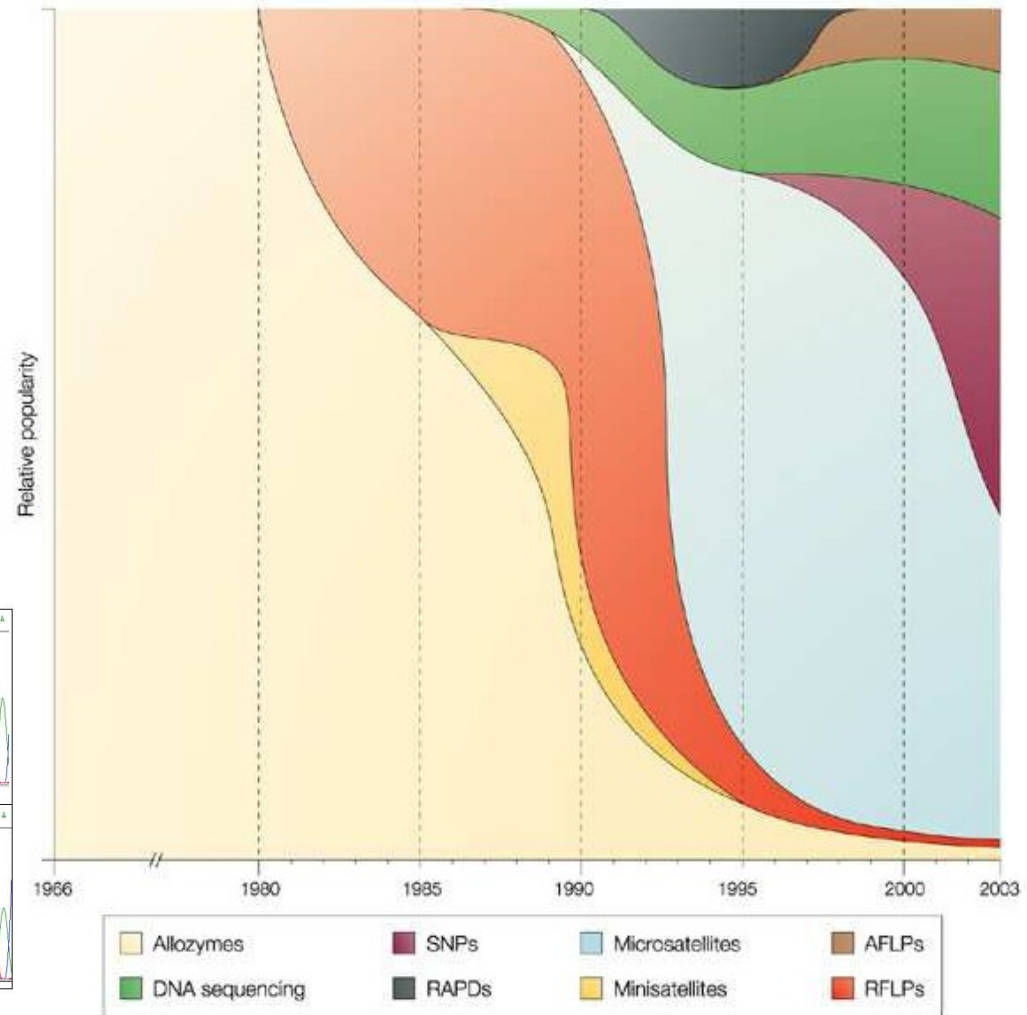
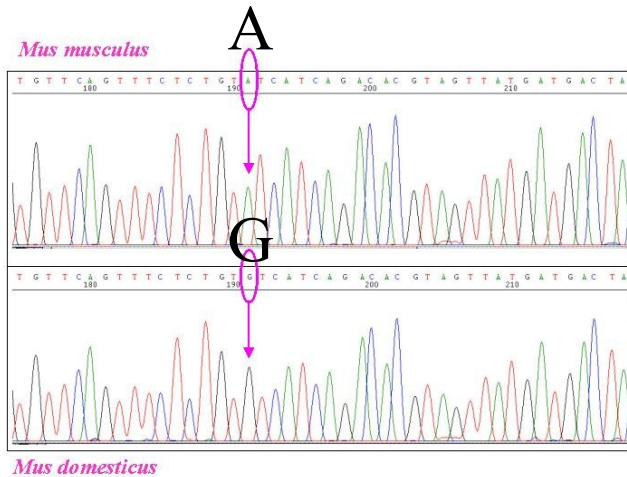
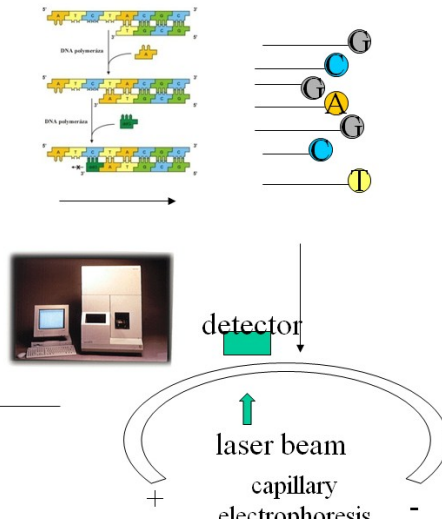
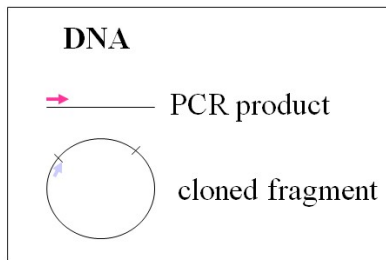


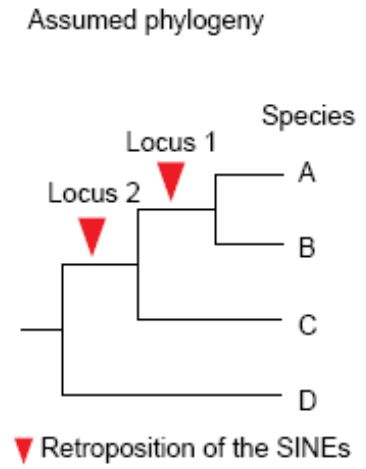
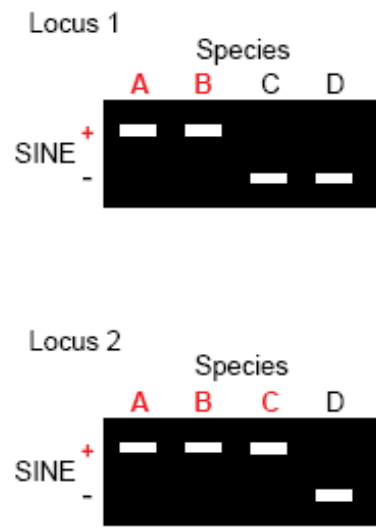
# SNP Fashion on the rise



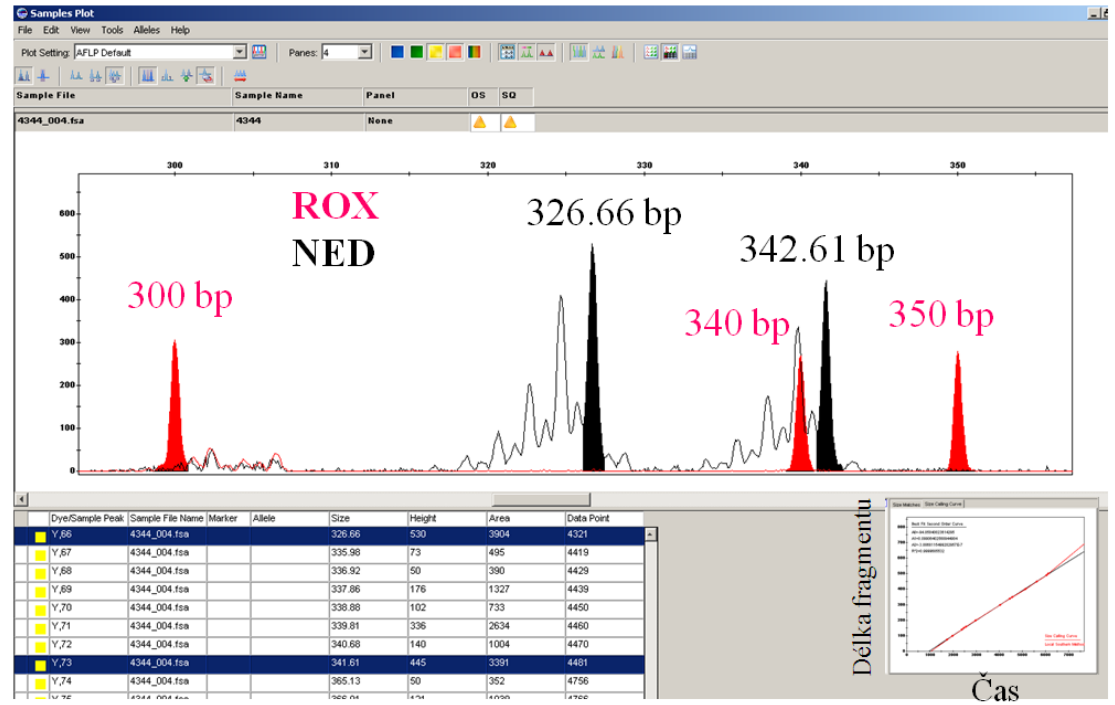
# Sekvencování DNA



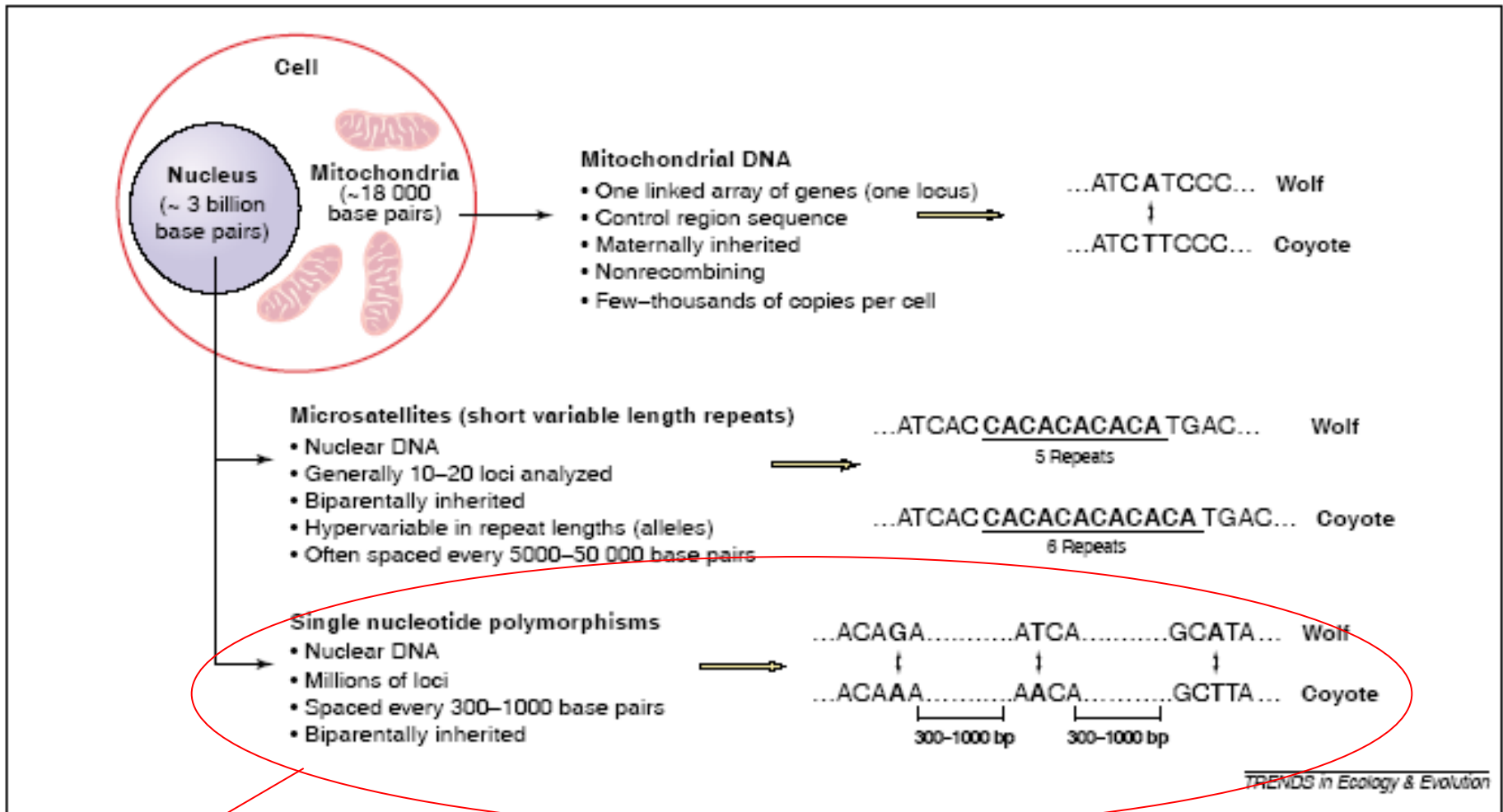
# SINE



# Mikrosateliti



# Single nucleotide polymorphisms (SNPs)



SNPs : nuclear genome (consensus)

