

Data QC and Exploratory Data Analysis (Morning session)

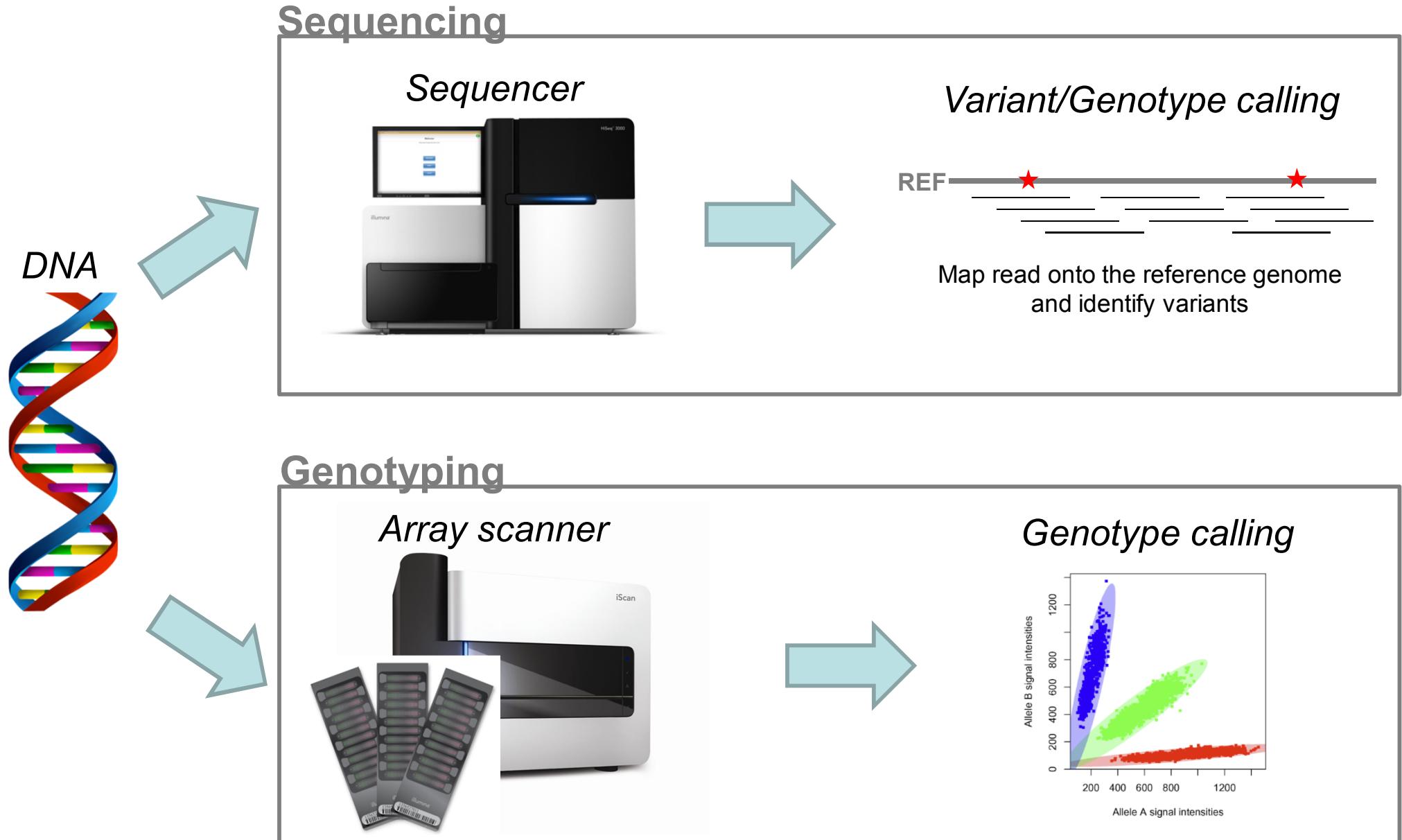
Olivier Delaneau
University of Geneva

19/05/2017

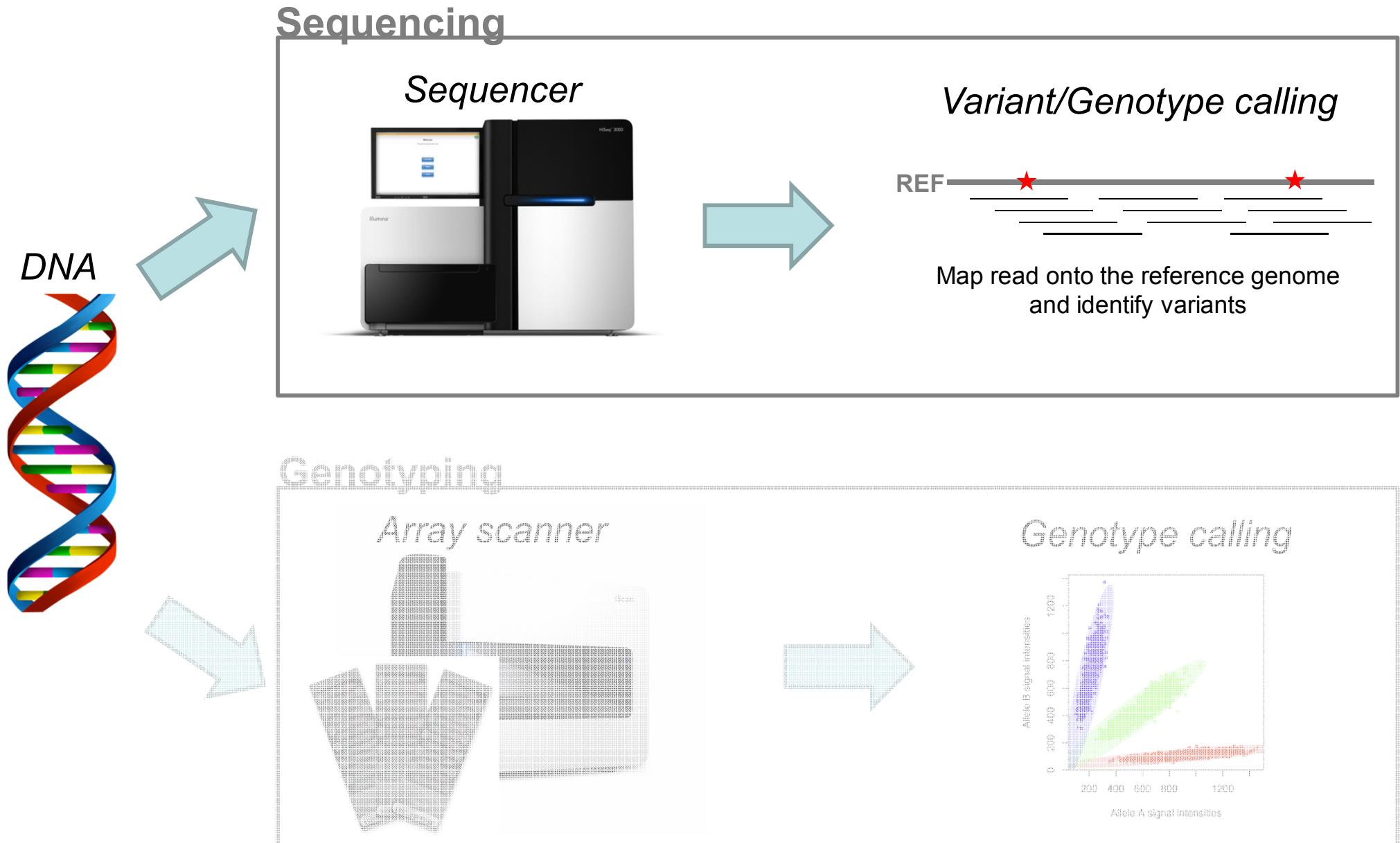
Outline

- Methods for genotyping
