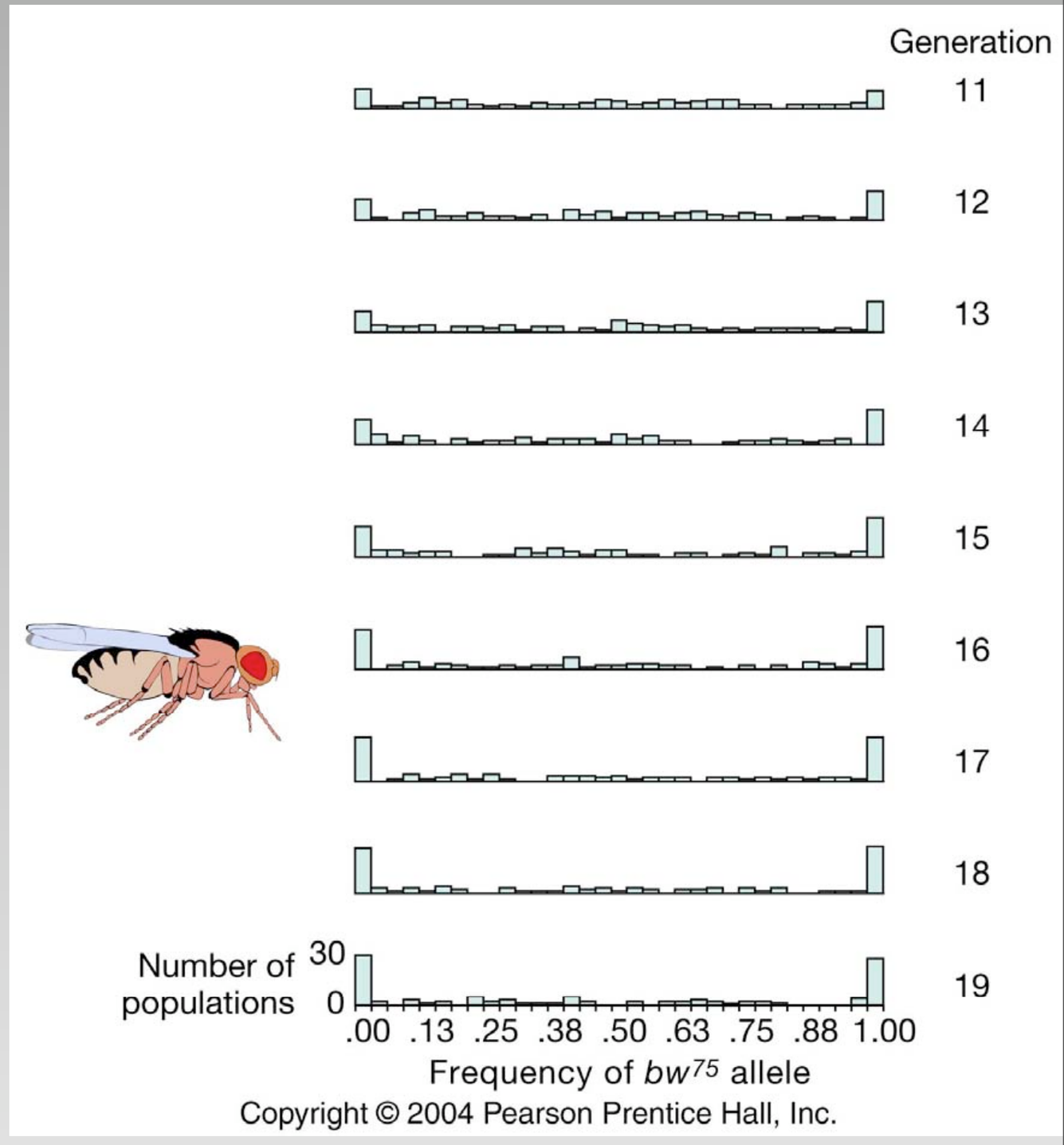
Nuclear marker analyses: *Deviations from Hardy-Weinberg (HW) equilibrium*

- causes
  - inbreeding
  - assortative and disassortative mating
  - fragmented populations
    - small isolated population fragments will differentiate at random due to genetic drift

      e. g. Buri 1956: evolution of heterozygosity in bw^{75} allele over 19 generations in 105 replicate populations maintained with 16 parents per generations



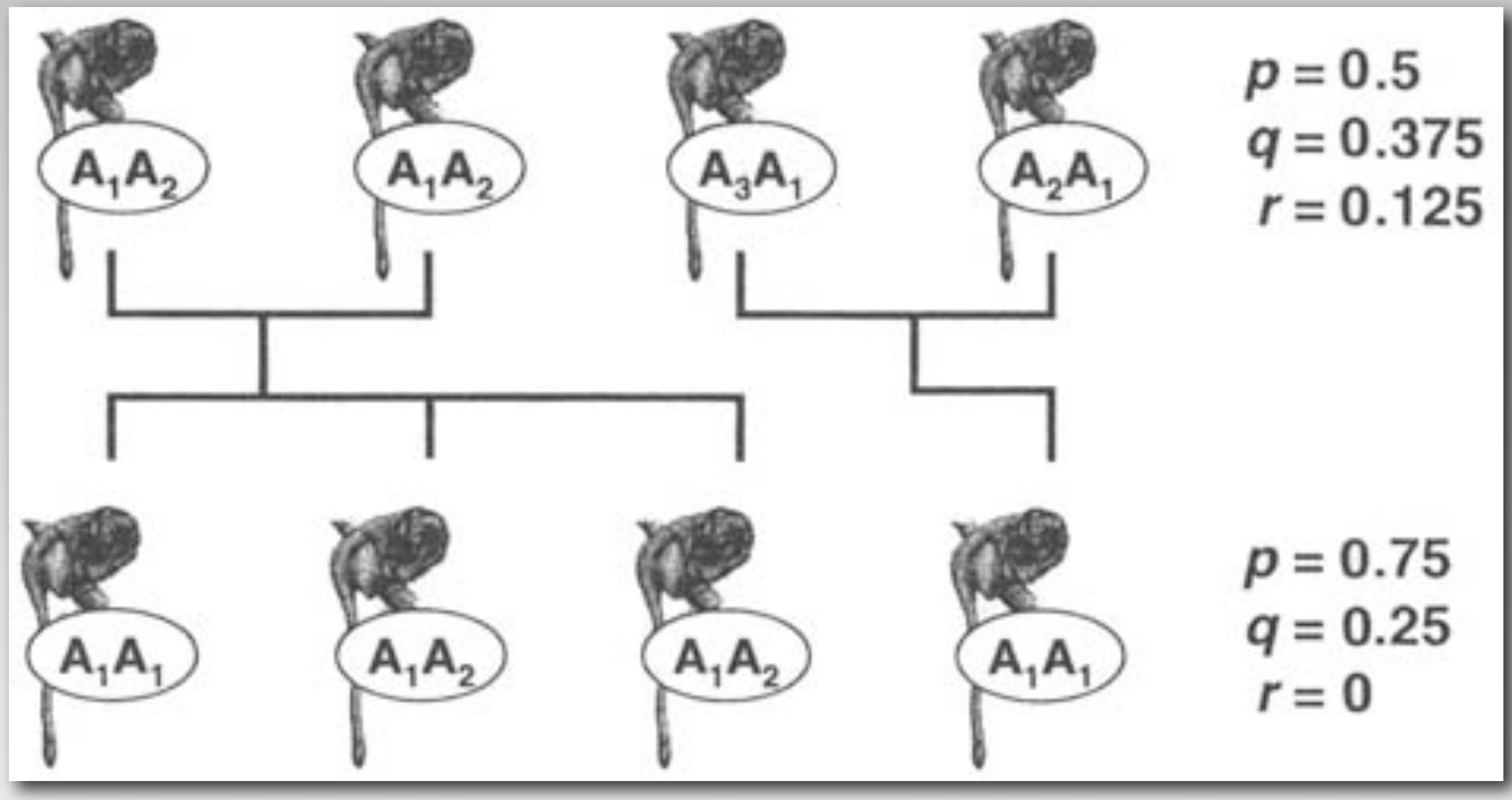




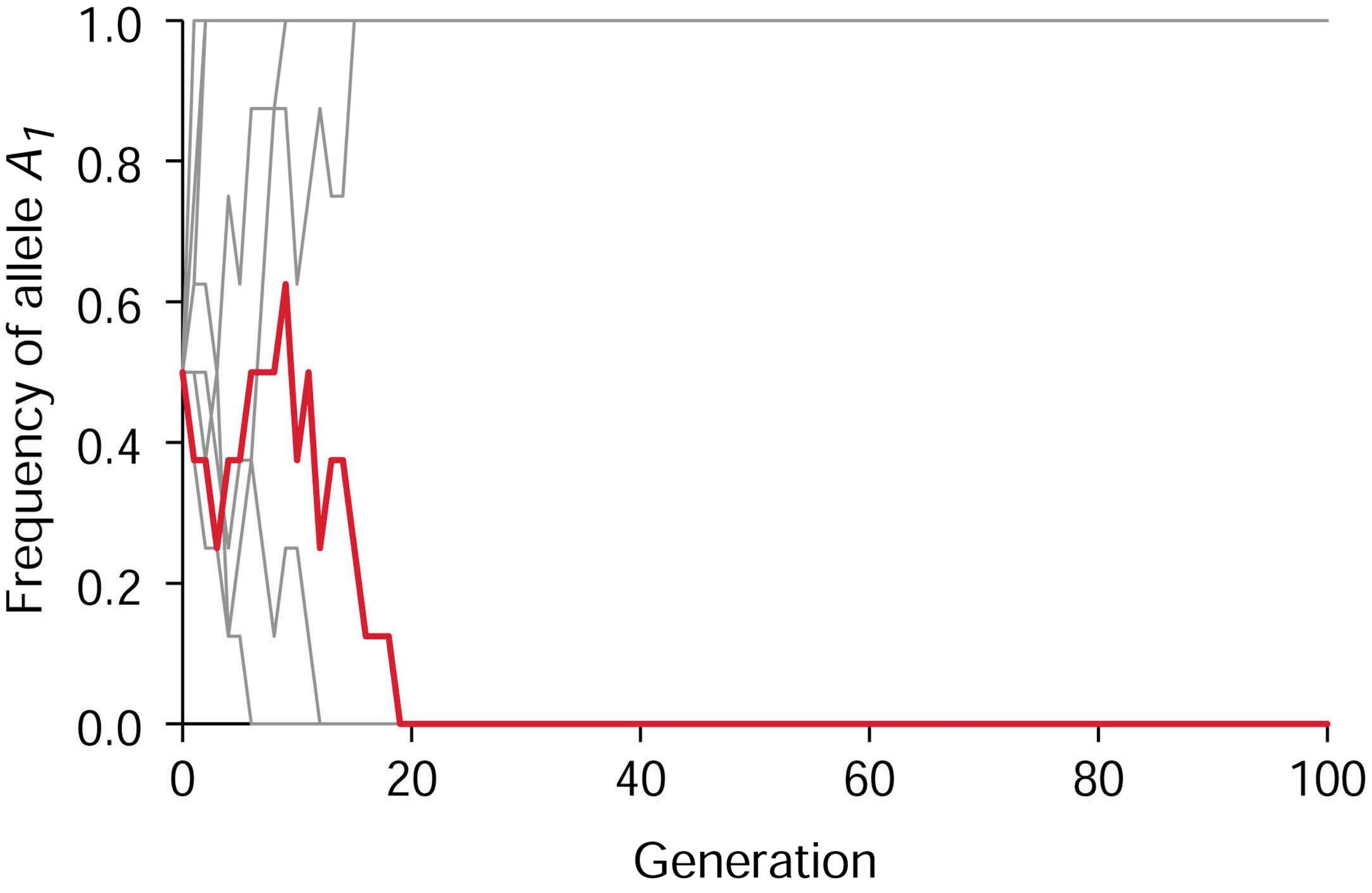
Buri, 1956: frequency distribution of the bw^{75} allele over 19 generations in 105 replicate populations maintained with 16 parents per generations