# Single-locus genetic markers

- SNPs (single nucleotide polymorphisms) – sekvenční polymorfismus
- kodominantní – je možné odlišit heterozygota (např. A/T) od homozygota (např. A/A)

CA**A**GTA

TG**G**ACG

CA**T**GTA

TG**C**ACG

**A/T**

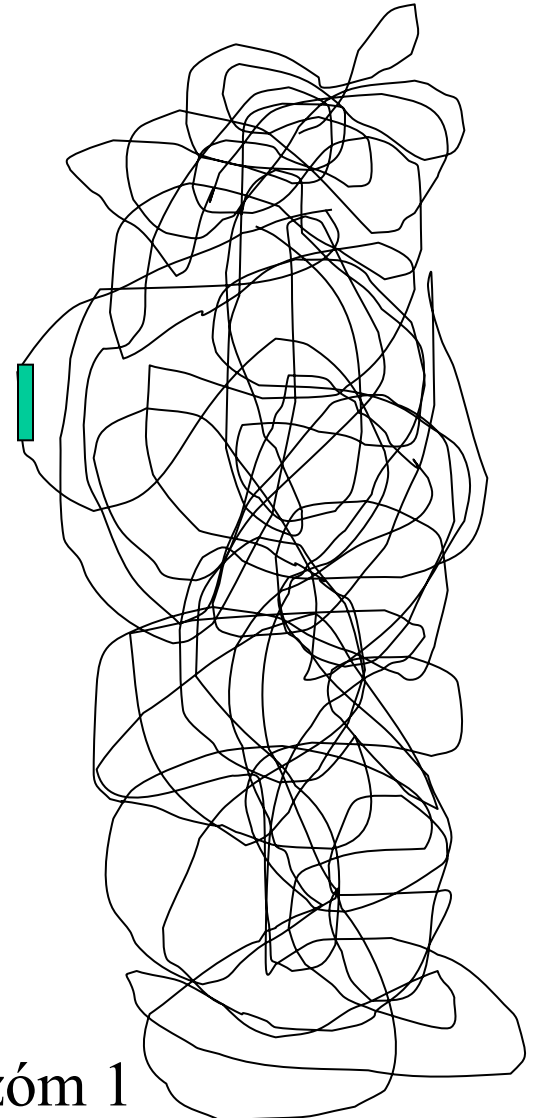
CA**A**GTA

TG**G**ACG

CA**A**GTA

TG**G**ACG

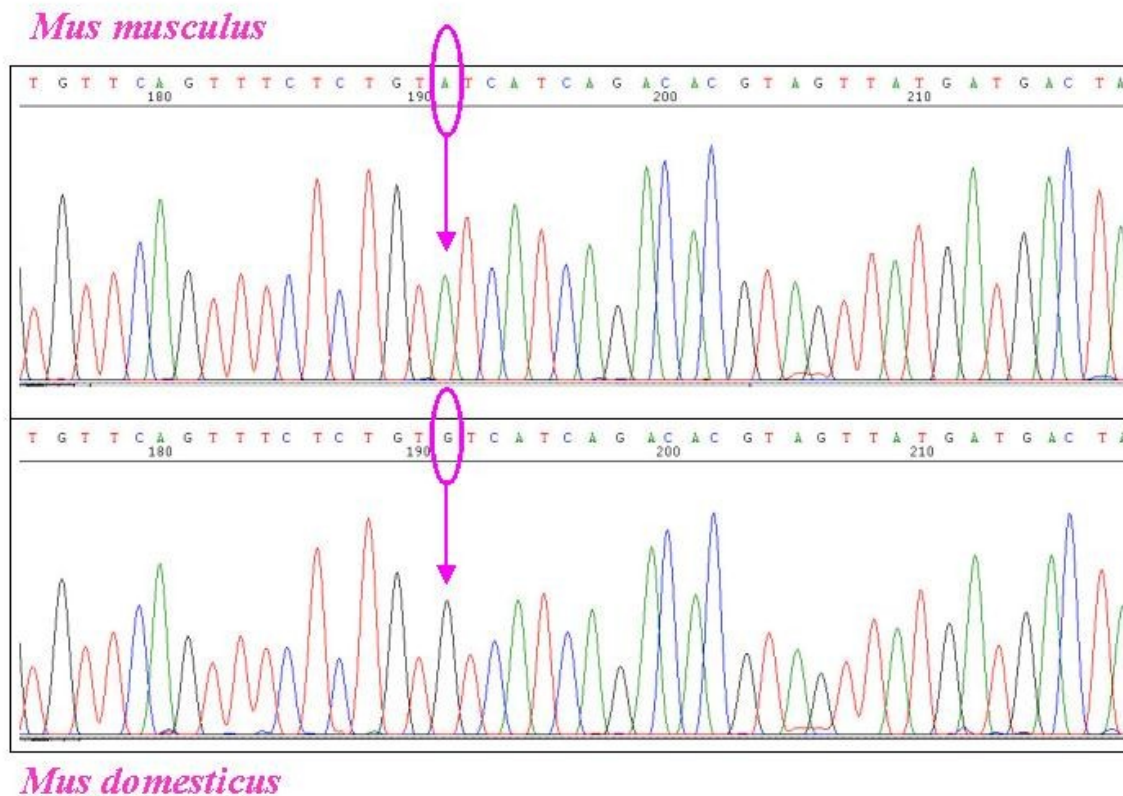
**A/A**



Př.: chromozóm 1

# Příklad informativního SNP znaku

- fixovaný polymorfismus (homozygoti) – využití např. při studiu hybridizací (hybridi = heterozygoti)



**transice**  
**A ↔ G**

transition: Pu → Pu or Py → Py

transversion: Pu → Py or Py → Pu

# Využití SNPs znaků

- obdobné jako u mikrosatelitů
- identifikace druhu (nebo genetické skupiny) - studium hybridizace
- fylogeografie
- populační genetika (genetická variabilita a struktura, tok genů, identifikace jedinců a vztahů mezi nimi, populační velikost a její změny atd.)

# Výhody

- početné a rozšířené v genomu (v kódujících i nekódujících oblastech) – milióny lokusů
- 1 SNP cca každých 300-1000 bp
- Mendelovská dědičnost (vs. mtDNA)
- evoluce je dobře popsitelná jednoduchým mutačním modelem (vs. microsatellites)
- jsou analyzovány kratší fragmenty DNA – neinvazivní genetika

# Nevýhody

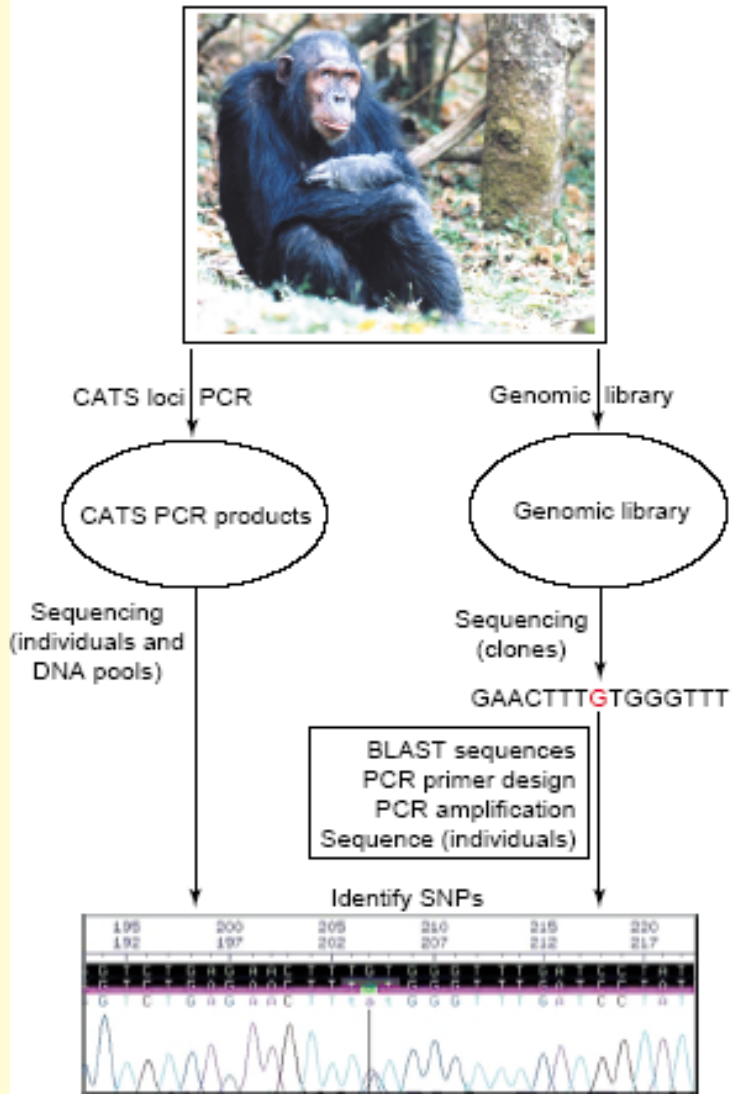
- „ascertainment bias“ – výběr znaků se provádí na základě jen malého počtu jedinců a nemusí být reprezentativní
- nízká variabilita na lokus (většinou jen 2 alely)
- pro populační genetiku je vyžadován větší počet lokusů (4-10 krát více než u mikrosatelitů)



# Metody analýzy

1. Nalezení lokusů („ascertainment“)
2. Genotypizace

# Nalezení SNPs

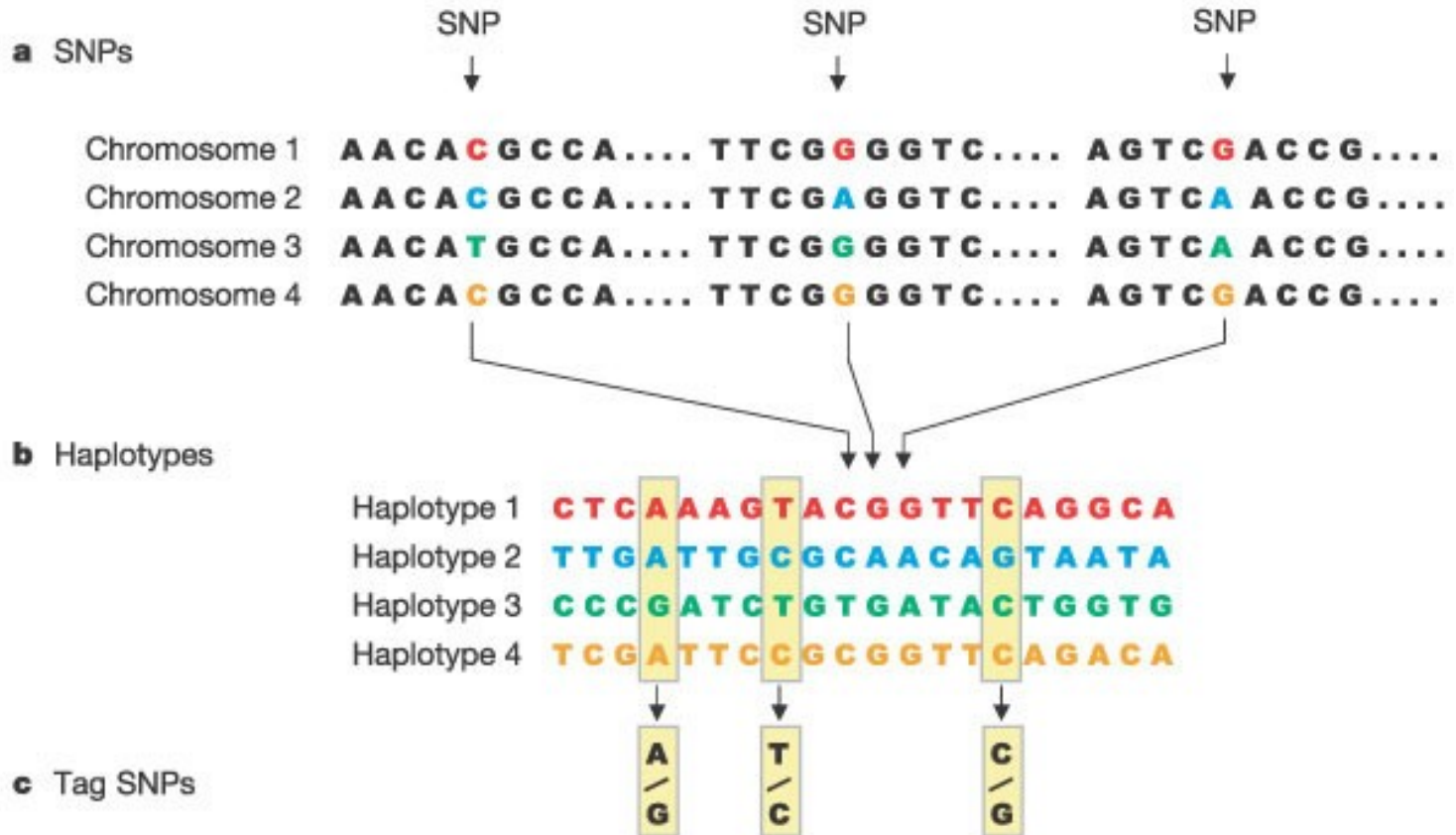


(1) CATS loci = comparative anchor tagged site loci (= cross amplification)