- File format for genotype data and standard data management tools
- QC at the genetic variant level
- QC at the sample level
- Population structure
- Information on the practical

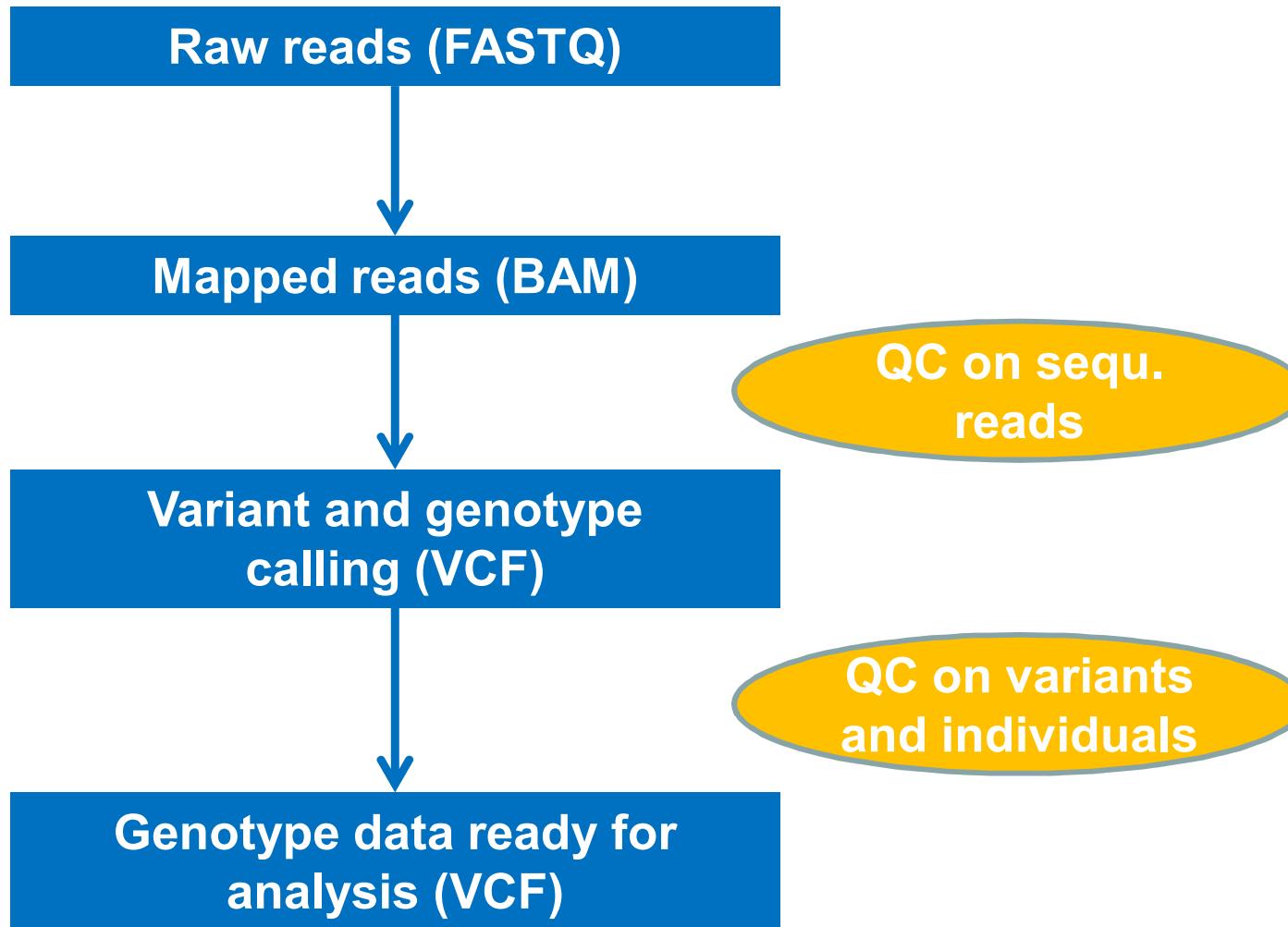
How to read genetic variants?