Small population problems: *impact of the population size on the genetic diversity*

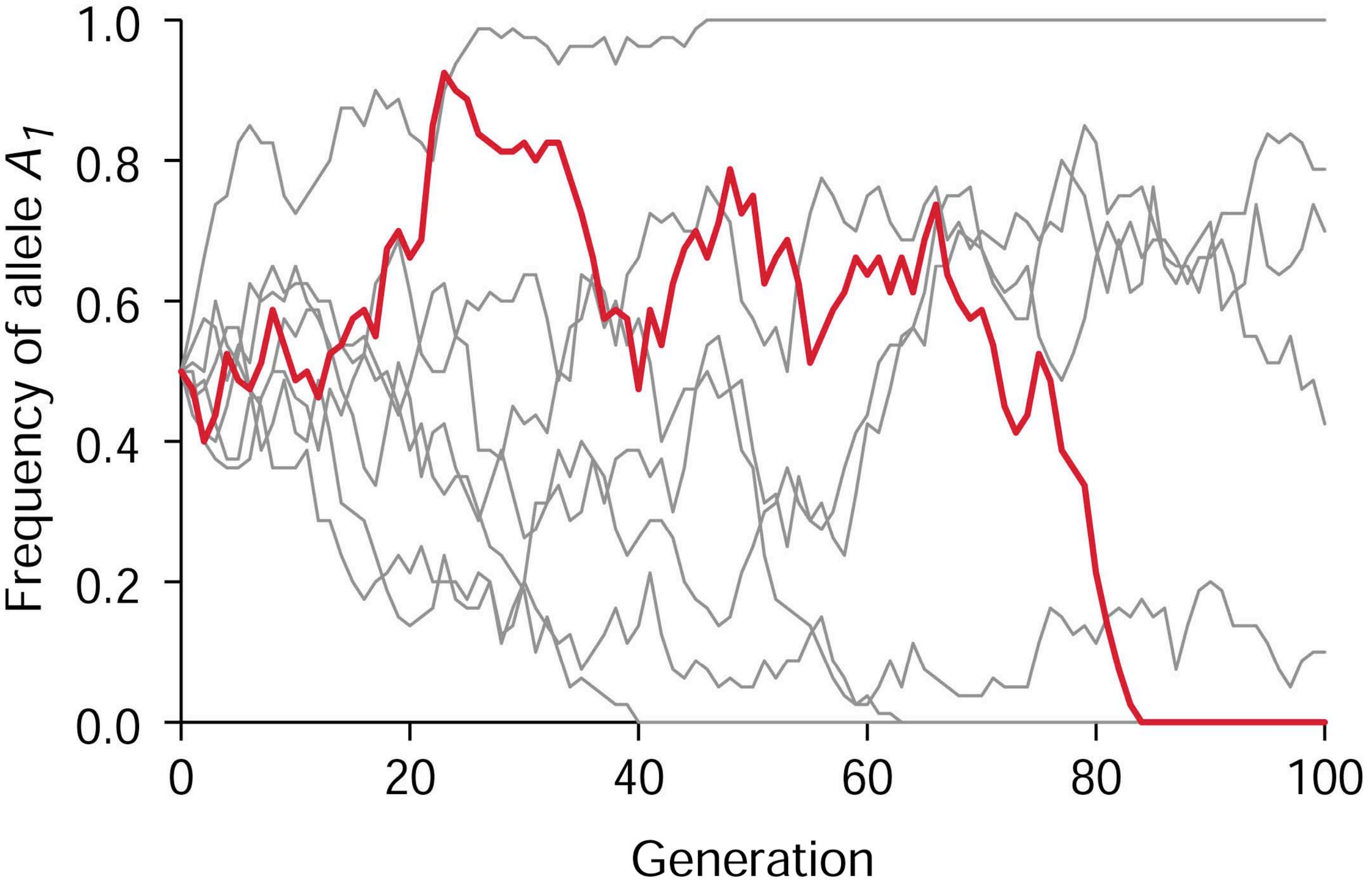
- stochasticity
  - just by chance, some alleles (especially the rare ones) may not be passed to the next generation and are consequently lost.
    ➡ frequency of alleles change over generation



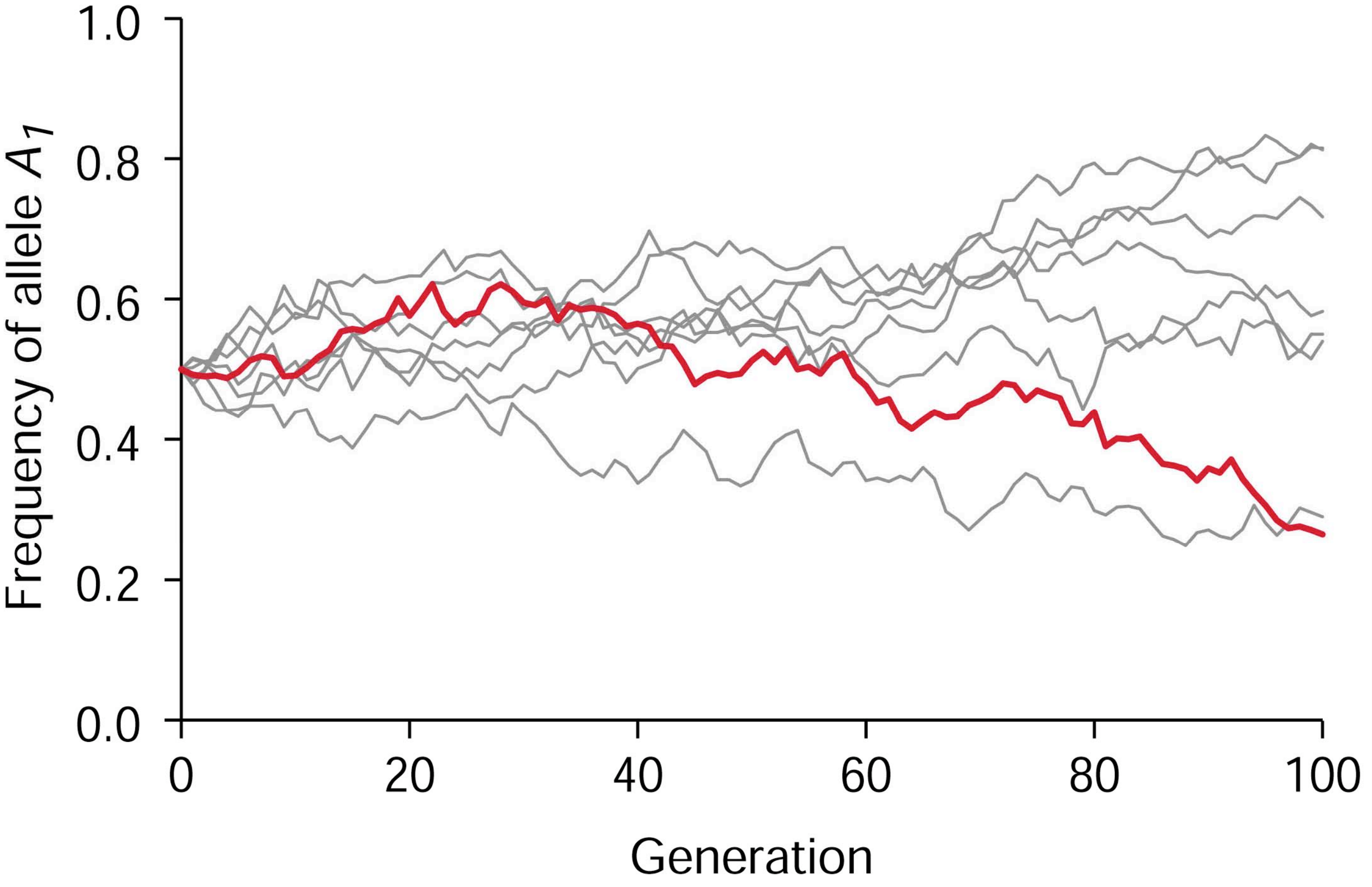
(a) Population size = 4



(b) Population size = 40

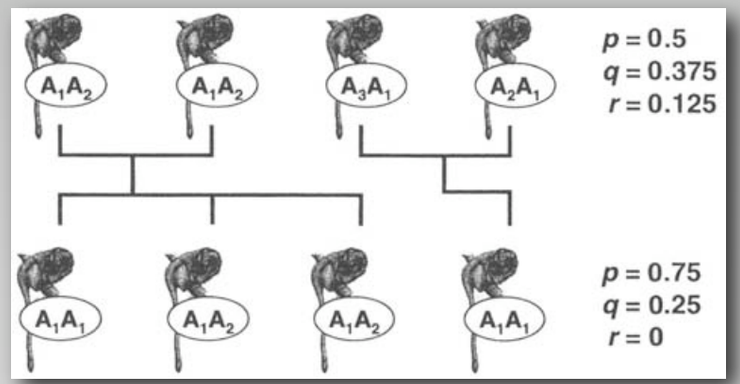


(c) Population size = 400



Small population problems: *impact of the population size on the genetic diversity*