(2) Genomic library = genome restriction + cloning

**Next-generation sequencing**  
– sekvenování genomu více jedinců a hledání polymorfismů

# Identifikace různých genotypů u různých jedinců (= homologních chromozómů, tj. variabilita alel)

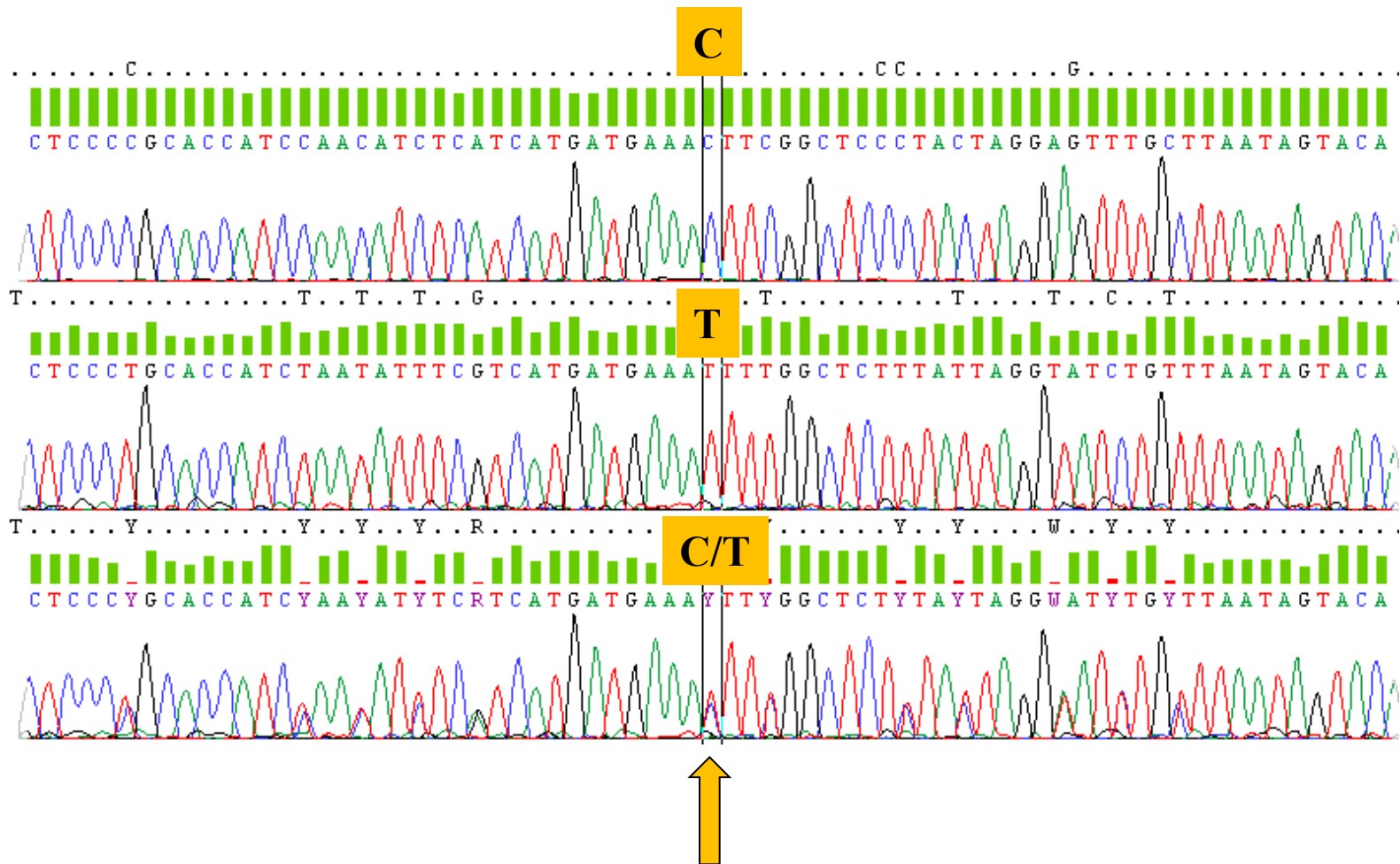


SNPs genotyping

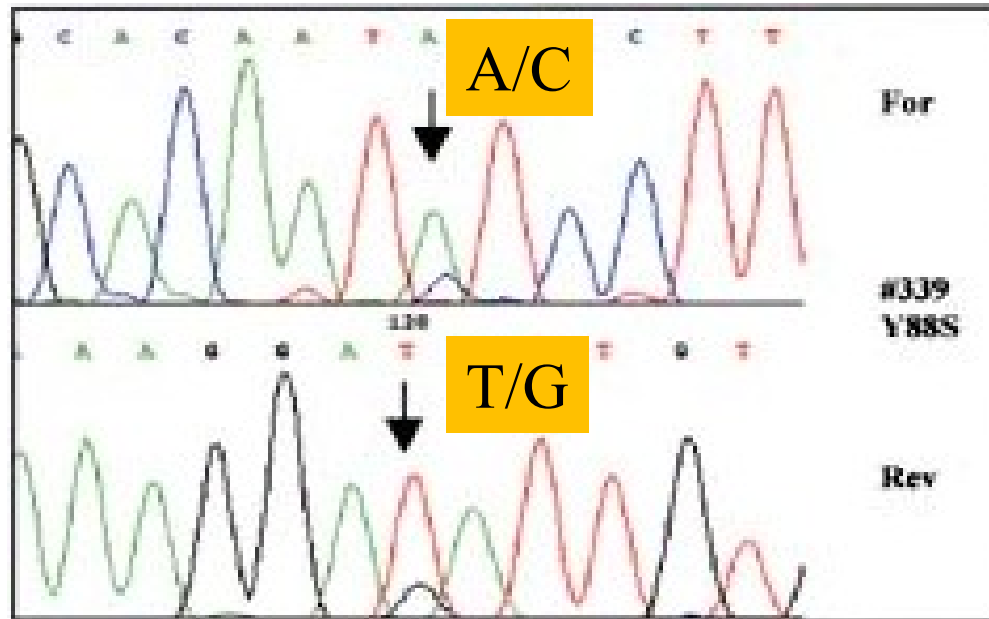
= zjištění genotypu daného  
jedince

# SNPs genotyping - sekvenování?

## Je drahé a nejasné u heterozygotů



# Heterozygotes?



Bi-directional sequencing - are you really sure?

SNPs genotyping - klonování a následné sekvenování?  
- separation of two (or more in duplicated genes) alleles

each clone contain the only allele

!!! cloning - 1000 Kč  
!!! sequencing 1 clone - 150 Kč

↑ ligation, transformation

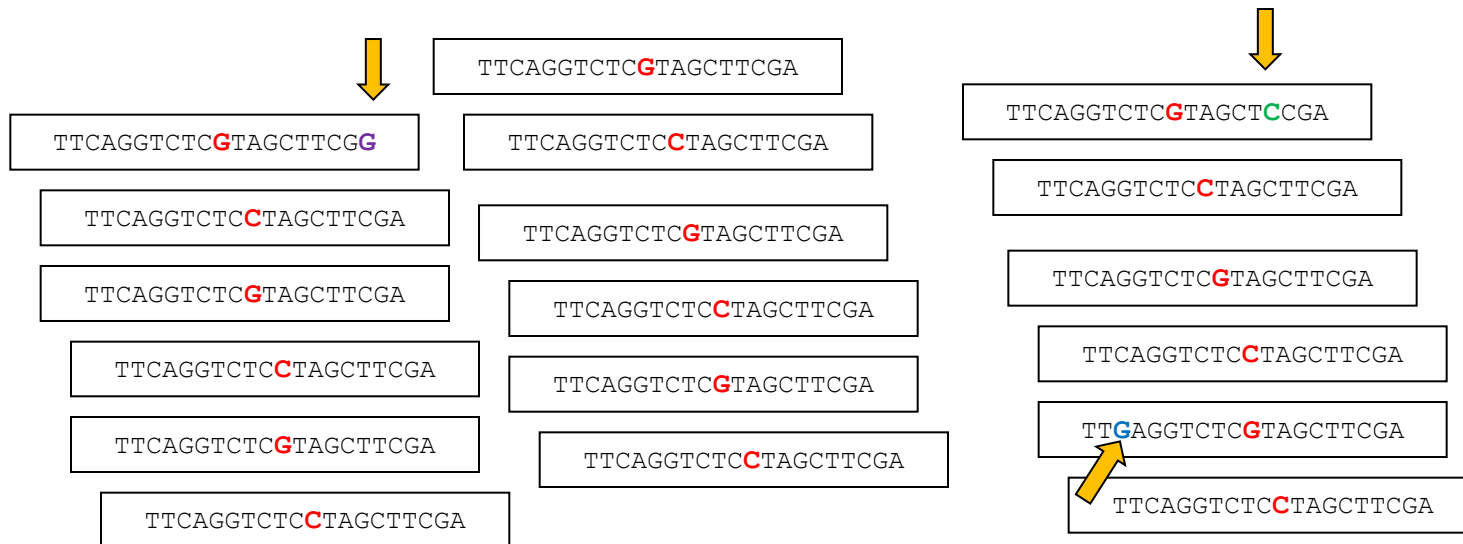
 Ex.: heterozygote = two diff. alleles

# PCR is making substitution errors that are visualised by cloning (!)

TTCAGGTCTC**G**TAGCTTCGA

TTCAGGTCTC**C**TAGCTTCGA

... před PCR = heterozygot G/C



PCR artefacts



# SNPs genotyping

## 1. Old standards (PCR-based)

- RFLP: PCR + štěpení + standardní elfo
- DGGE, TGGE, SSCP: PCR + nestandardní elfo
- původně detekce geneticky podmíněných chorob, např. cystická fibróza

## 2. New methods (not based on standard PCR)

- HRM: high-resolution melting (real-time PCR)
- real-time PCR se specifickými sondami (TaqMan, molecular beacon)
- ASPE: allele-specific primer extension
- SBE: single base extension
- SNP microarrays (GeneChip method)

# SNP genotyping - old standards

## PCR-RFLP

(restriction fragments length polymorphism)

## Enzyme Site Recognition

- Each enzyme digests (cuts) DNA at a specific sequence = restriction site
- Enzymes recognize 4- or 6- base pair, palindromic sequences (eg GAATTC)

**Restriction site**

**Palindrome**

G T A G A A T T C A T T C A C G C A  
C A T C T T A A G T A A G T G C G T

G T A G                      A A T T C A T T C A C G C A  
C A T C T T A A                      G T A A G T G C G T

**Fragment 1**

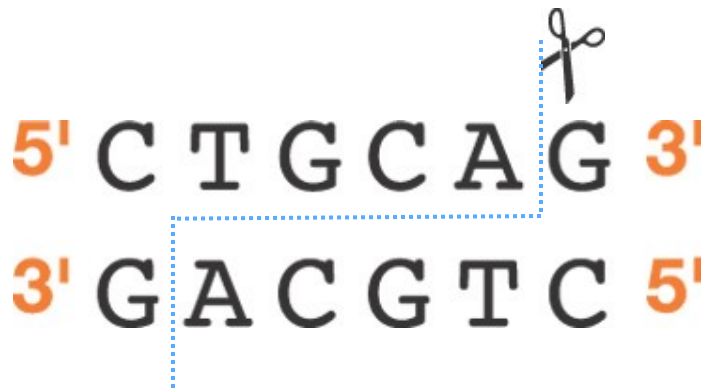
**Fragment 2**

# Common Restriction Enzymes



## ***EcoRI***

- *Escherichia coli*
- 5 prime overhang



## ***PstI***

- *Providencia stuartii*
- 3 prime overhang

# SNP genotyping - old standards

## PCR-RFLP

### Allele A

CCGATCA**A**TGCGGCAA  
GGCTAGT**T**ACGCCGTT

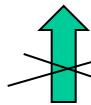


cutting by restriction endonuclease

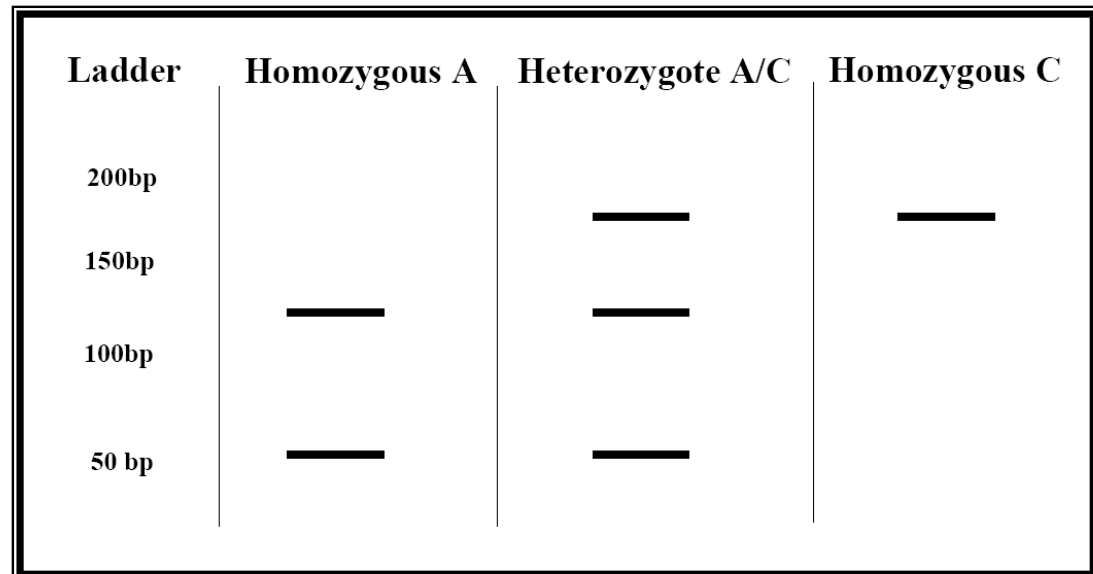
- neumožní nalézt novou variantu daného SNP (odliší pouze 2 formy daného znaku: +/- )