How to read genetic variants?



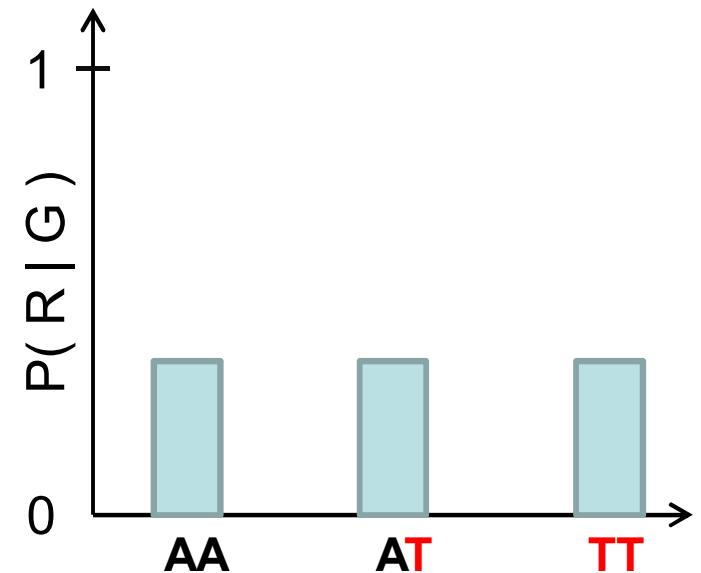
Calling genetic variations



Computing genotype likelihoods

REF

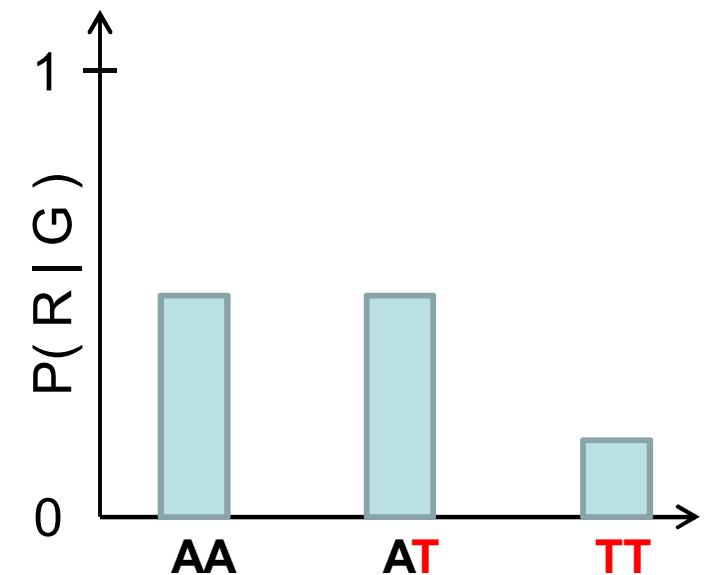
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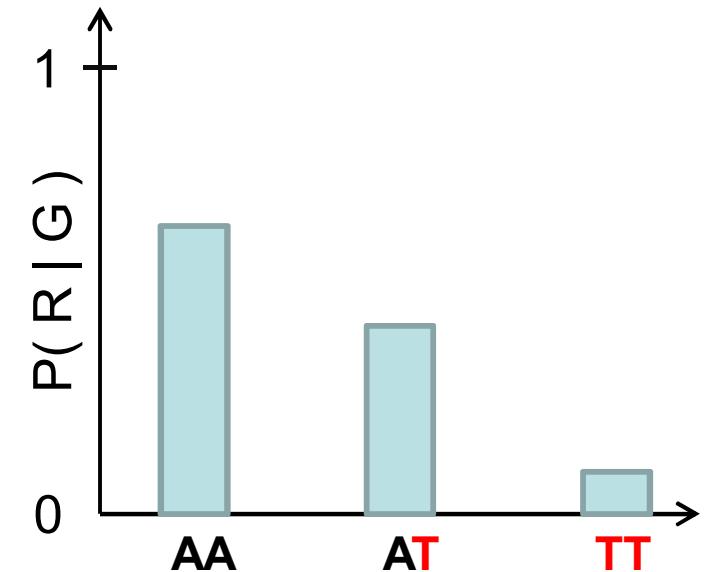
Computing genotype likelihoods