- stochasticity
  - just by chance, some alleles (especially the rare ones) may not be passed to the next generation and are consequently lost.
    ➡ frequency of alleles change over generation



  - genetic drift: allele frequency change over generation, with a general reduction of the global genetic diversity

    consequences:
    - random changes in allele frequencies from one generation to the next one
    - loss of genetic diversity and fixation of alleles within populations
    - diversification among replicate population from the same original sources (e. g. fragmented populations)

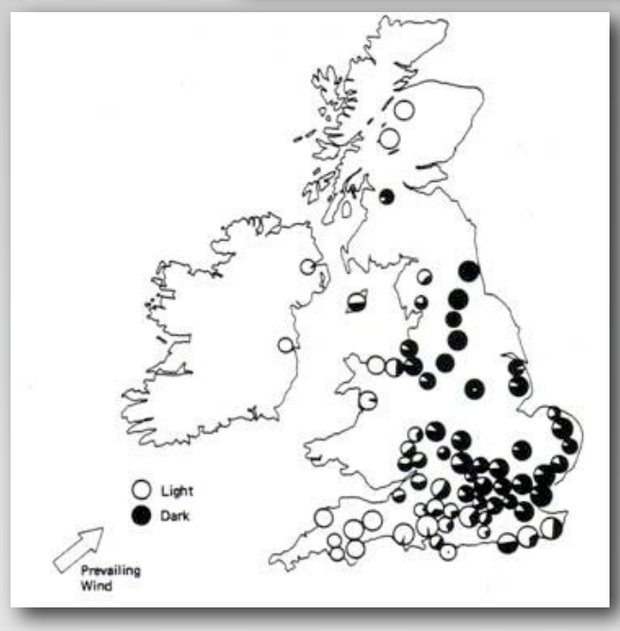
Small population problems: *lost of genetic diversity*

- reasons of the lost of genetic diversity in small populations
  - genetic drift
  - inbreeding reducing heterozygosity
  - selection reducing genetic diversity by favouring one allele at the expense of another ➡ fixation
- impact:
  - reduce the ability to evolve in response to environmental changes

    e.g.: peppered moth in UK / resistance to myxoma virus in Australian rabbits

Introduction: *Why genetic diversity is important in populations...*

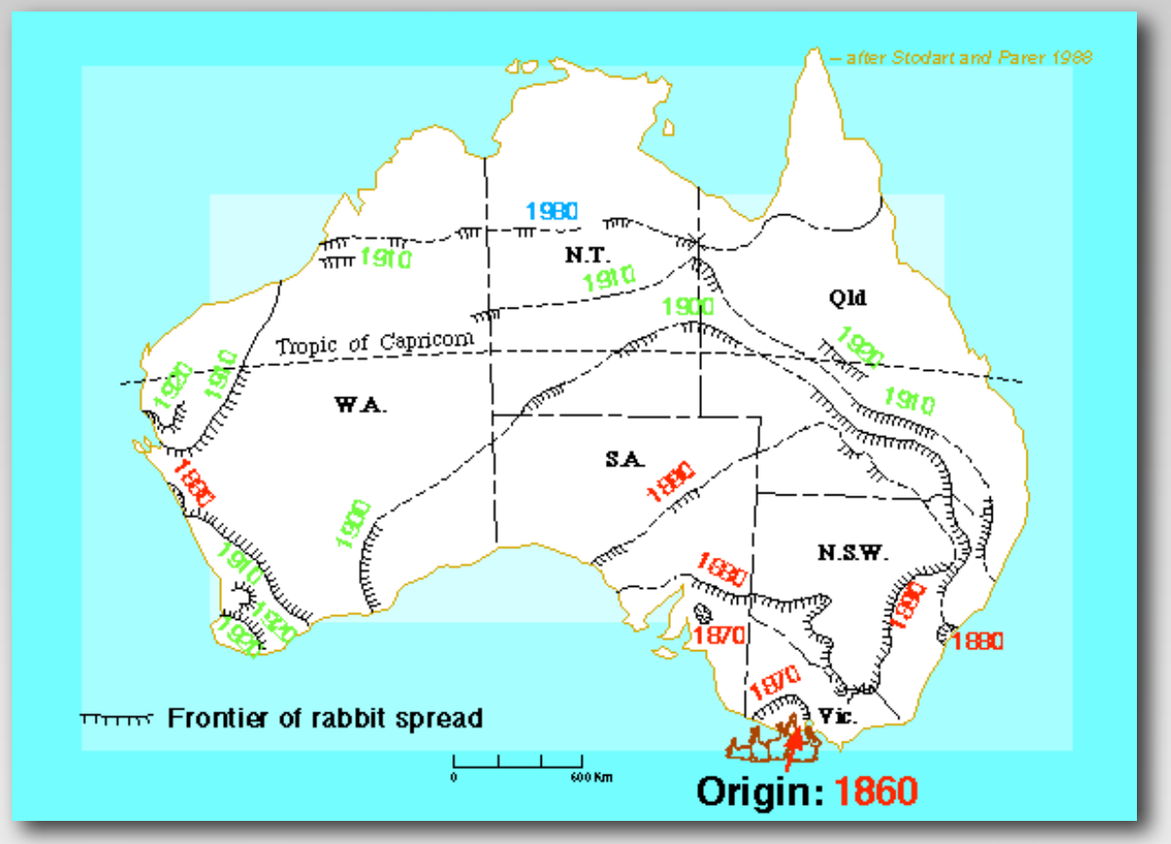
- genetic diversity reflect evolutionary potential
  - example 1 - habitat selection: peppered moth in UK
    - dark and light forms
    - night: active / day: resting on trees
    - ➡ camouflage critical for survival
    - light form: camouflaged on lichen-covered tree trunks
    - Industrialisation: kill lichen by sulphur pollution
    - ➡ light form: visible / dark form: camouflaged

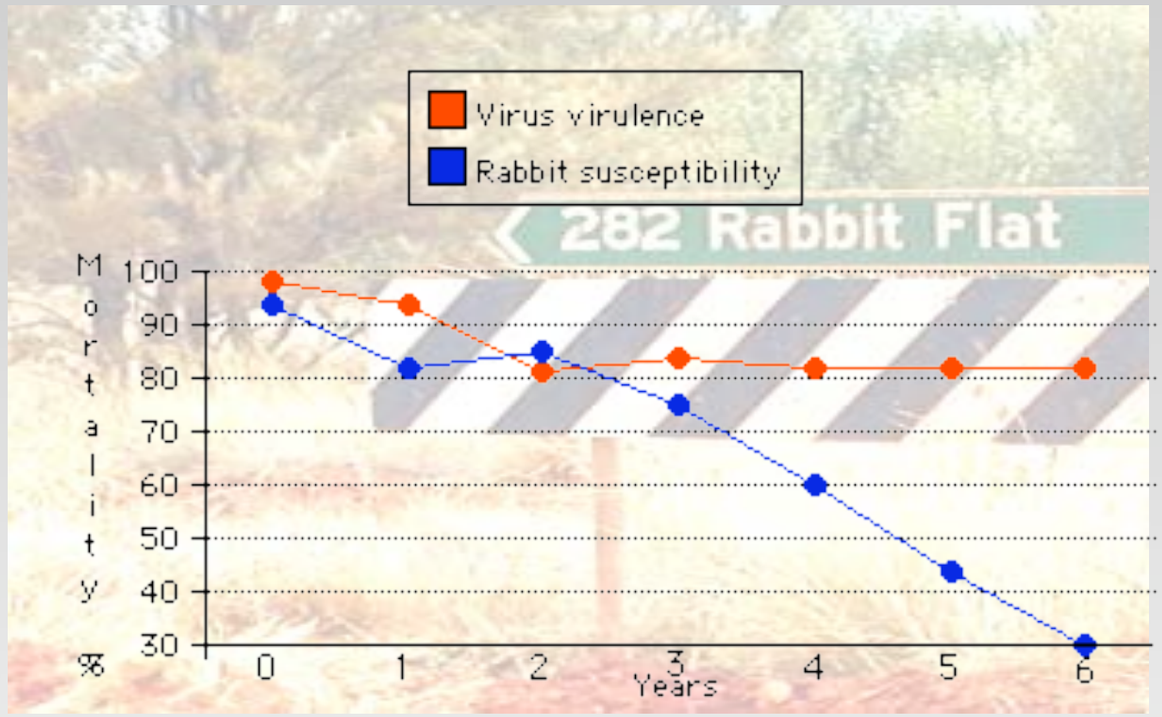


Grant (1999) Fine tuning the peppered moth paradigm, Evolution 53, 980-984
Kettlewell (1973) The Evolution of Melanism, Clarendon Press, Oxford, UK
Majerus (1998) Melanism: Evolution in Action. Oxford University Press
Kettlewell (1958) A survey of the frequencies of *Biston betularia* (L.) (Lep.) and its melanic forms in Great Britain, Heredity 12, 551-572
but see also: Rudge (2006) Myths about moths: a study in contrasts, Endeavour 30, 19-23

Introduction: *Why genetic diversity is important in populations...*

- genetic diversity reflect evolutionary potential
  - example 2 - disease resistance: resistance to myxoma virus in Australian rabbits
    - introduction of rabbits in Australia: 1860
    - control measure: introduction of myxoma in 50'
    - ➡ high mortality rate first years
    - high selection for resistance





Small population problems: *lost of genetic diversity*

- reasons of the lost of genetic diversity in small populations
  - genetic drift
  - inbreeding reducing heterozygosity
  - selection reducing genetic diversity by favouring one allele at the expense of other ➡ fixation
- impact:
  - reduce the ability to evolve in response to environmental changes
  - reduce the fitness

Analysis of the relationship between allozyme heterozygosity and fitness in the rare Gentiana pneumonanthe L.