### Allele C

CCGATCA**C**TGCGGCAA  
GGCTAGT**G**ACGCCGTT



no cut



# SNPs genotyping - old standards electrophoresis methods of mutation detection

- Thermal gradient gel electrophoresis (TGGE)
- Denaturing gradient gel electrophoresis (DGGE)
- Single-strand conformation polymorphism (SSCP)

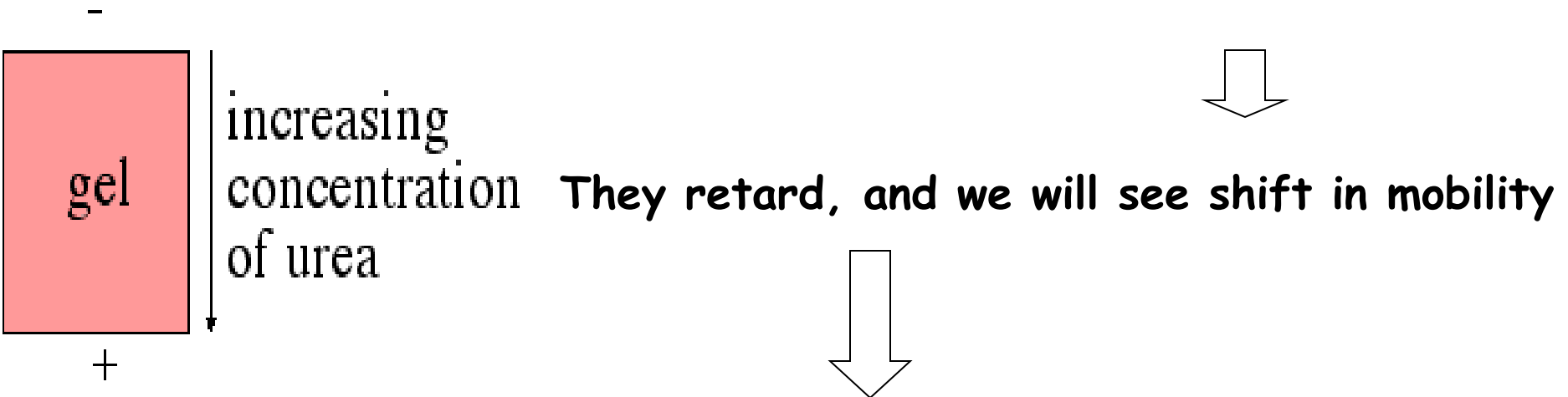
= special electrophoresis methods based on differences in mobility of different DNA sequences

# Denaturing gradient gel electrophoresis (DGGE) (TGGE - podobné, ale gradient teploty)

The small (200-700 bp) genomic fragments are run on a low to high denaturant GRADIENT acrylamide gel

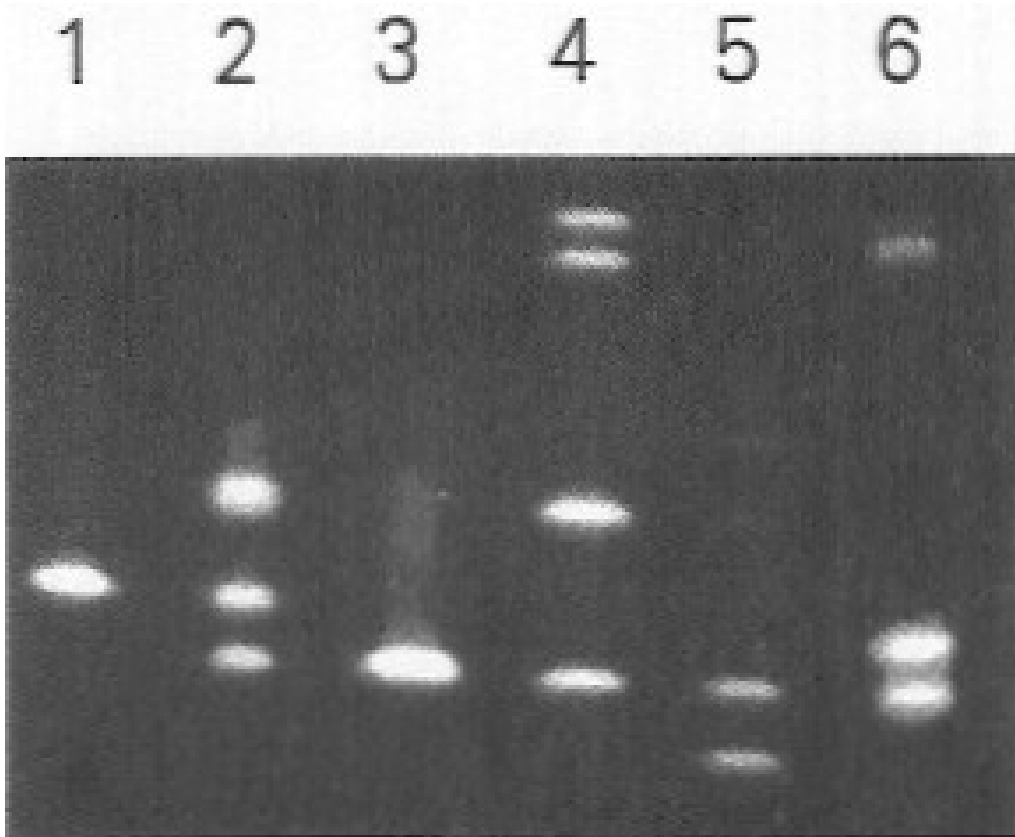
Each fragments move according to molecular weight, but as they progress into more denaturing conditions, each (depending on its sequence composition) reaches

**A POINT where the DNA BEGINS TO MELT**



We will see different shifts in mobility for differing products

Detekce nových mutací – např. v diagnostice genetických chorob  
nebo při analýzách MHC



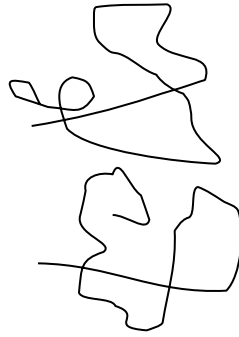
1 - normal homozygote  
3 - homozygous mutations  
will yield one band  
on a different position  
2, 4, 5, 6 - heterozygous  
mutations will yield 4  
bands (2 homozygous and 2  
heterozygous)

**NOT ALL BANDS ARE  
SEEN !!!!!**

# Single strand conformation polymorphism (SSCP)

## Allele 1 - C

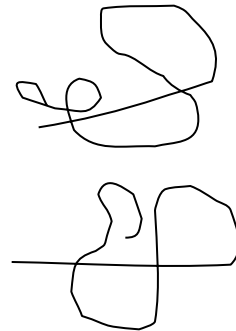
...CGCTT**C**AGG ...  
...GCGAA**G**TCC...



heating - denaturation  
snap-cooling → partial renaturation

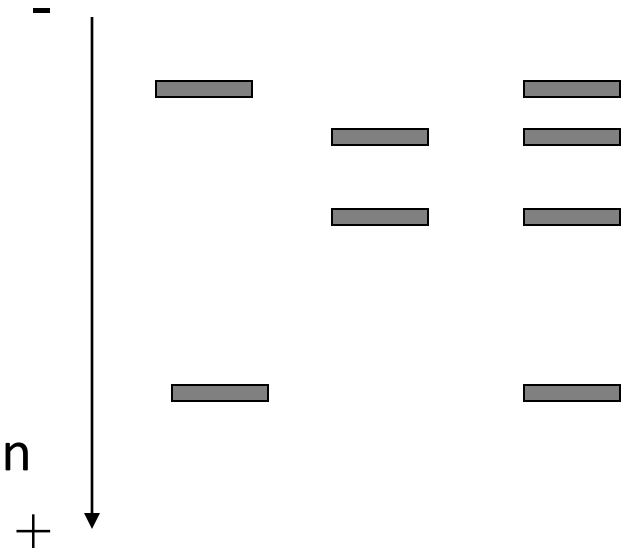
## Allele 2 - A

...CGCTT**A**AGG ...  
...GCGAA**T**TCC...



sequence-specific  
ssDNA conformations

Homo1 Homo2 Hetero



!!! non-denaturing PAGE

radioisotopes  
silver-staining  
fluorescent dyes (SYBR gold)





# Použití automatických sekvenátorů

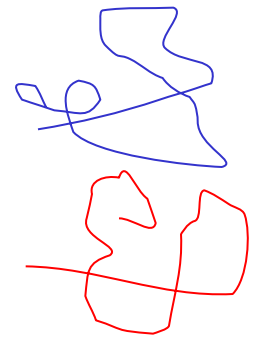
Why not non-denaturing electrophoresis?  
e.g. CAP (conformation analysis polymer)



- well controlled electrophoresis
- two fluorescent labels
- high sensitivity

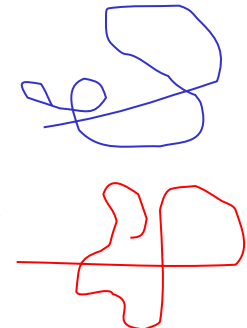
## Allele 1

*FAM*... CGCTTCAGG ...  
... GCGAAGTCC ...*HEX*



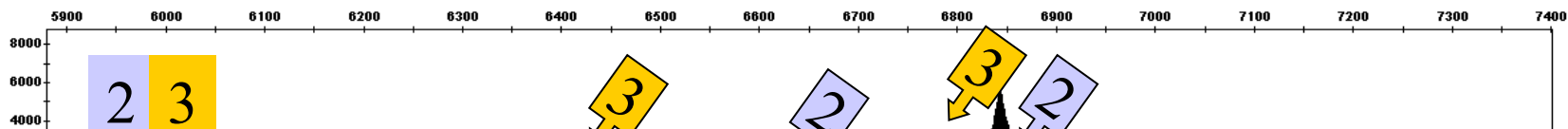
## Allele 2

*FAM*... CGCTTAAGG ...  
... GCGAATTCC ...*HEX*



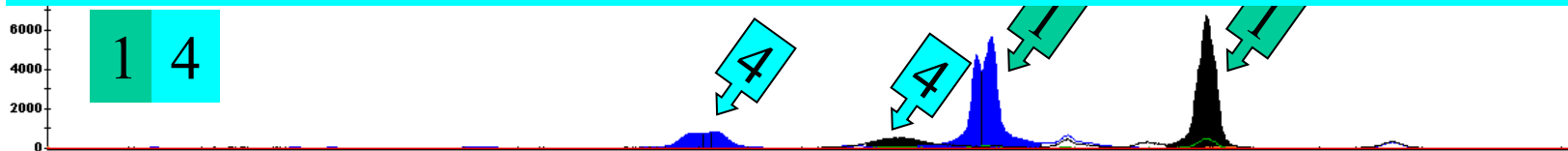
## MHC Class II (DQA gene) – mice HZ

Sample File	Sample Name	Panel	DS	SQ
hz319_004.fsa	hz319	None		



1 hour, ~ 100 Kč/4 samples incl. PCR

Information about all alleles (vs. cloning-sequencing)



	Dye/Sample Peak	Sample File Name	Marker	Allele	Size	Height	Area	Data Point
	B.65	hz701_003.fsa			6537.54	788	6803	6530
	B.66	hz701_003.fsa			6542.55	830	17081	6535

# Data analysis

- GeneMapper (Applied Biosystems)
- different „Size Standard“ for each temperature
- alignment of more samples
- allows detection of short sequences with several SNPs (very useful for e.g. MHC genotyping)