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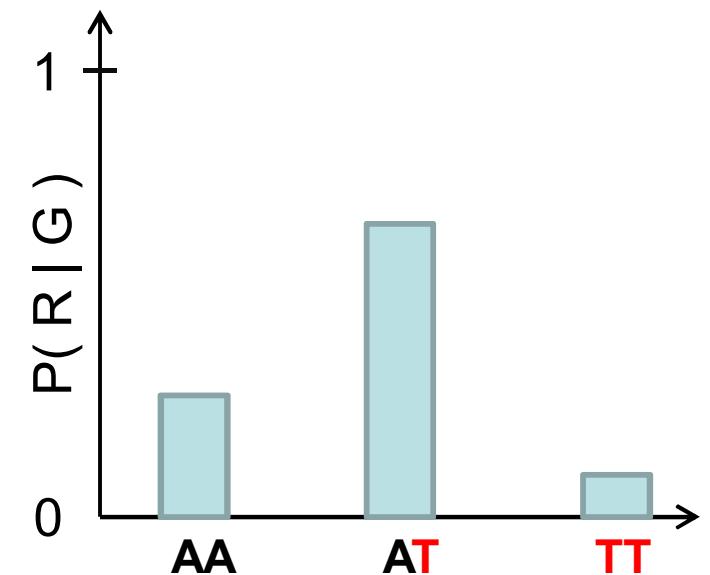
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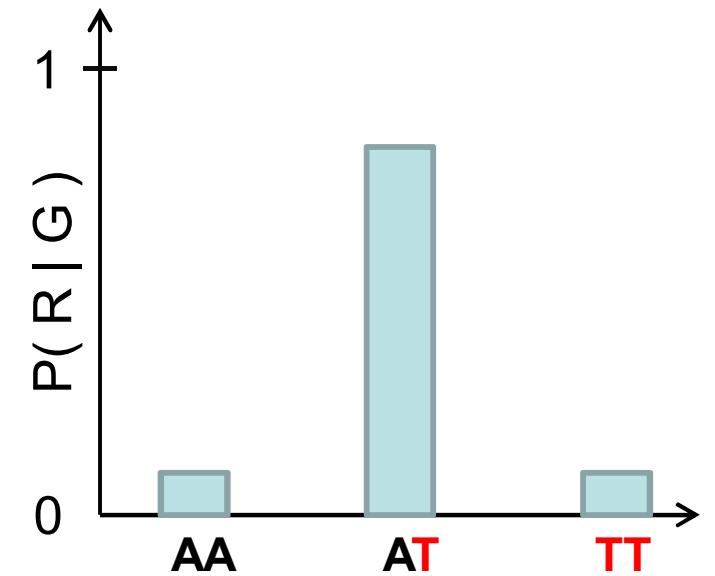
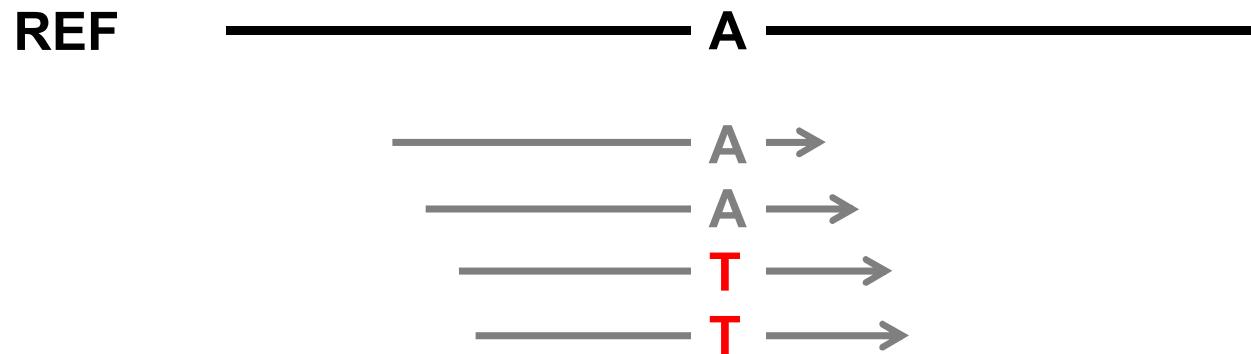
Computing genotype likelihoods