Oostermeijer et al. (1995) **J. Evol. Biol.** 8: 739-759 (1995)

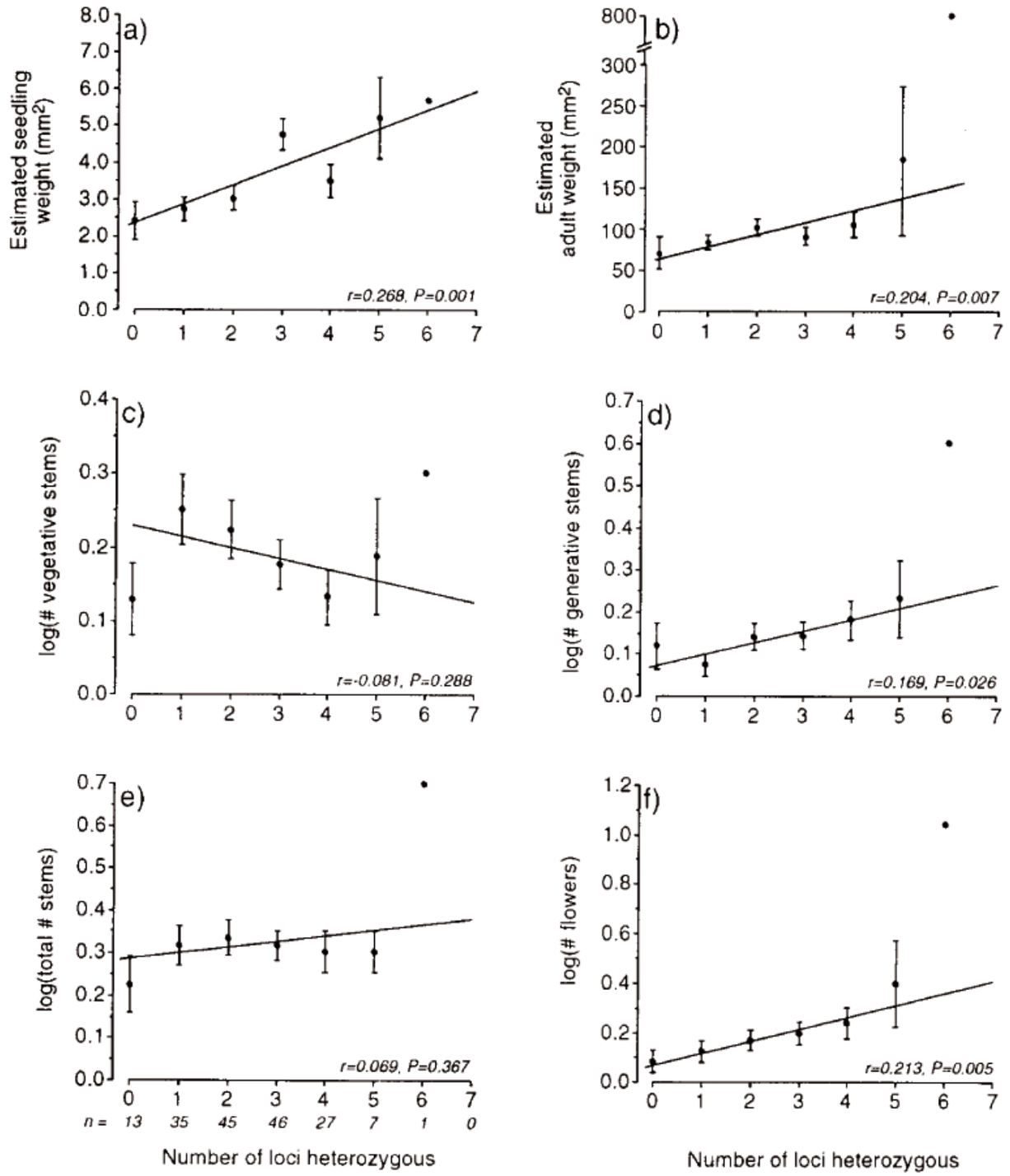
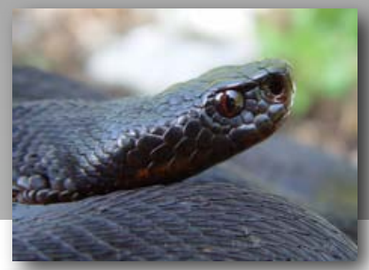


Fig. 1. Relationship between the number of heterozygous loci per individual (out of seven assayed polymorphic loci) and six components of individual fitness, (a) seedling weight, (b) adult weight, (c) number of vegetative stems, (d) number of generative stems, (e) total number of stems, and (f) number of flowers. Note that parameters (c) to (f) have been ln-transformed. Regression lines are based on values of individual plants and not on the class means shown in these graphs with their standard errors. In the right hand corner of each graph, the correlation coefficient (r) and its probability (P) is given. Below graph (e) the number of individuals per heterozygosity class (n) is shown. Only one individual was heterozygous for 6 of the seven loci (hence this class has no standard error), and none were heterozygous for all seven.

Small population problems: *lost of genetic diversity*

- reasons of the lost of genetic diversity in small populations
  - genetic drift
  - inbreeding reducing heterozygosity
  - selection reducing genetic diversity by favouring one allele at the expense of another ➡ fixation
- impact:
  - reduce the ability to evolve in response to environmental changes
  - reduce the fitness
- consequences:
  - extinction of alleles
  - extinction of populations or species
    - e.g. Madsen et al. (1996, 1999, 2004) - near extinction



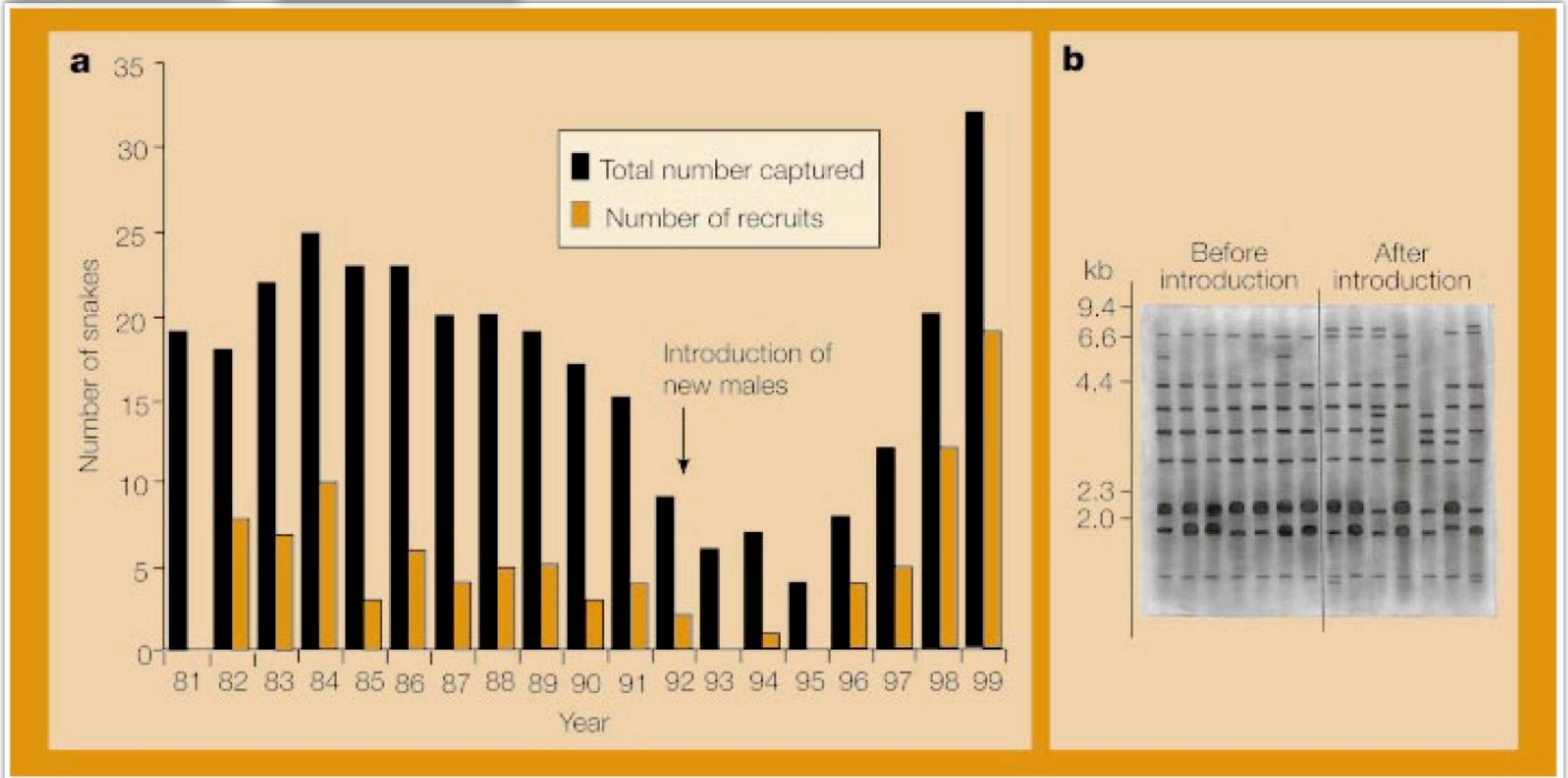
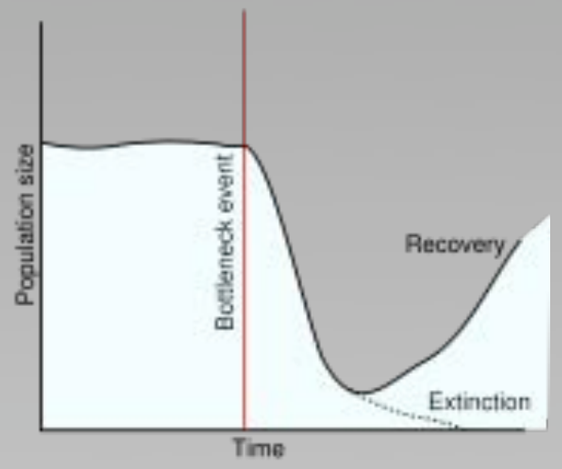


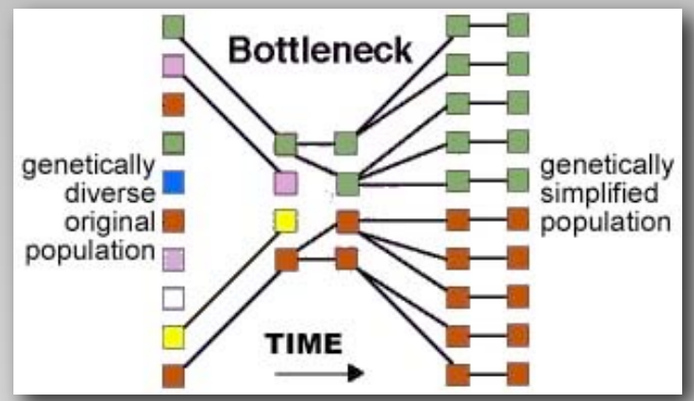
Figure 1 Introducing new males increases the genetic diversity and enables the adder population to recover. **a,** Total number and number of recruited male adders captured in Smygehuk from 1981 to 1999. **b,** Southern-blot analysis of major histocompatability complex (MHC) class I genes in seven males sampled before the introduction of new males (left) and in seven recruited males sampled in 1999 (right).