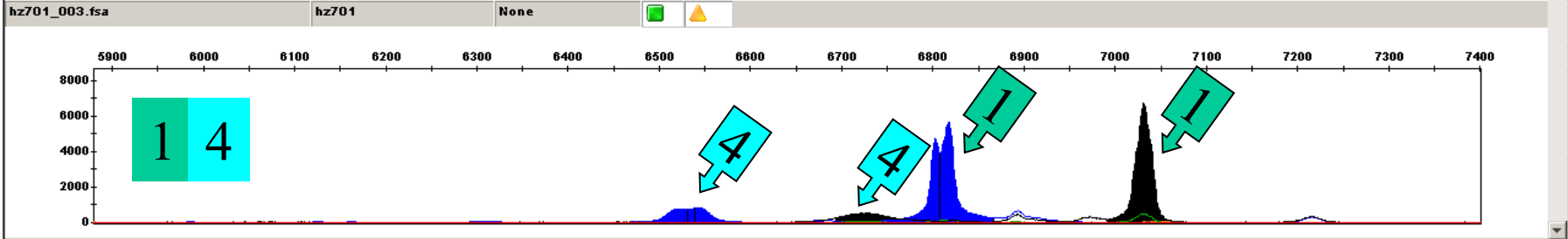
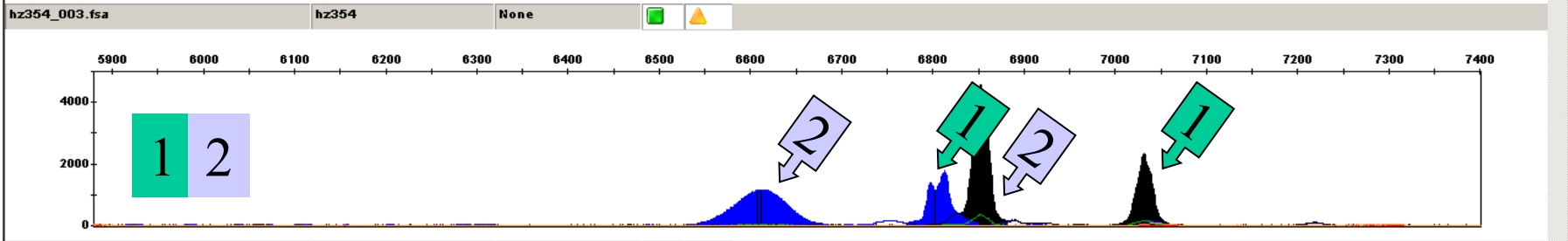
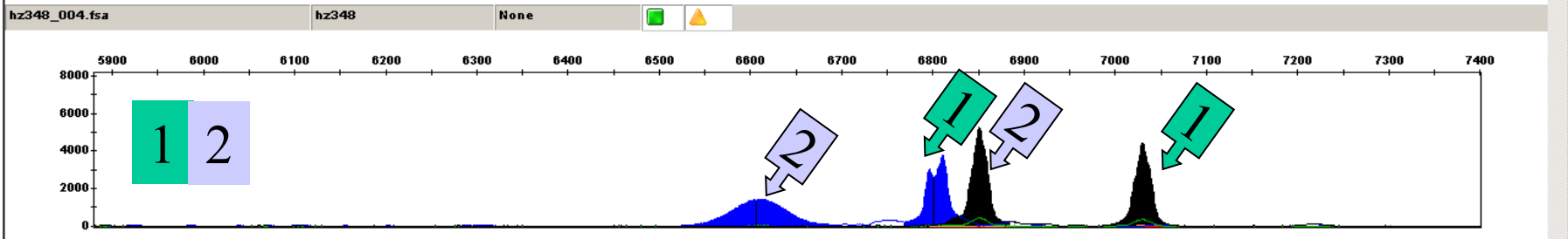
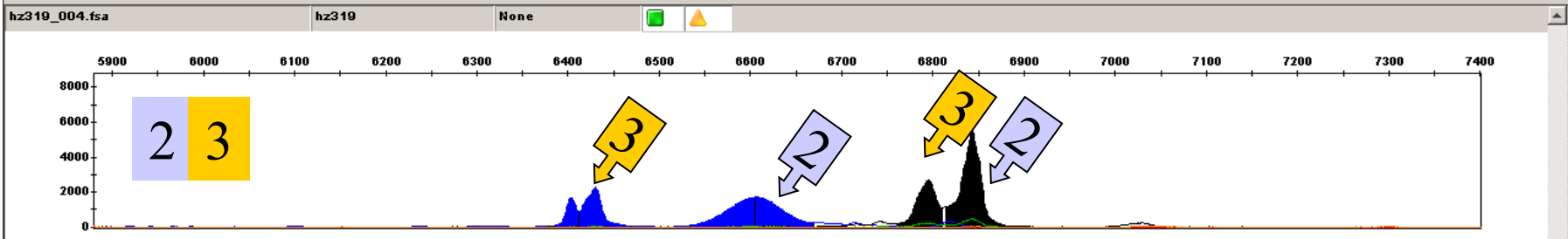
# Applications

- 1) Genotyping of codominant markers  
(e.g. single copy MHC genes)

# MHC Class II (DQA gene) – house mice

Plot Setting: AFLP Default Panes: 4

Sample File	Sample Name	Panel	OS	SQ
hz319_004.fsa	hz319	None	Green	Yellow
hz348_004.fsa	hz348	None	Green	Yellow
hz354_003.fsa	hz354	None	Green	Yellow
hz701_003.fsa	hz701	None	Green	Yellow



Dye/Sample Peak	Sample File Name	Marker	Allele	Size	Height	Area	Data Point
B,65	hz701_003.fsa			6537.54	788		
B,66	hz701_003.fsa			6542.55	830		

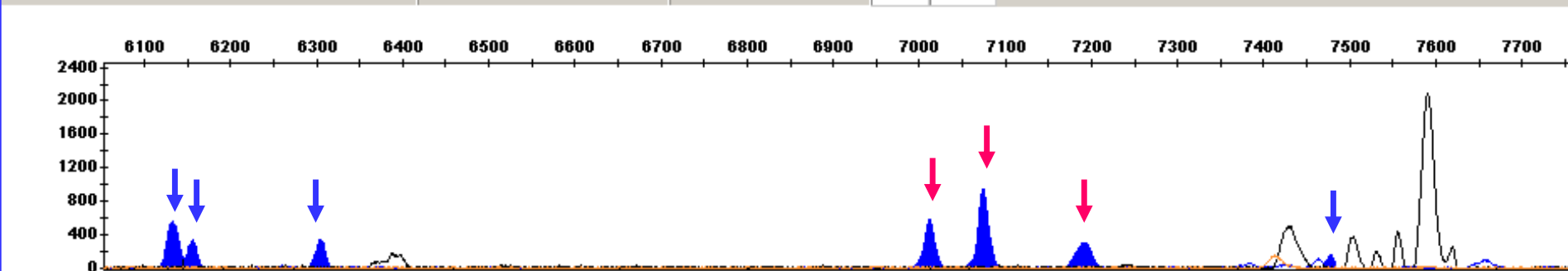
... even shape of the peaks is important !!!

# Applications

- 1) Genotyping of codominant markers  
(e.g. single copy MHC genes)
- 2) Identification of number of genes  
(e.g. duplicated MHC genes)



Seven peaks in one colours =  
= At least four amplified copies !!!



SSCP of three individuals:

↓ - different alleles

↓ same alleles



# Applications

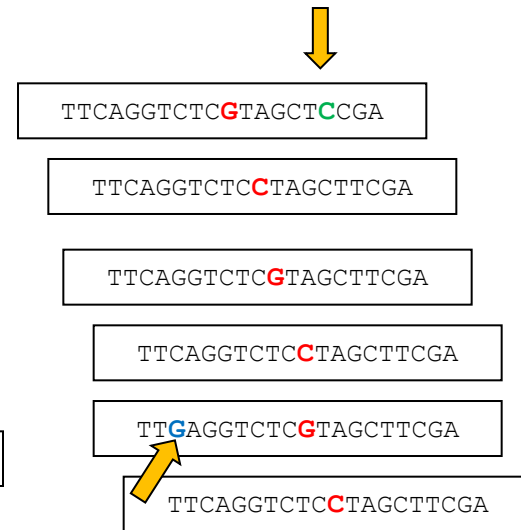
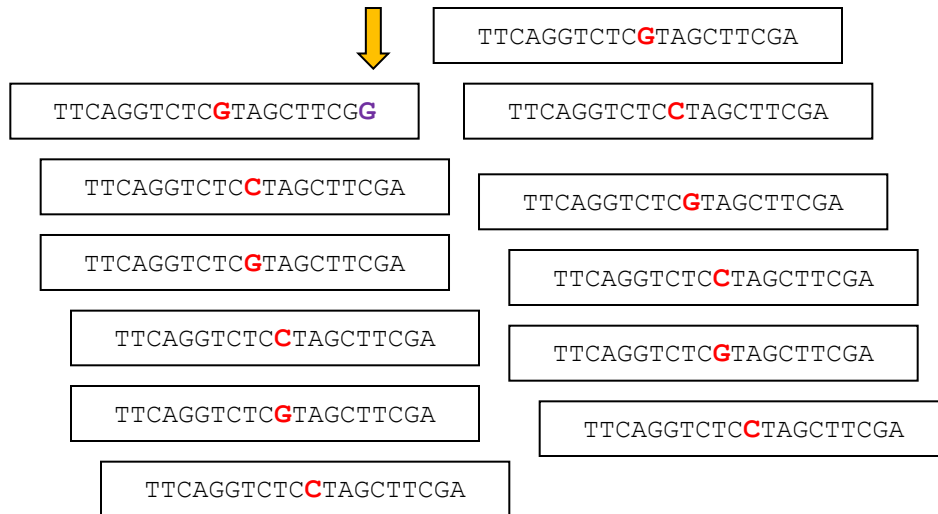
- 1) Genotyping of codominant markers  
(e.g. single copy MHC genes)
- 2) Identification of number of genes  
(e.g. duplicated MHC genes)
- 3) Detection of PCR artefacts during  
cloning

# Detection of PCR artefacts during cloning

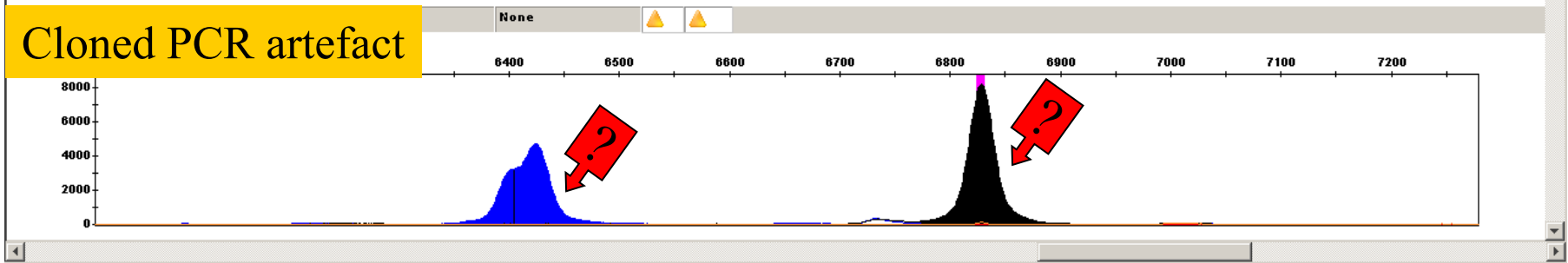
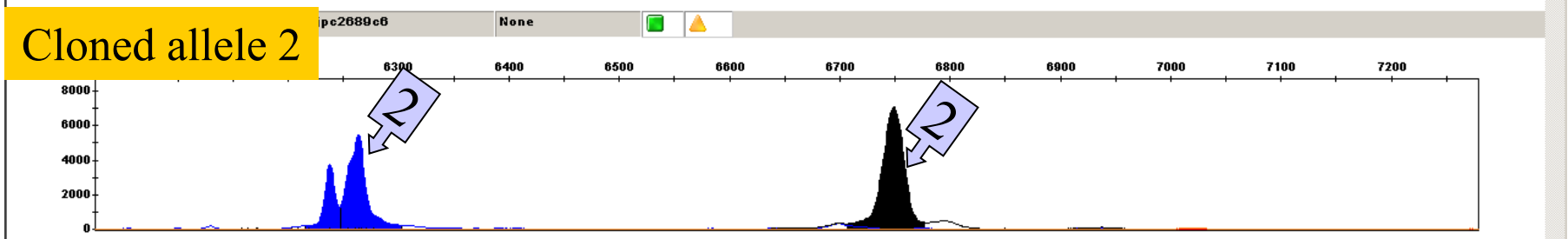
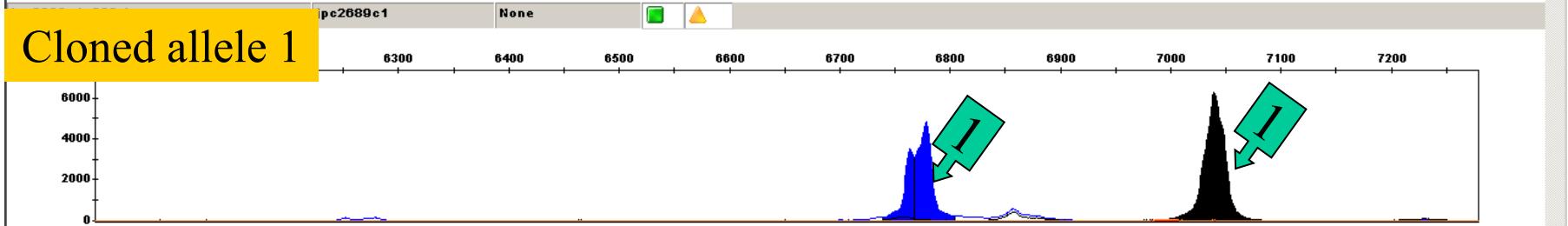
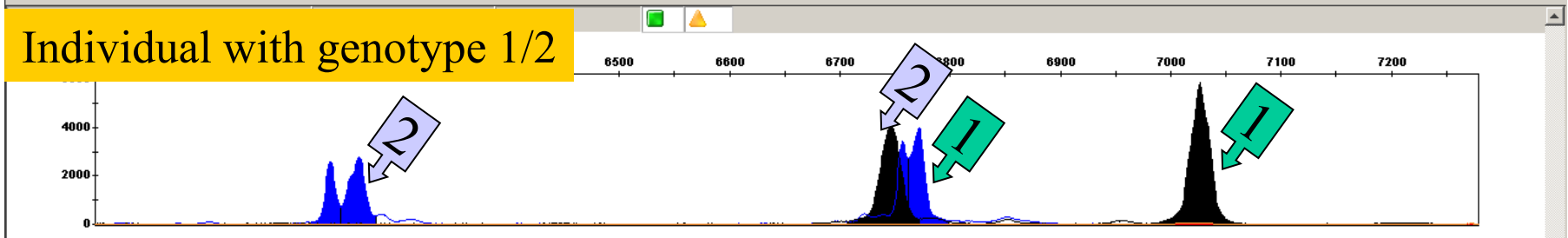
TTCAGGTCTC**G**TAGCTTCGA