Computing genotype likelihoods



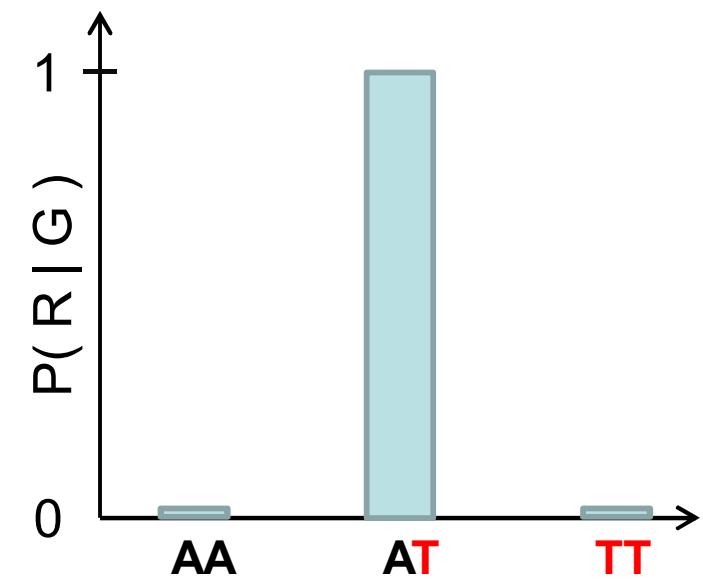
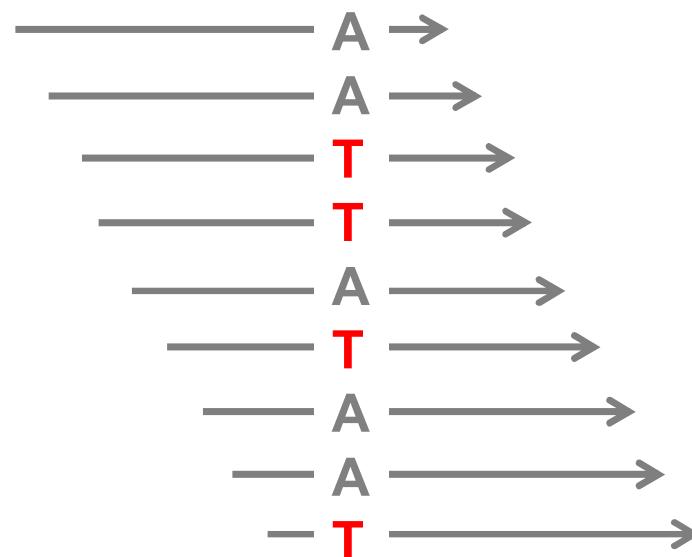
Computing genotype likelihoods