Madsen et al. (1999) Restoration of an inbred adder population, Nature 402, 34-35

Small population problems: *bottleneck*



- bottleneck: large reduction of N_e in a period of time
  - consequence: lost of genetic diversity, especially rare alleles
  - impact depends on the population size during the bottleneck and the duration of it (nb generation)
  - e.g.: northern elephant seal (*Mirounga angustirostris*)
    - large reduction of the population size due to hunting
    - 20-30 survived in Isla Guadalupe (probably only a single harem)
    - mtDNA:
      - before 1892: ≥4 haplotypes (only 5 samples)
      - after 1892: only 2 haplotypes (>150 samples)
    - 20 allozymes:
      - no diversity in the northern elephant seal
      - normal level for the southern elephant seal (*Mirounga leonina*)

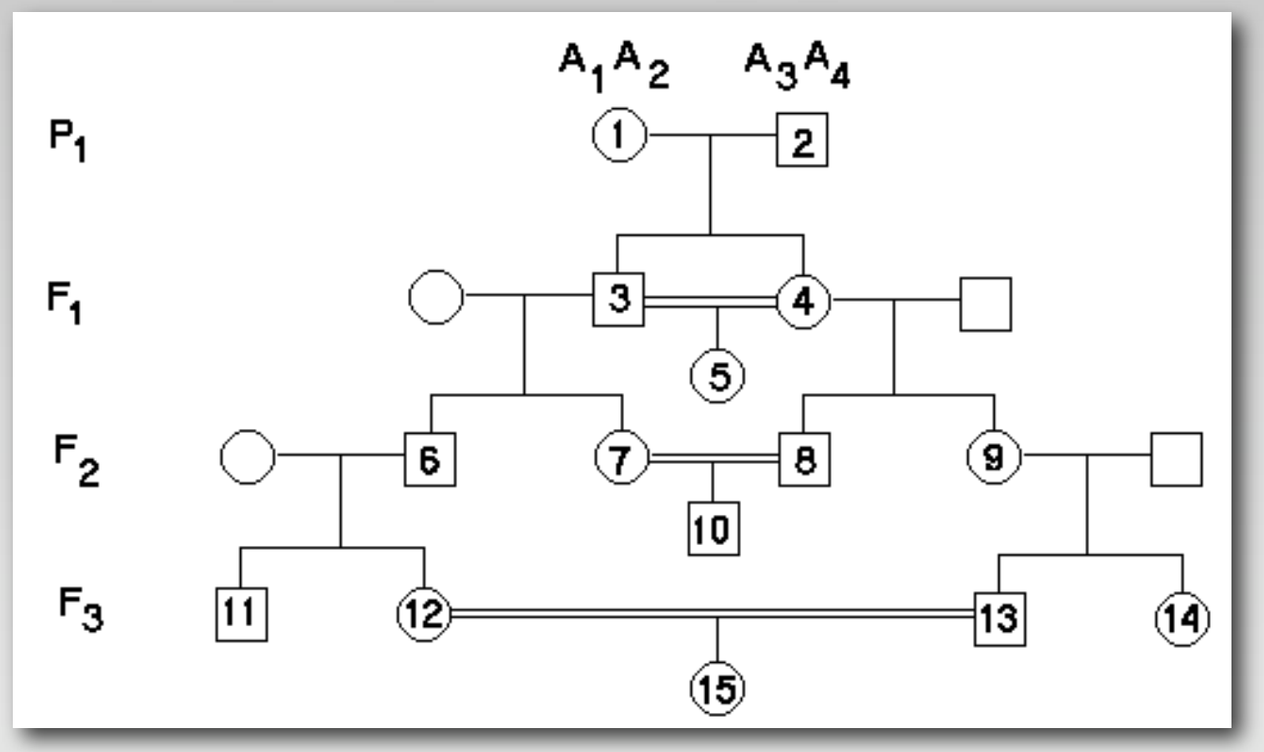


Small population problems: *Inbreeding estimations*

regarder l'estimation de r et F

- inbreeding: mating of individuals related by ancestry measured as the probability that two alleles at a locus ar by descent (*F*). Recent copies of the same allele are refer identical by descent, or autozygote

- also named as pedigree inbreeding



Relationship	***Description***	***Example***	***r***	***F of offspring***
Parent / Offspring	mother or father, to son or daughter	2 & 4	1/2	1/2
Full sibs	offspring of same parents	3 & 4	1/2	1/4
Half sibs	offspring with one parent in common	not shown	1/4	1/8
1st cousins	offspring of full sibs	7 & 8	1/4	1/16
2nd cousins	offspring of 1st cousins	12 & 13	1/8	1/64

Small population problems: *Theory of inbreeding in small populations*

in an hermaphroditic species

N = nb individuals

2N ancestral alleles

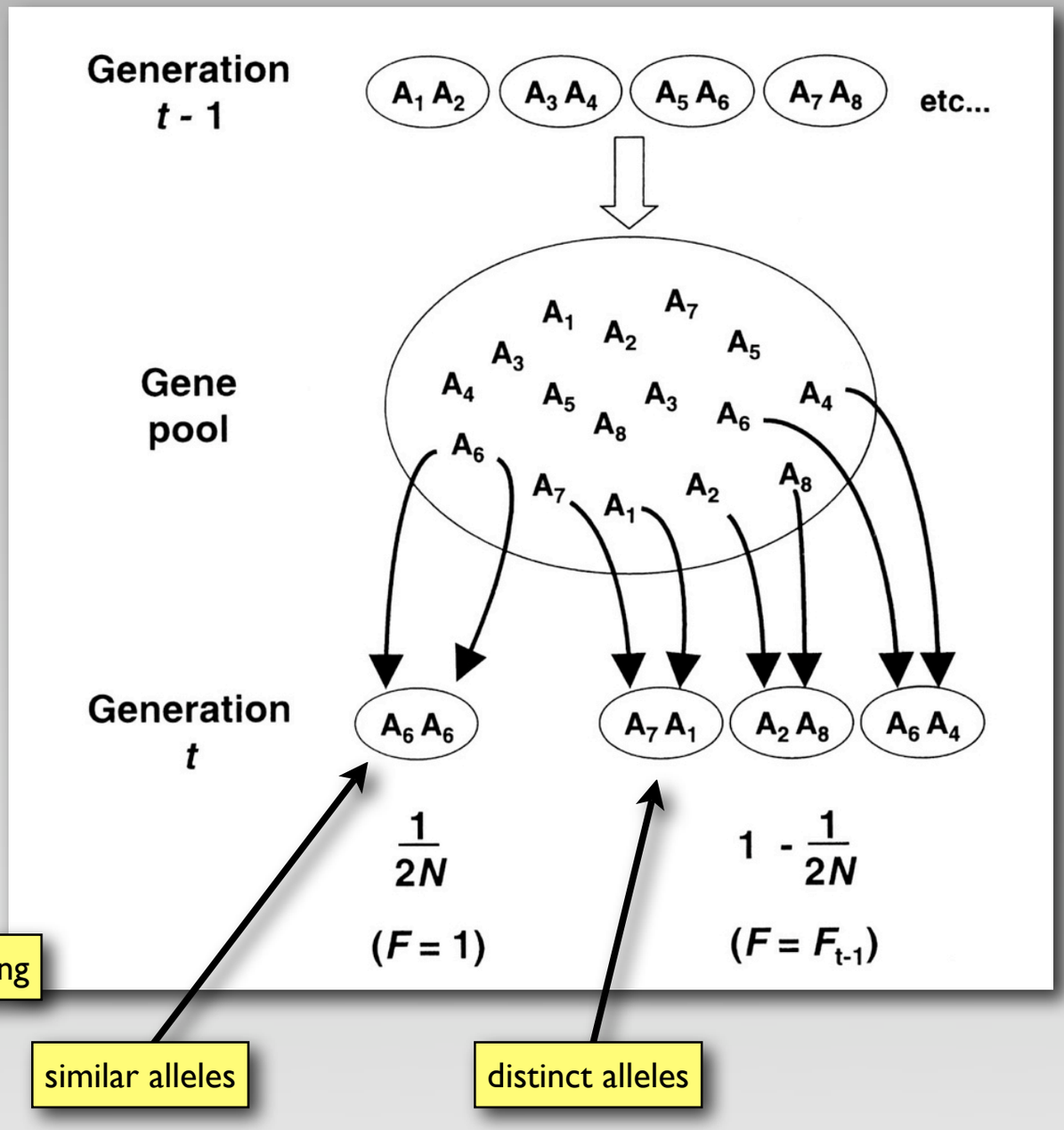
each individual at t: randomly sampling with replacement of two alleles

e.g. A6 first sampled:
prob. that the second is A6 for 1 individual:
= 1/2N

probability of sampling distinct alleles:
= 1 - 1/2N

➡ probability of creating a zygote with both alleles identical by descendent (F_t):
$F_t = 1/2N + [1-1/2N]F_{t-1}$ ← previous inbreeding