TTCAGGTCTC**C**TAGCTTCGA

... před PCR = heterozygot G/C



# MHC Class II (DQA gene) – house mice



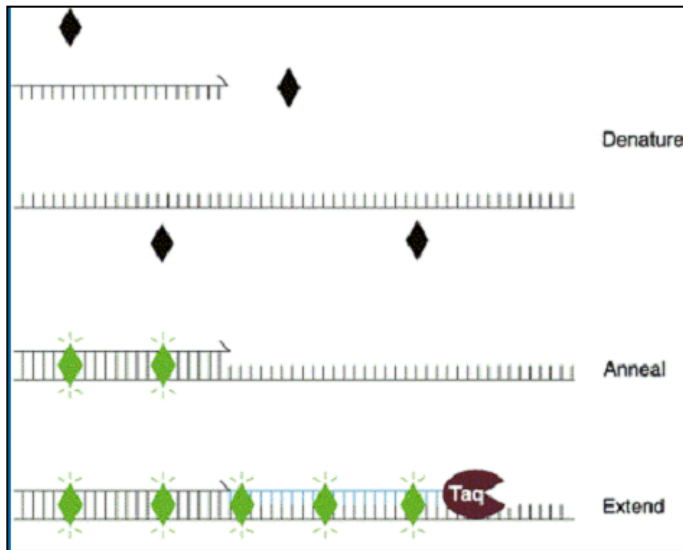
## Detection of PCR artefacts during cloning of heterozygotes

# SNP genotyping - new methods

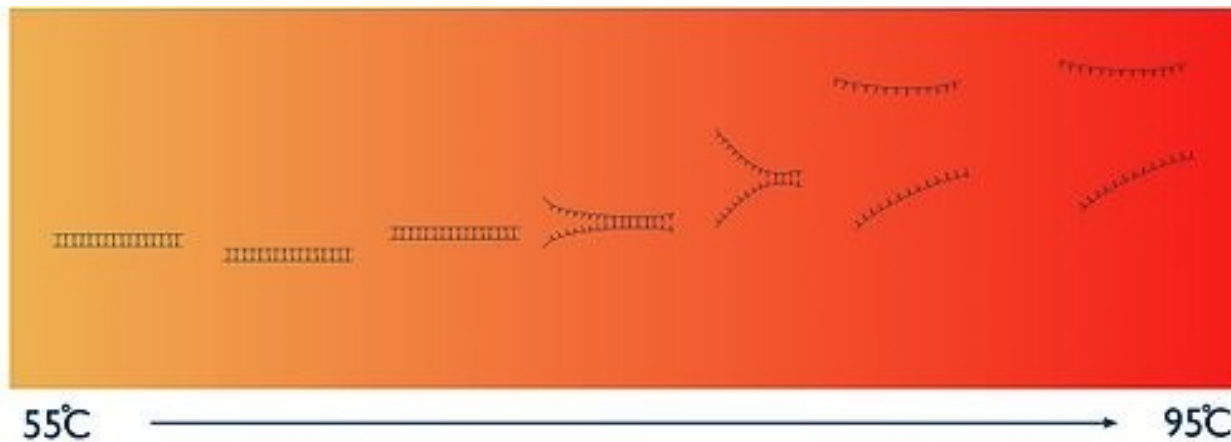
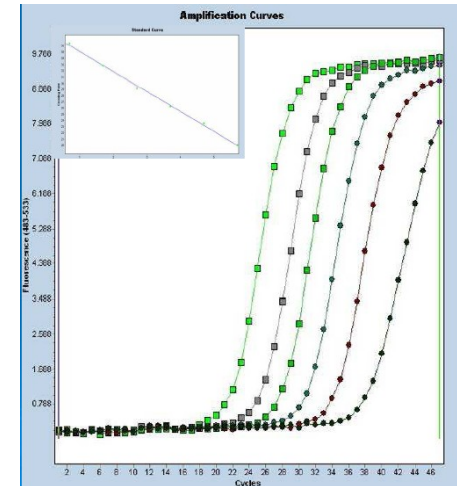
= not based on standard PCR

1. high-resolution melting temperature (HRMT)
2. real-time PCR se specifickými sondami (TaqMan, molecular beacon)
3. ASPE: allele-specific primer extension
4. SBE: single base extension
5. SNP microarrays (GeneChip method)

# High-resolution melting temperature (HRMT)

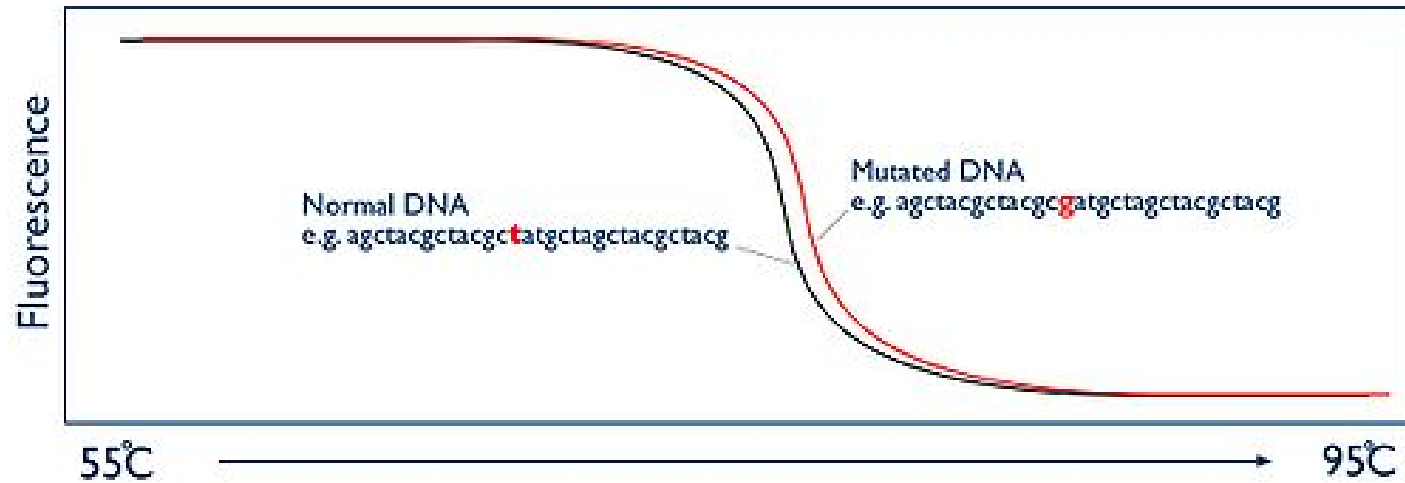


Step 1: real-time PCR = increase of fluorescence

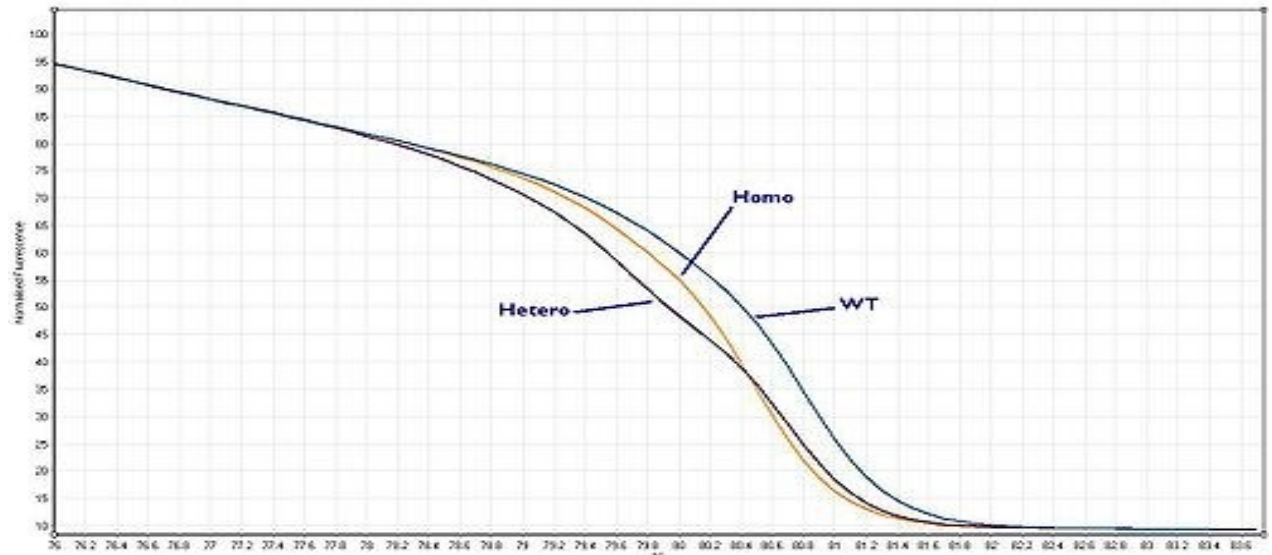


Step 2: measuring melting after PCR = decrease of fluorescence

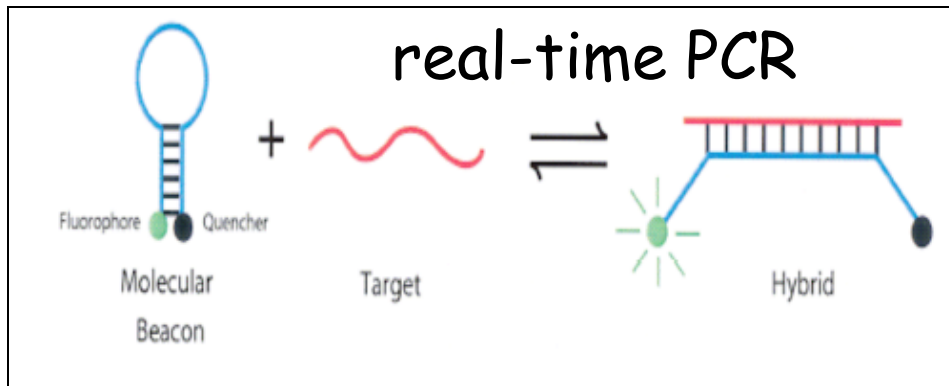
# HRMT genotyping



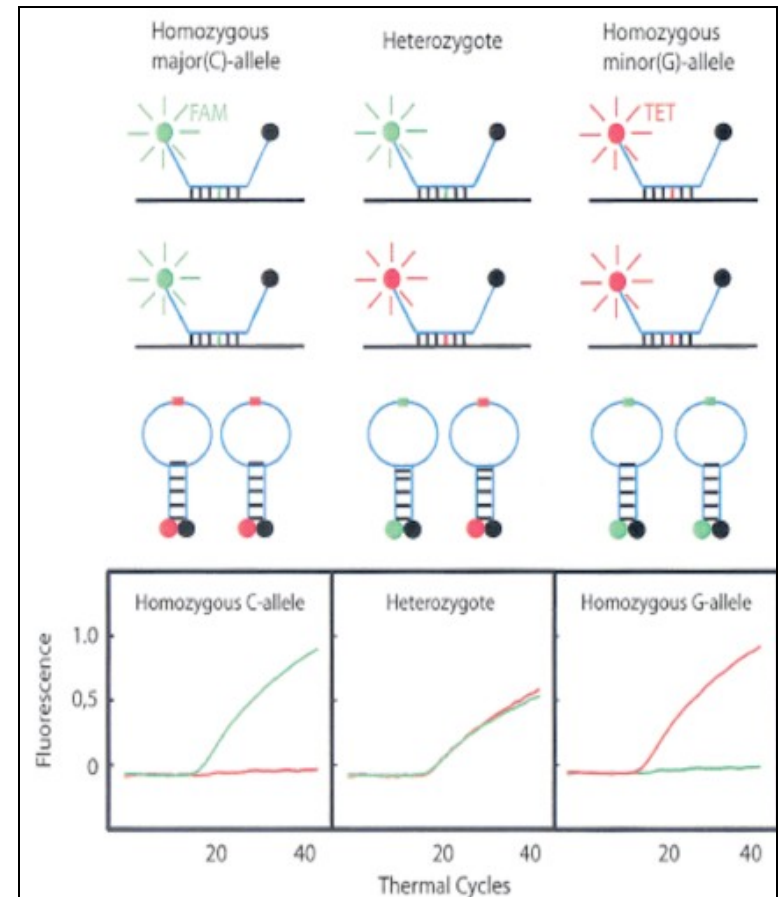
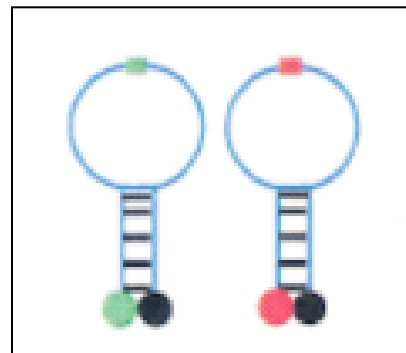
Detekce  
heterozygotů