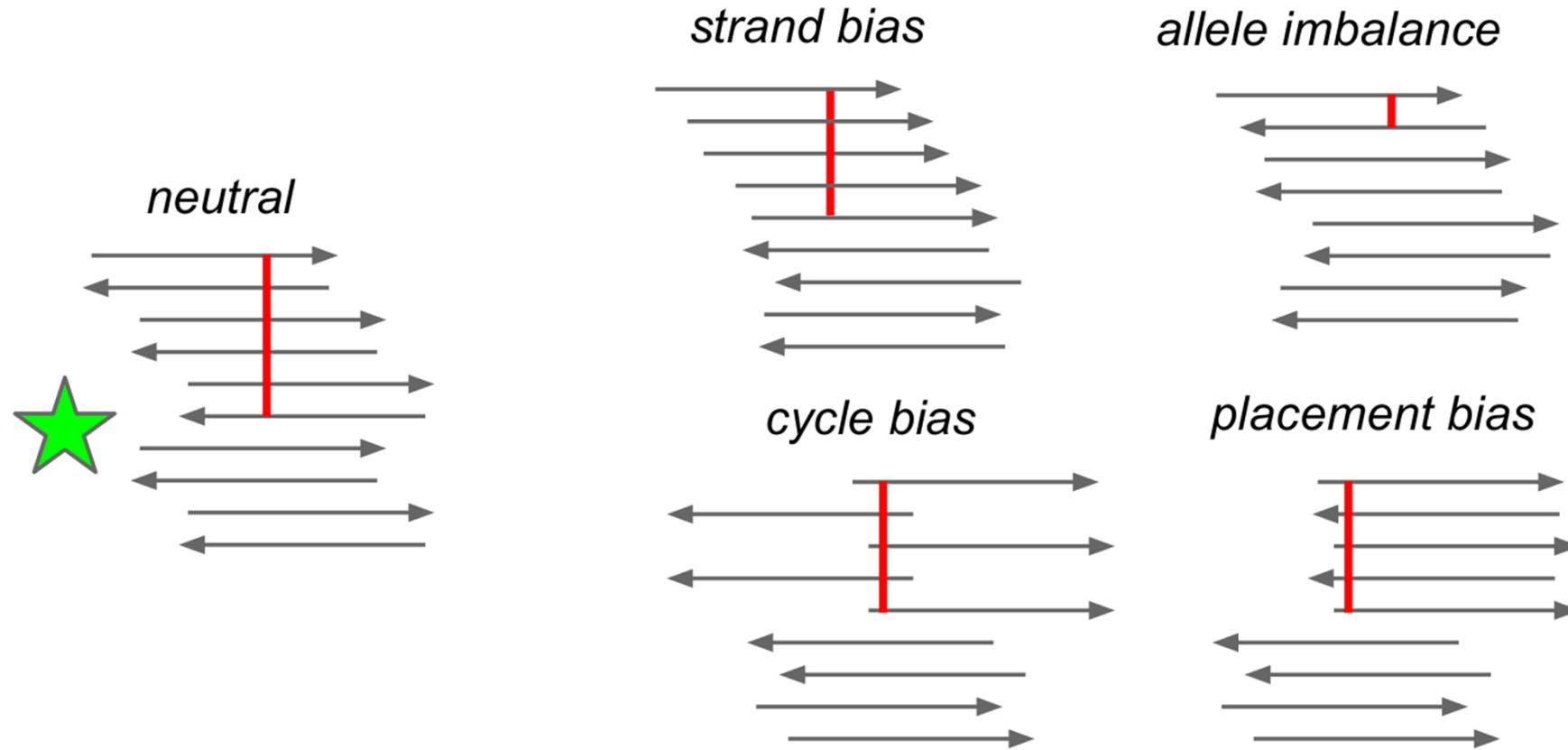
Computing genotype likelihoods

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Accounting for sequencing biases



These biases are either modeled in the genotype likelihoods (e.g. *FreeBayes*) or variants can be filtered out after calling (e.g. *samtools*) by statistical testing.

Refining genotype calls

$$P(R | G)$$

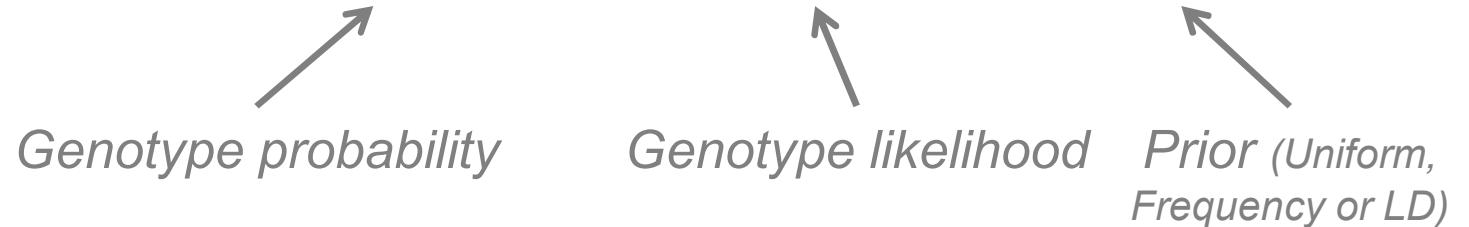


Genotype likelihood

Refining genotype calls

$$P(G | R) \sim P(R | G) \times P(G)$$

Genotype probability *Genotype likelihood* *Prior (Uniform, Frequency or LD)*

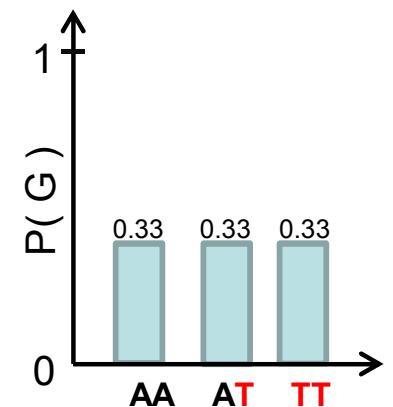
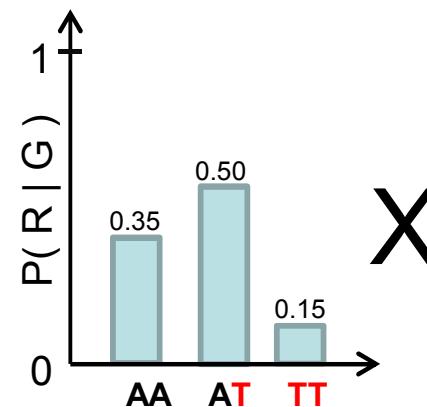
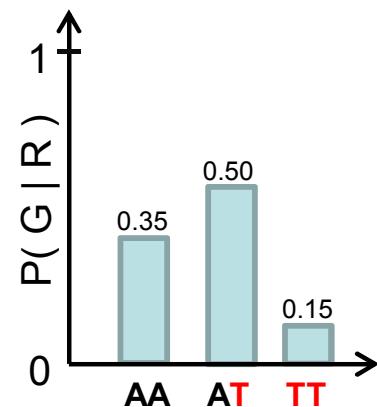


Refining genotype calls

$$P(G|R) \sim P(R|G) \times P(G)$$

Genotype probability Genotype likelihood Prior (Uniform, Frequency or LD)

Uniform prior:



Refining genotype calls

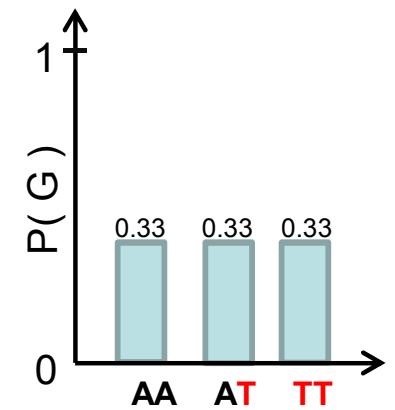
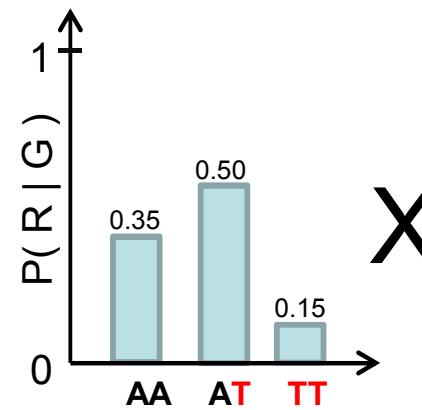
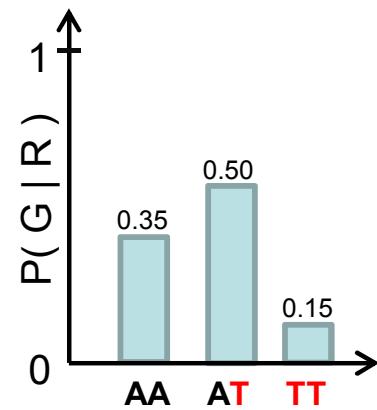
$$P(G|R) \sim P(R|G) \times P(G)$$

Genotype probability

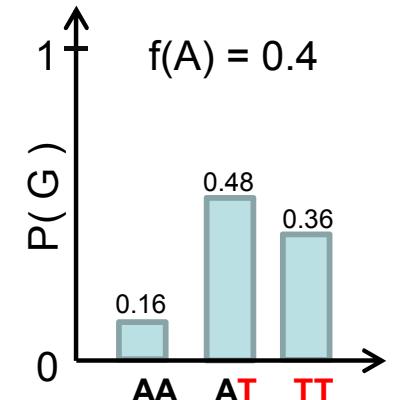
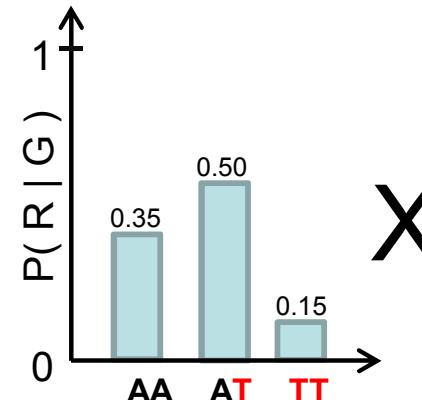
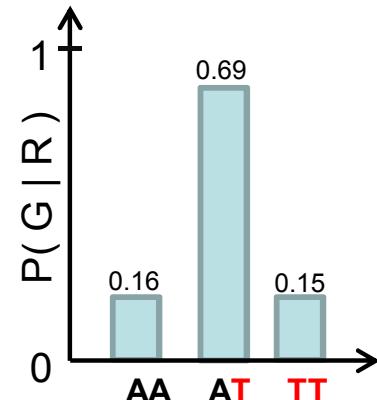
Genotype likelihood

Prior (Uniform,
Frequency or LD)