➡ increase of inbreeding per generation:
$\Delta F = 1/(2N)$



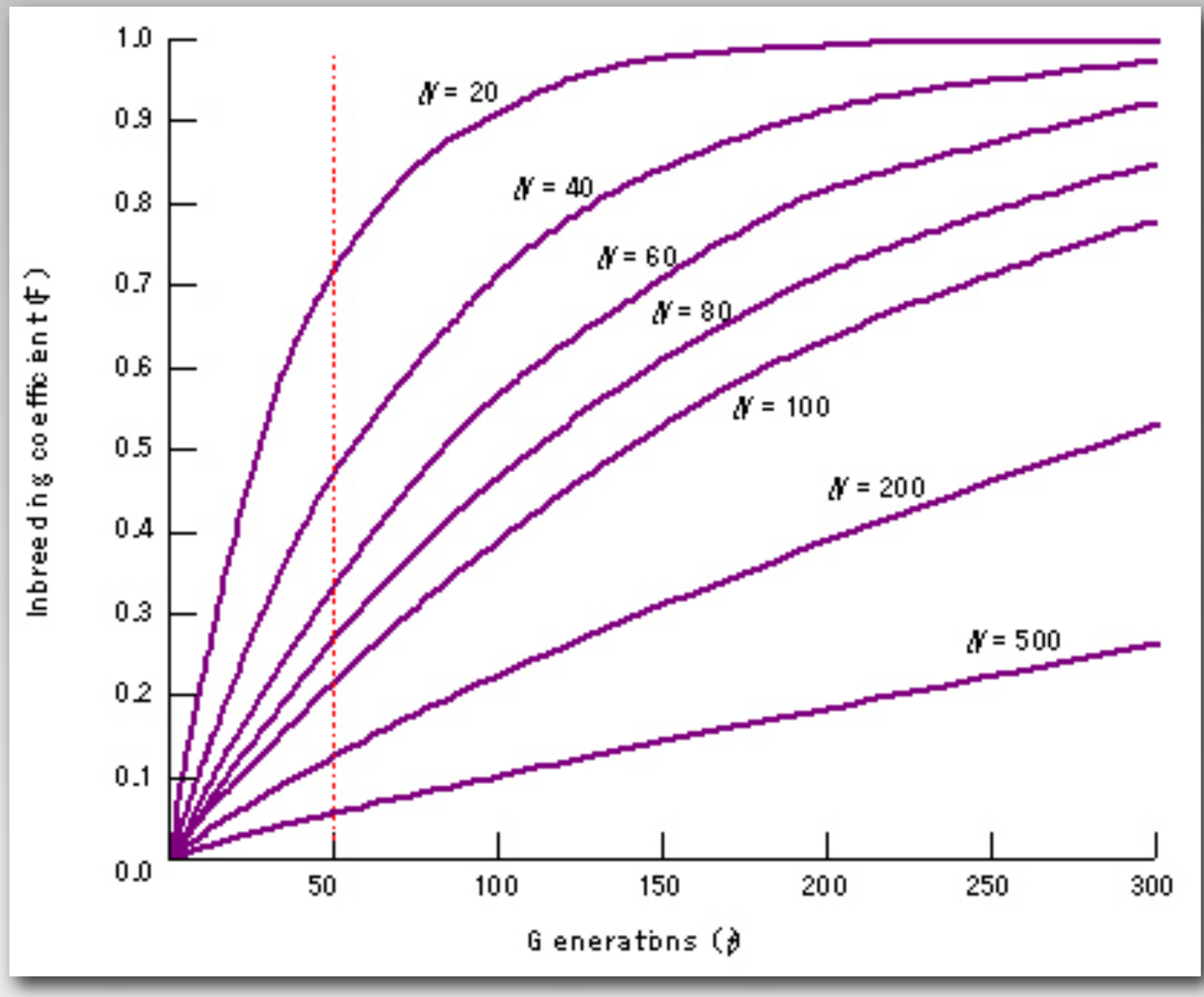
Small population problems: *Theory of inbreeding in small populations*

➡ probability of creating a zygote with both alleles identical by descendent (F_t):

$F_t = 1/2N + [1-1/2N]F_{t-1}$

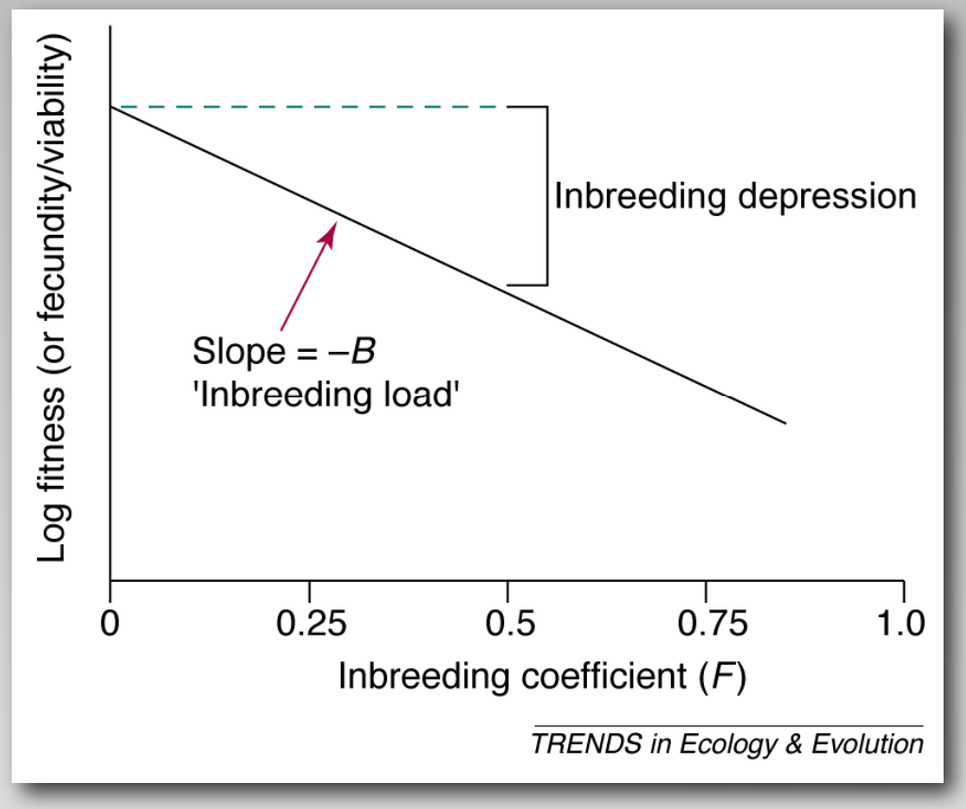
➡ increase of inbreeding per generation:

$\Delta F = 1/(2N)$



Small population problems: *Inbreeding depression*

- population size reduction increase inbreeding rate in closed populations ➡ inbreeding results in a decline of the global fitness, named as inbreeding depression



- purging
  - elimination due to a strong negative selection on rare deleterious recessive alleles
    purging highly effective for alleles with large effects (e. g. lethal)

Small population problems: *Inbreeding depression*

Charpentier et al. (2006), Life history correlates of inbreeding depression in mandrills (*Mandrillus sphinx*), **Molecular Ecology** 15:21-28

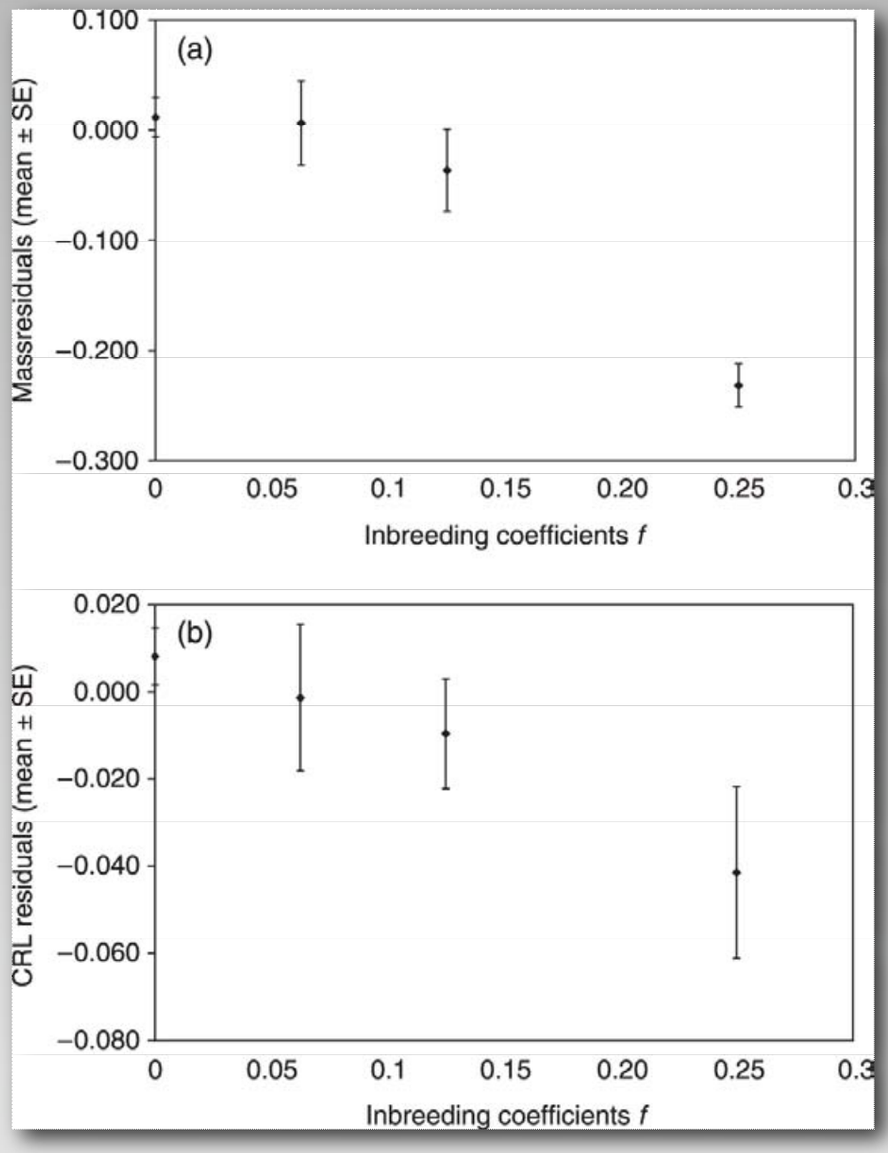


Fig. 1 Relationship between inbreeding coefficients and growth in females. Figures show mean ± SE for each inbreeding value. (a) Mass-for-age; (b) Crown-rump length -for-age (= embryos length)