# Real-time PCR se specifickou sondou

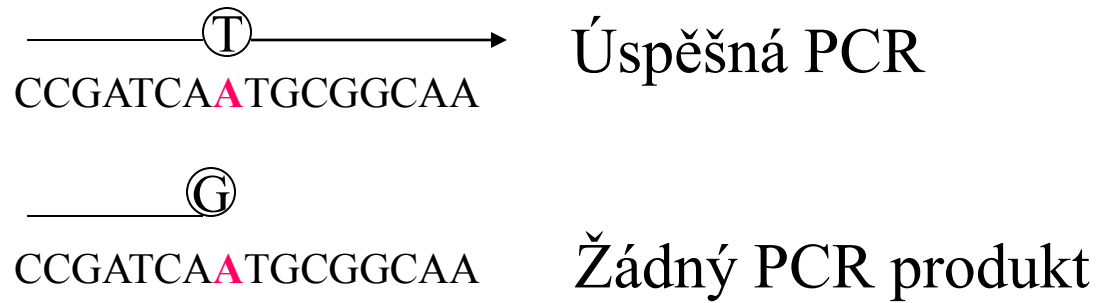


sondy  
specifické pro  
jednotlivé alely



- 1) TaqMan sondy
- 2) Molecular Beacons („maják“)

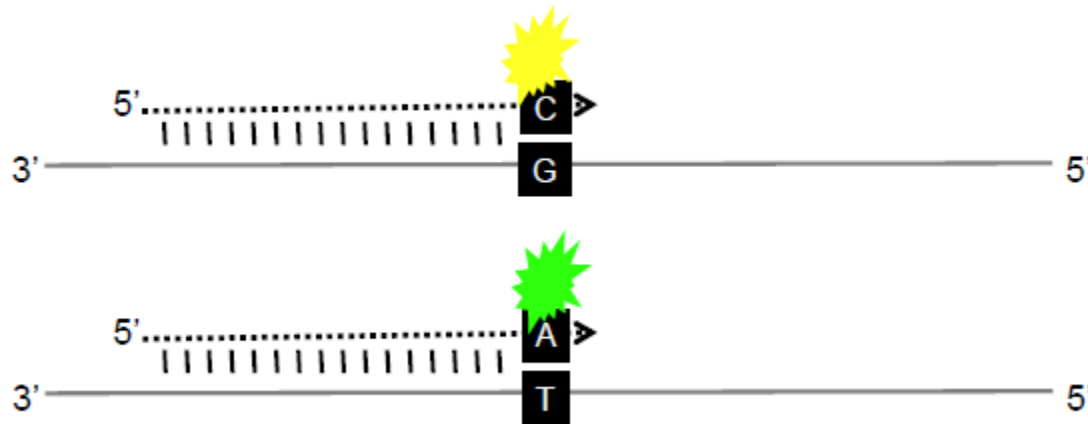
# ASPE: allele-specific primer extension



- dvě PCR se specifickými primery
- 3' terminální nukleotid na primerech je komplementární k SNP nukleotidu
- alelově-specifická amplifikace je umožněna vysoce specifickou polymerázou

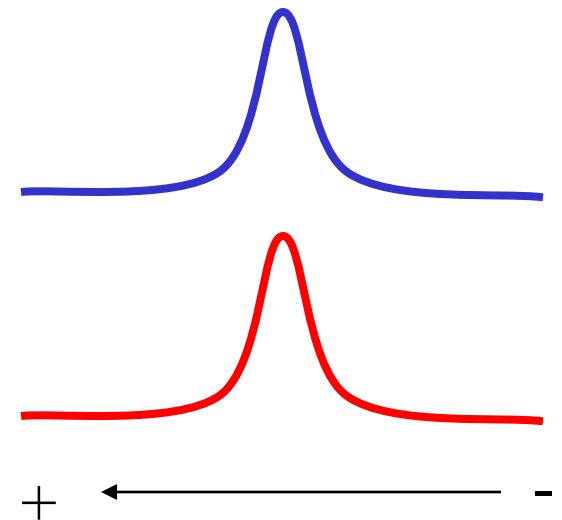
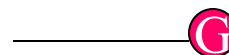
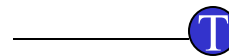


# ASPE: allele-specific primer extension (automatizovaná verze)



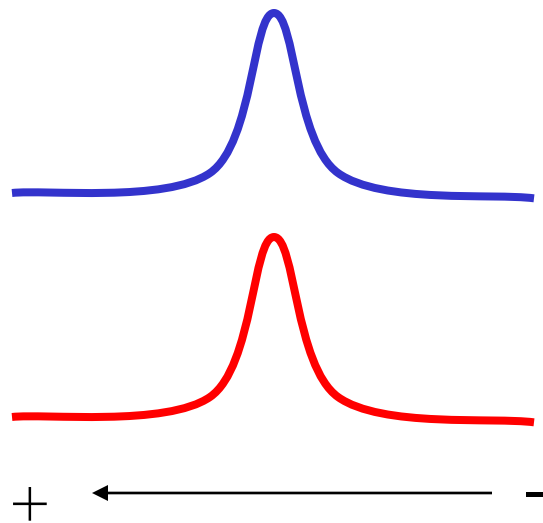
- existují zoptimalizované multiplexy pro modelové druhy (např. člověk 1536 SNPs)
- fluorescenční detekce (Illumina)

# (3) SBE: single base extension

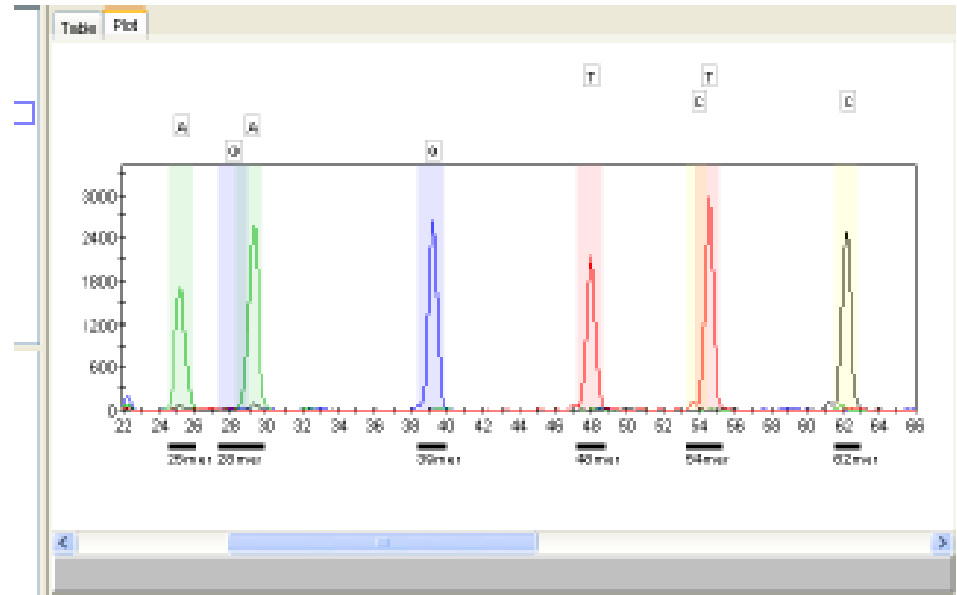


- pouze jeden dideoxynukleotid je přidán k primeru
- detekce různými metodami

# Detection of SBE products



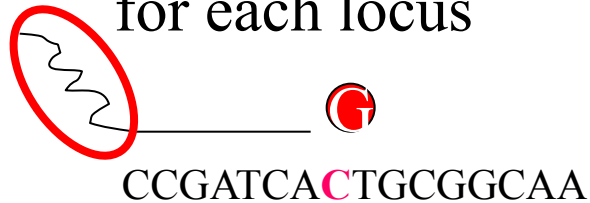
electrophoresis in a capillary  
SNaPShot Multiplex Kit  
(Life Technologies)



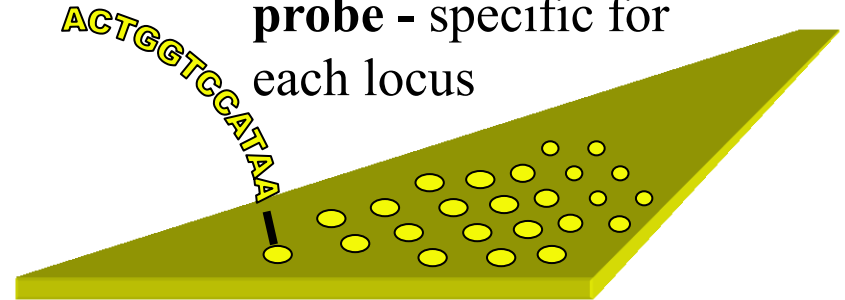
„multiplex version“ - různé  
dlouhé primery, aby bylo  
možné odlišit různé lokusy

# Microarray detection of multiple SBE products

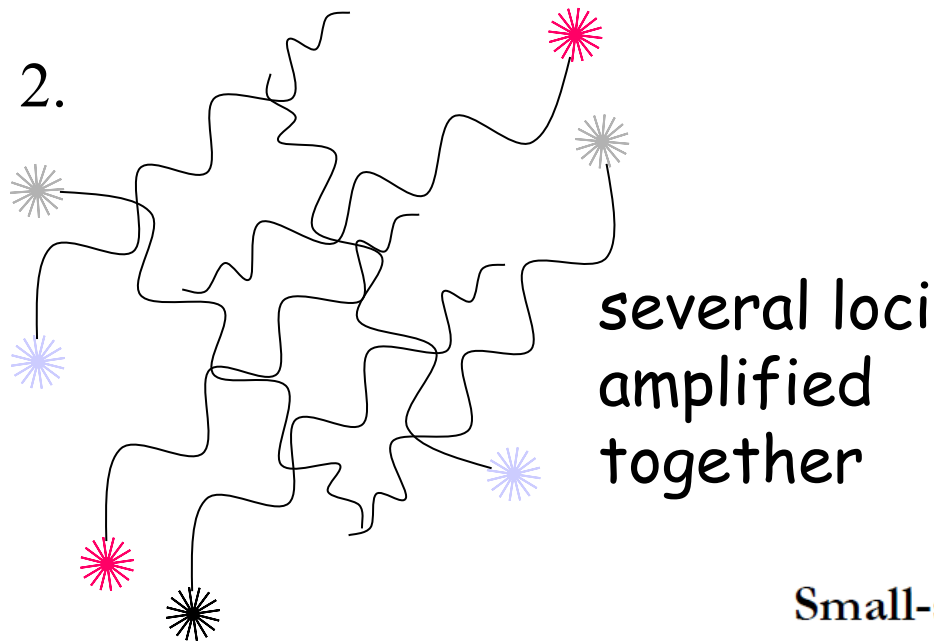
1. **tag** – specific for each locus



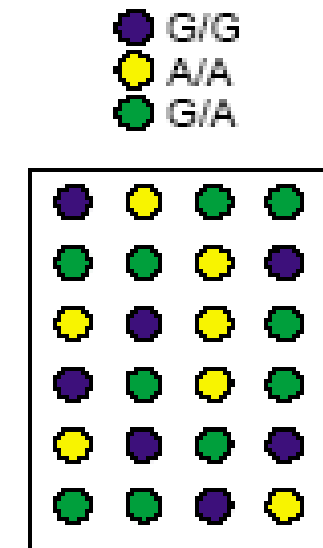
3. **tag-complementary probe** - specific for each locus



- 2.