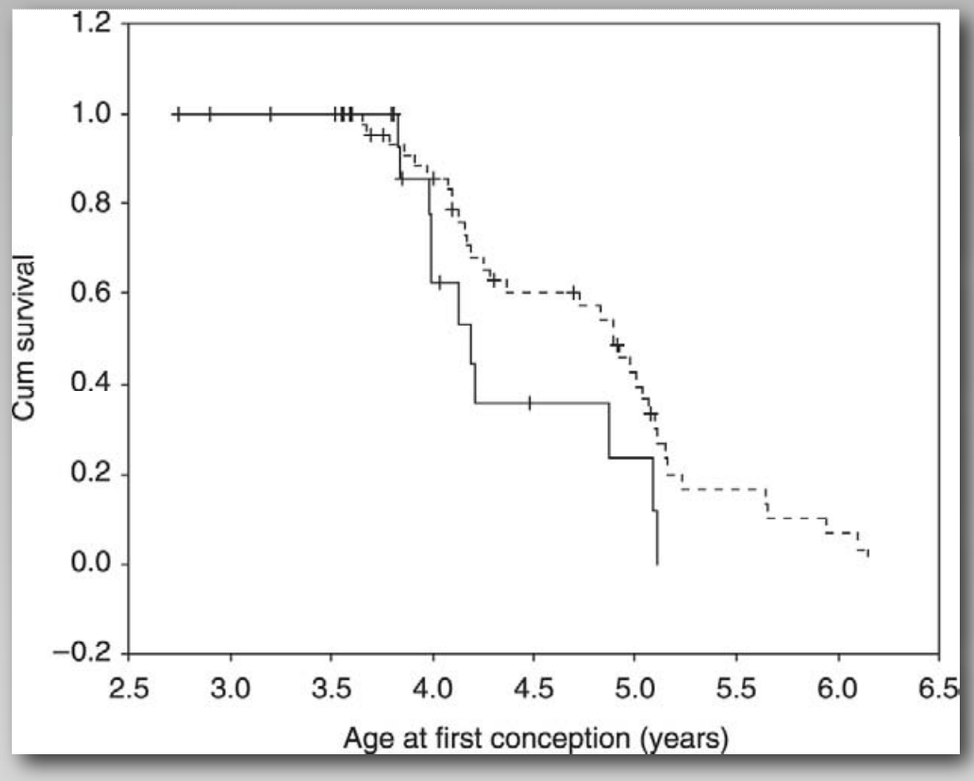


Fig. 2 Cumulative survival curve showing age at first reproduction in inbred (solid line) and noninbred (dashed line) female mandrills. Crosses indicate censored cases.

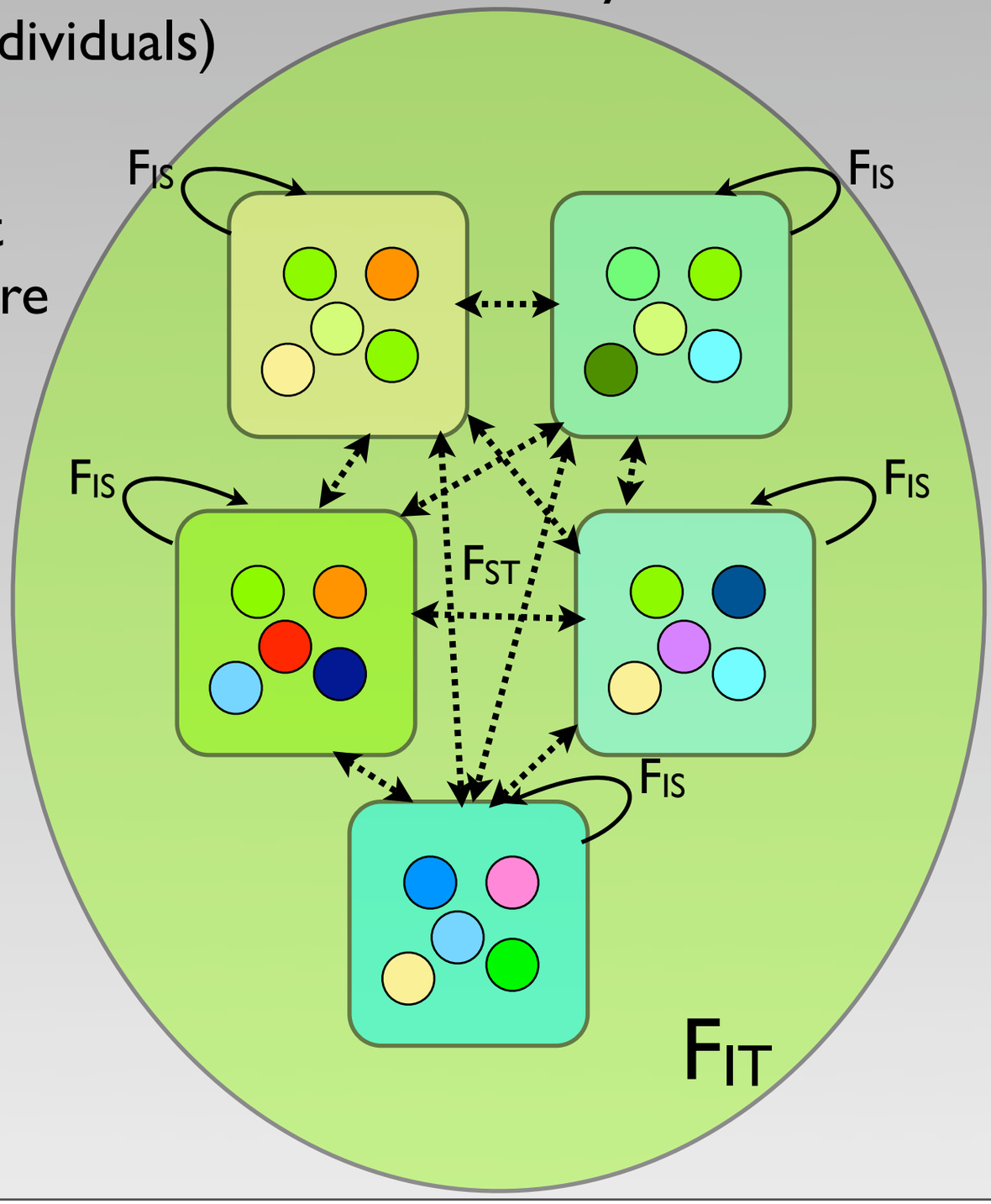
Population differentiation

- high fragmentation of habitats
  - instead of one continuous habitat (panmixia) ➡ separated populations without or with limited migration between them
- genetic differentiation between populations
  - due to genetic drift, stochasticity, selection, etc...
- measuring population fragmentation: *F*-statistics (Wright, 1969)

panmictic population is one where all individuals are potential partners

Population differentiation: *F-statistics*

- F_{IS}: probability that two alleles in an individual are identical by descent (≈ *F* averaged across all individuals) intra-population
- F_{ST}: fixation index - probability that two alleles from two populations are identical by descent between population structure between populations
- F_{IT}: general genetic structure
- $F_{IT} = F_{IS} + F_{ST} - (F_{IS})(F_{ST})$



Population differentiation: *F-statistics*

- $F_{IT} = F_{IS} + F_{ST} - (F_{IS})(F_{ST})$

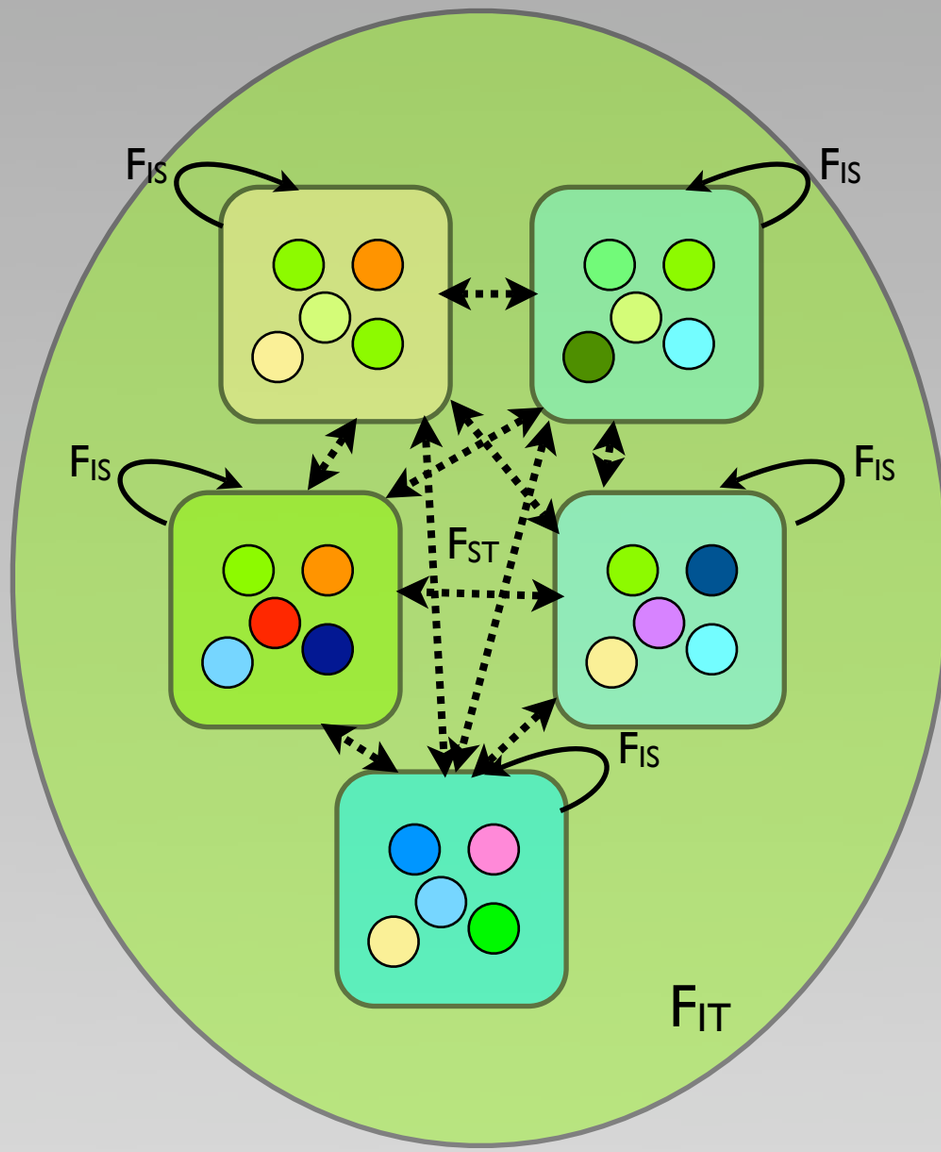
  or $F_{ST} = (F_{IT}-F_{IS})/(1-F_{IS})$

- but inbreeding and heterozygosity related:
  $F = 1-(H_O/H_E)$

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

  $F_{ST} = 1-(H_S/H_T)$

  $F_{IT} = 1-(H_I/H_T)$



H_I = observed heterozygosity averaged across all population fragments

H_S = expected heterozygosity averaged across all population fragments

H_T = expected heterozygosity for the total population (=H_e)

Population differentiation: *F-statistics*

- example 1:

	Genotypes						
Population	A_1A_1	A_1A_2	A_2A_2	Allele frequency	H_o	H_e =2pq	F =1-(H_O/H_E)
1	0.25	0.50	0.25	A_1: p=0.5 A_2: q=0.5	0.5	0.5	0
2	0.4	0.2	0.4	A_1: p=0.5 A_2: q=0.5	0.2	0.5	0.6

combined: H_I = 0.35 A_1: p=0.5, A_2: q=0.5 H_S = 0.5, H_T = 0.5

F_{ST} = 0, F_{IT} = 0.3, F_{IS} = 0.3

$H_T = 2*p*q$

$1-H_S/H_T$

$1-H_I/H_T$

$=1-H_I/H_S$

Population differentiation: *F-statistics*

- example 2:

	Genotypes						
Population	A_1A_1	A_1A_2	A_2A_2	Allele frequency	H_o	H_e =2pq	F =1-(H_O/H_E)
1	0.25	0.50	0.25	A_1: p=0.5 A_2: q=0.5	0.5	0.5	0
2	0.14	0.12	0.74	A_1: p=0.2 A_2: q=0.8	0.12	0.32	0.625

combined: H_I = 0.31 A_1: p=0.35 A_2: q=0.65 H_S = 0.41 H_T = 0.455

F_{ST} = 0.099 F_{IT} = 0.319 F_{IS} = 0.244

p = 2*A_1A_1 +A_1A_2

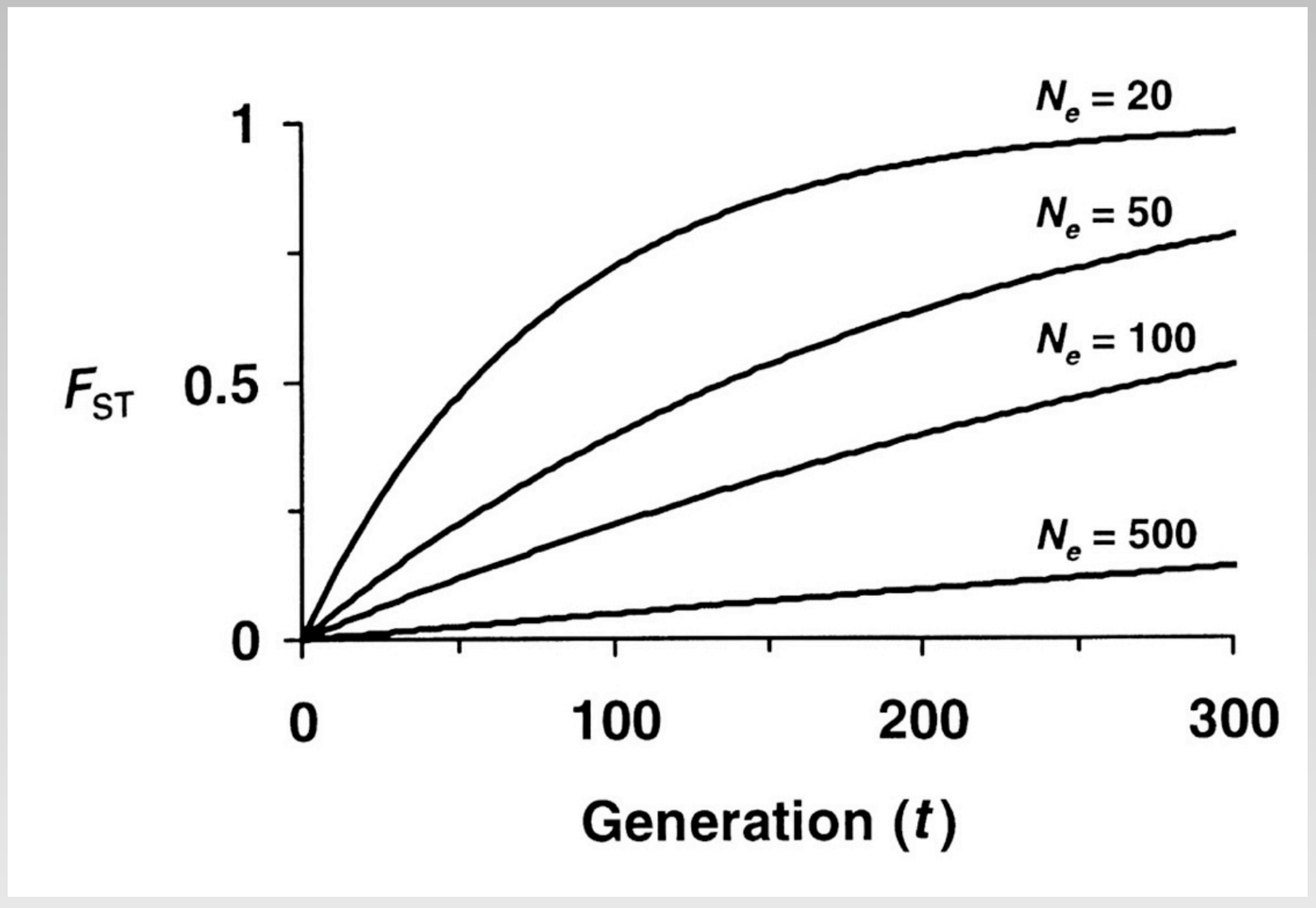
1-H_S/H_T

1-H_I/H_T

=1-H_I/H_S

Population differentiation: *evolution over time*

- when populations are isolated (no gene-flow):
  increase of the genetic differentiation between populations (F_{ST})



Population differentiation: *gene flow*

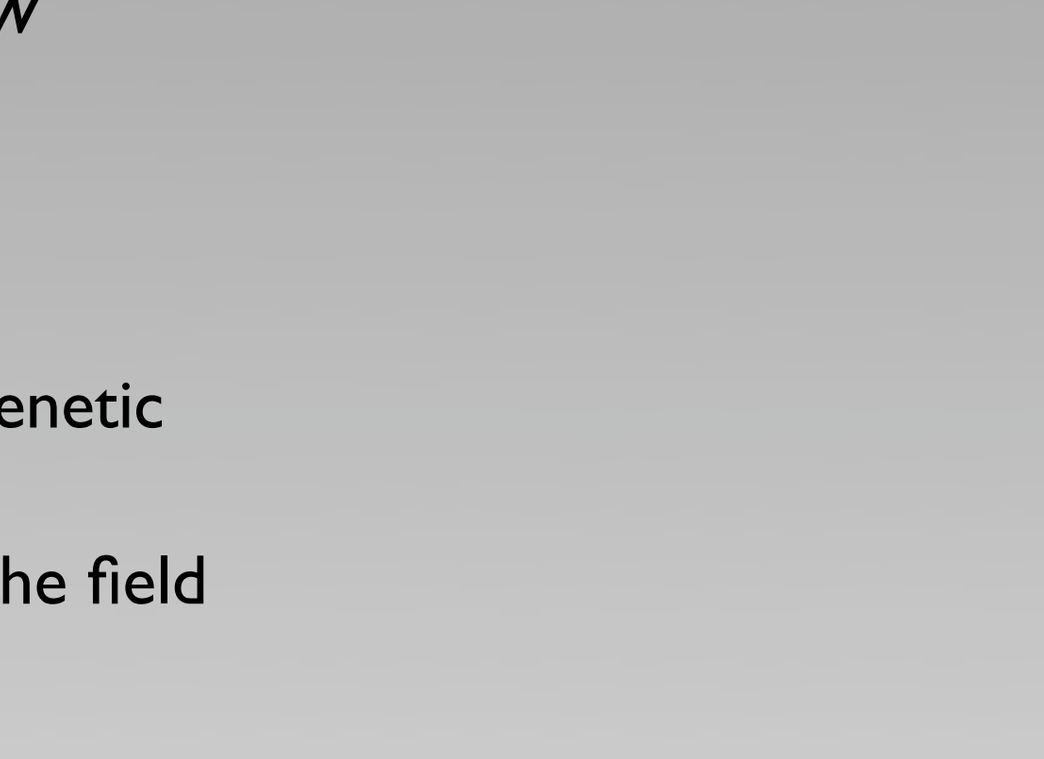
- gene flow reduce the isolation
- gene flow must be sufficient to avoid genetic differentiation
- measuring gene flow: very difficult on the field
  rough estimation using the function:

  $F_{ST} = 1/(4N_em+1)$

  N_e= effective population size

  m = migration rate

  N_em = number of migrant per generation



Relationship between inbreeding, heterozygosity, genetic diversity and population size

- numerous relationships between these parameters
- theory (for random mating populations)
  - relationship between inbreeding and heterozygosity
    $F = 1-(H_t/H_o)$
  - relationship between increase of inbreeding per generation and population size
    $\Delta F = 1/(2N)$
  - loss of genetic diversity ≈ inbreeding coefficient
- in practice (rarely completely random mating in all pop.)
  - large plant populations doing selfing: high inbreeding coefficient, low heterozygosity but high overall genetic diversity (alleles randomly distributed in the population but not within the individuals)
- relationship between inbreeding and loss of genetic diversity more complex in species with regularly high level of inbreeding

supplementary information

- books
  - Frankham, Ballou & Briscoe (2002) Introduction to Conservation Genetics, Cambridge University Press
  - Allendorf & Luikart (2007) Conservation and the Genetics of Populations, Blackwell Publishing
- articles
  - inbreeding: Keller & Waller (2002) Inbreeding effects in wild populations, **TRENDS in Ecology & Evolution** 17: 230-241
  - analyses softwares: Excoffier & Heckel (2006) Computer programs for population genetics data analysis: a survival guide, **Nature Reviews Genetics** 7:745-758
- technical and analyses
  - DNA manipulation (PCR, sequencing, etc.): http://www.dnai.org/b/index.html
  - softwares: e. g. http://www.biology.lsu.edu/general/software.html
    http://evolution.genetics.washington.edu/phylip/software.html