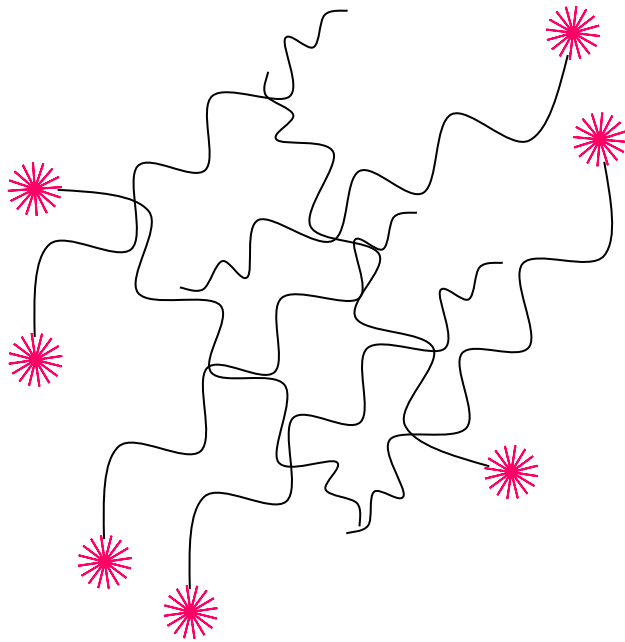
- 4.



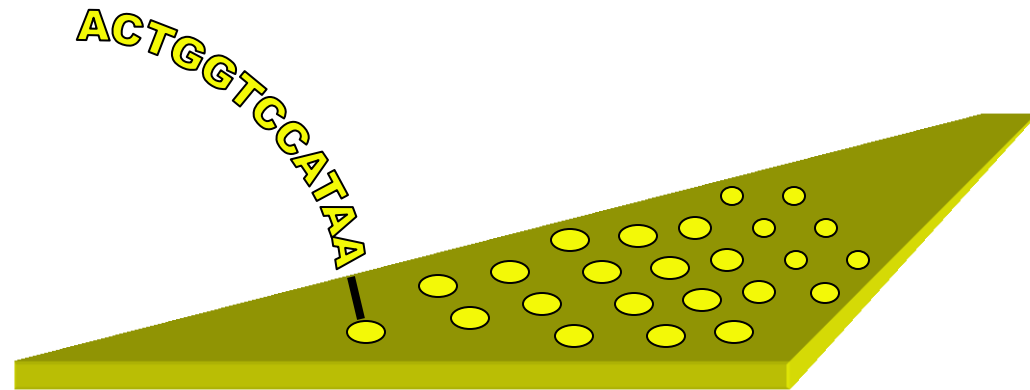
Small-scale “in house” SNP genotyping

multicolor detection (using of 5' oligonucleotide tags on SBE primers)

# (4) Microarray analysis of SNPs (whole genome approach - „chip technology“)

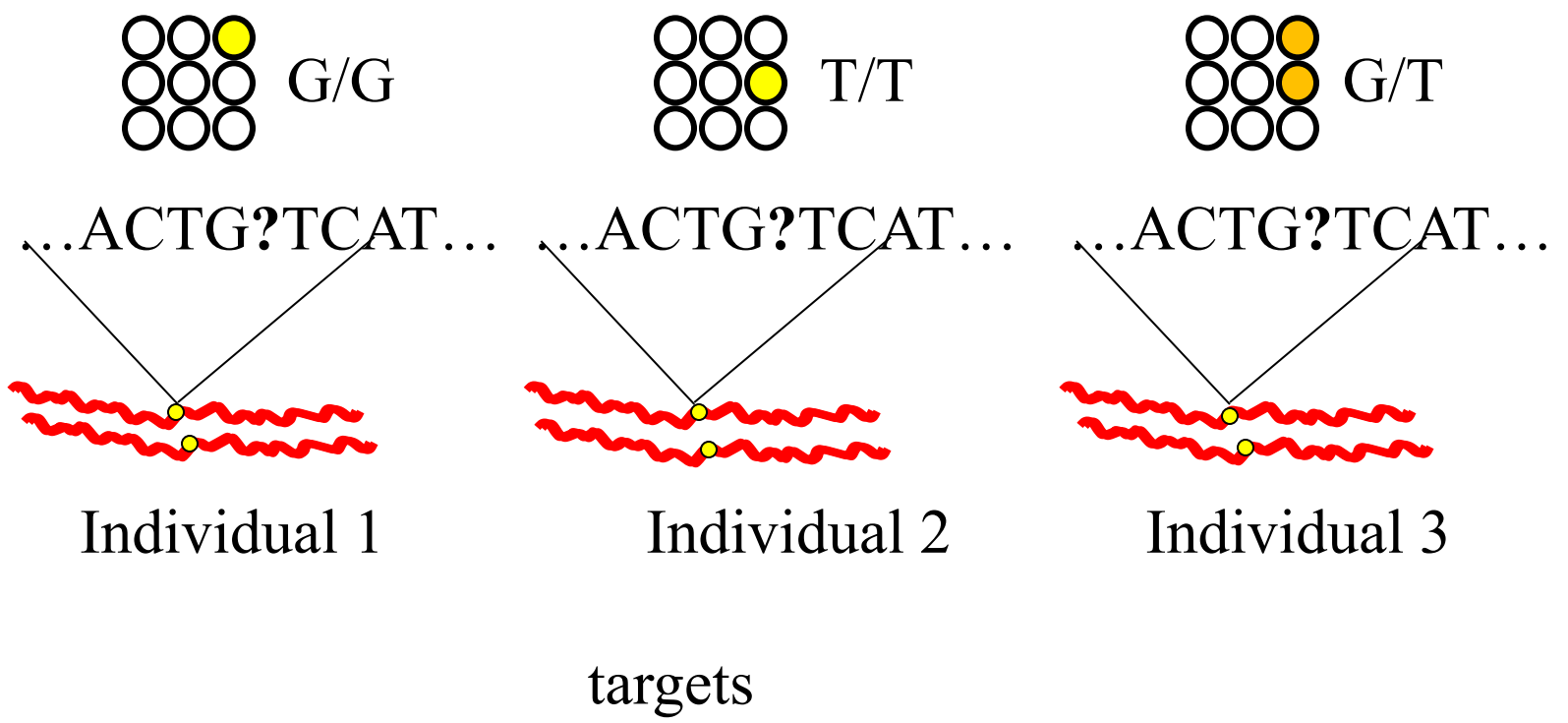


**Target** (genomic DNA  
fragmented by e.g.  
restriction enzymes)



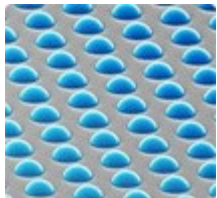
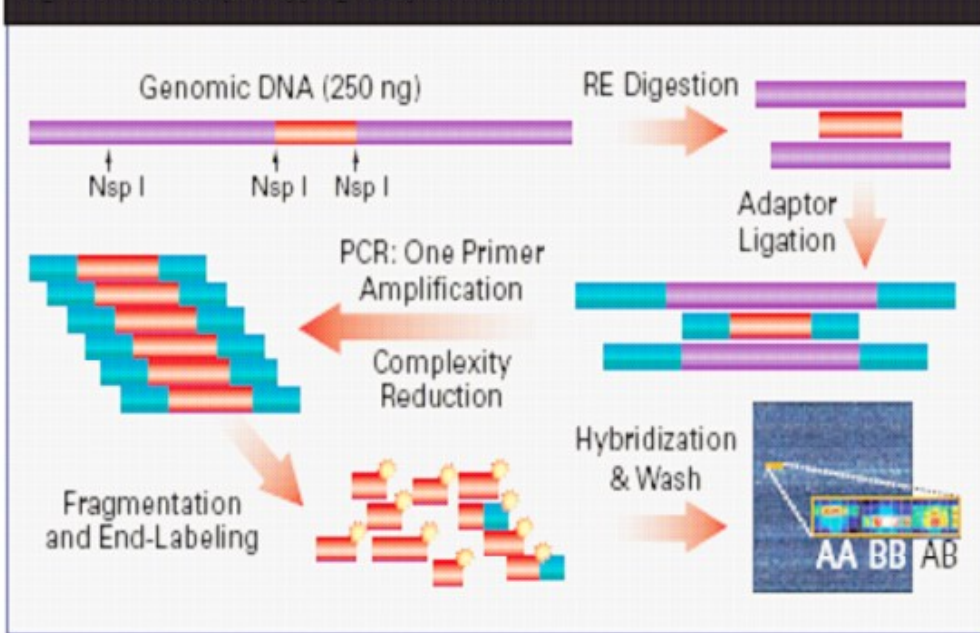
**Probe**  
(specific probes for each allele)

# Microarray SNP Genotyping

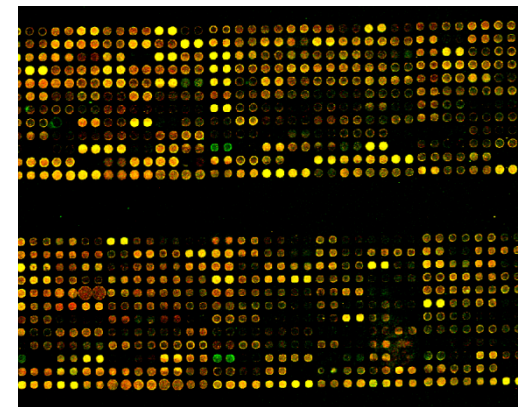
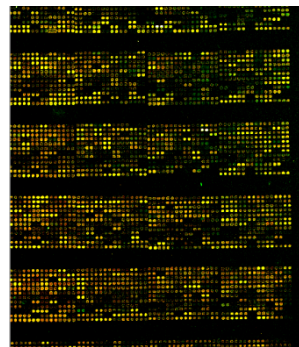


# Detekce: Affymetrix, Illumina

Figure 1: GeneChip® Mapping Assay Overview.



BeadArray  
(Illumina)



10 – 500 tisíc SNP znaků najednou – „chip technology“

Fees - Whole Genome Genotyping									
Platform	SNP multiplex	# samples per array	# genotypes	array \$	reagent \$	core fee \$	Project price per sample	Project price per genotype	volume discount bins
Affymetrix 10K	10,000	1	10,000	185	50	255	\$490.00	\$0.0490	
Affymetrix 50K	50,000	1	50,000	210	50	255	\$515.00	\$0.0103	
Affymetrix 100K (50K x2)	100,000	1	100,000	420	100	510	\$920.00	\$0.0092	
Affymetrix 250K	250,000	1	250,000	470	55	255	\$780.00	\$0.0031	
Affymetrix 500K (250K x2)	500,000	1	500,000	940	110	510	\$1,560.00	\$0.0031	
Affymetrix 500K (250K x2)	500,000	1	500,000	800	110	510	\$1,420.00	\$0.0028	1000-2000 samples
Affymetrix 500K (250K x2)	500,000	1	500,000	700	110	510	\$1,320.00	\$0.0026	2001-5000 samples
Illumina Human-1	109,000	1	109,000	800	na	110	\$910.00	\$0.0083	1-256 samples
Illumina Human-1	109,000	1	109,000	720	na	110	\$830.00	\$0.0076	257-496 samples
Illumina Human-1	109,000	1	109,000	640	na	110	\$750.00	\$0.0069	497-736 samples
Illumina Human-1	109,000	1	109,000	560	na	110	\$670.00	\$0.0061	737-976 samples
Illumina Human-1	109,000	1	109,000	480	na	110	\$590.00	\$0.0054	977+ samples
Illumina HumanHap300	317,000	1	317,000	1100	na	110	\$1,210.00	\$0.0038	1-256 samples
Illumina HumanHap300	317,000	1	317,000	990	na	110	\$1,100.00	\$0.0035	257-496 samples
Illumina HumanHapS	240,000	1	240,000	700	na	110	\$810.00	\$0.0034	737-976 samples
Illumina HumanHapS	240,000	1	240,000	600	na	110	\$710.00	\$0.0030	977+ samples
Illumina HumanHap550	550,000	1	550,000	1600	na	110	\$1,710.00	\$0.0031	1-256 samples
Illumina HumanHap550	550,000	1	550,000	1440	na	110	\$1,550.00	\$0.0028	257-496 samples
Illumina HumanHap550	550,000	1	550,000	1280	na	110	\$1,390.00	\$0.0025	497-736 samples
Illumina HumanHap550	550,000	1	550,000	1120	na	110	\$1,230.00	\$0.0022	737-976 samples
Illumina HumanHap550	550,000	1	550,000	960	na	110	\$1,070.00	\$0.0019	977+ samples
HumanHap300 + HumanHapS	550,000	1	550,000	1750	na	220	\$1,970.00	\$0.0036	1-256 samples
HumanHap300 + HumanHapS	550,000	1	550,000	1575	na	220	\$1,795.00	\$0.0033	257-496 samples
HumanHap300 + HumanHapS	550,000	1	550,000	1400	na	220	\$1,620.00	\$0.0029	497-736 samples
HumanHap300 + HumanHapS	550,000	1	550,000	1225	na	220	\$1,445.00	\$0.0026	737-976 samples
HumanHap300 + HumanHapS	550,000	1	550,000	1050	na	220	\$1,270.00	\$0.0023	977+ samples

Použití u příbuzných druhů je možné, ale je tam velmi silný „ascertainment bias“