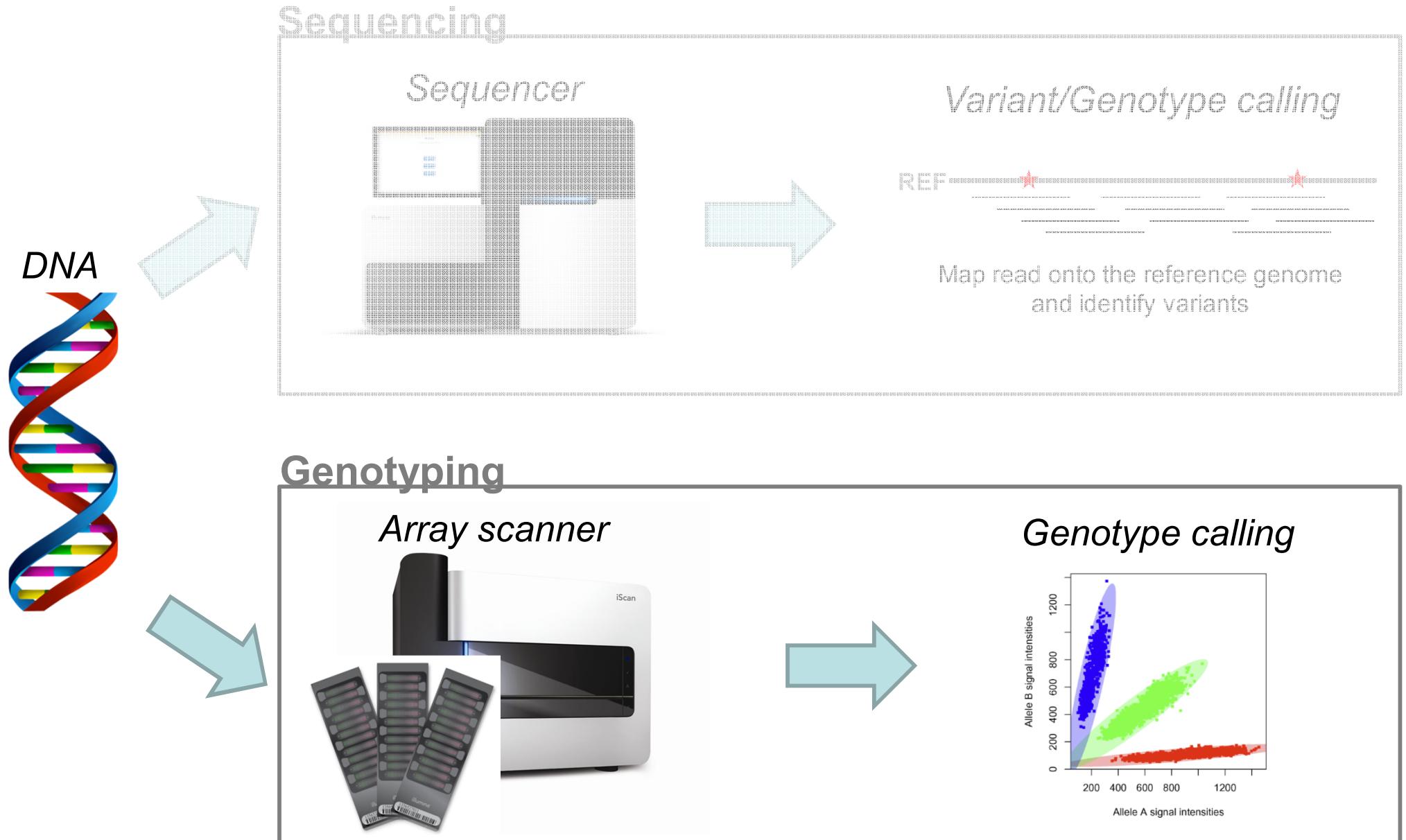
Uniform prior:



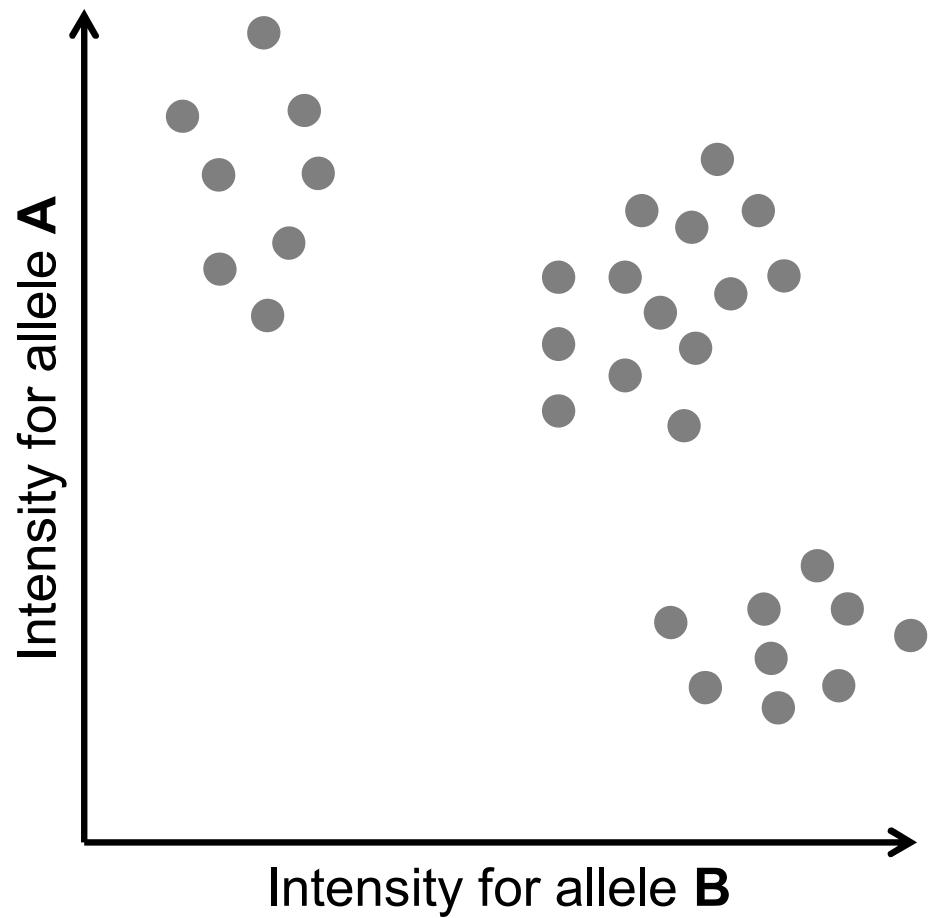
Frequency prior:
(samtools)



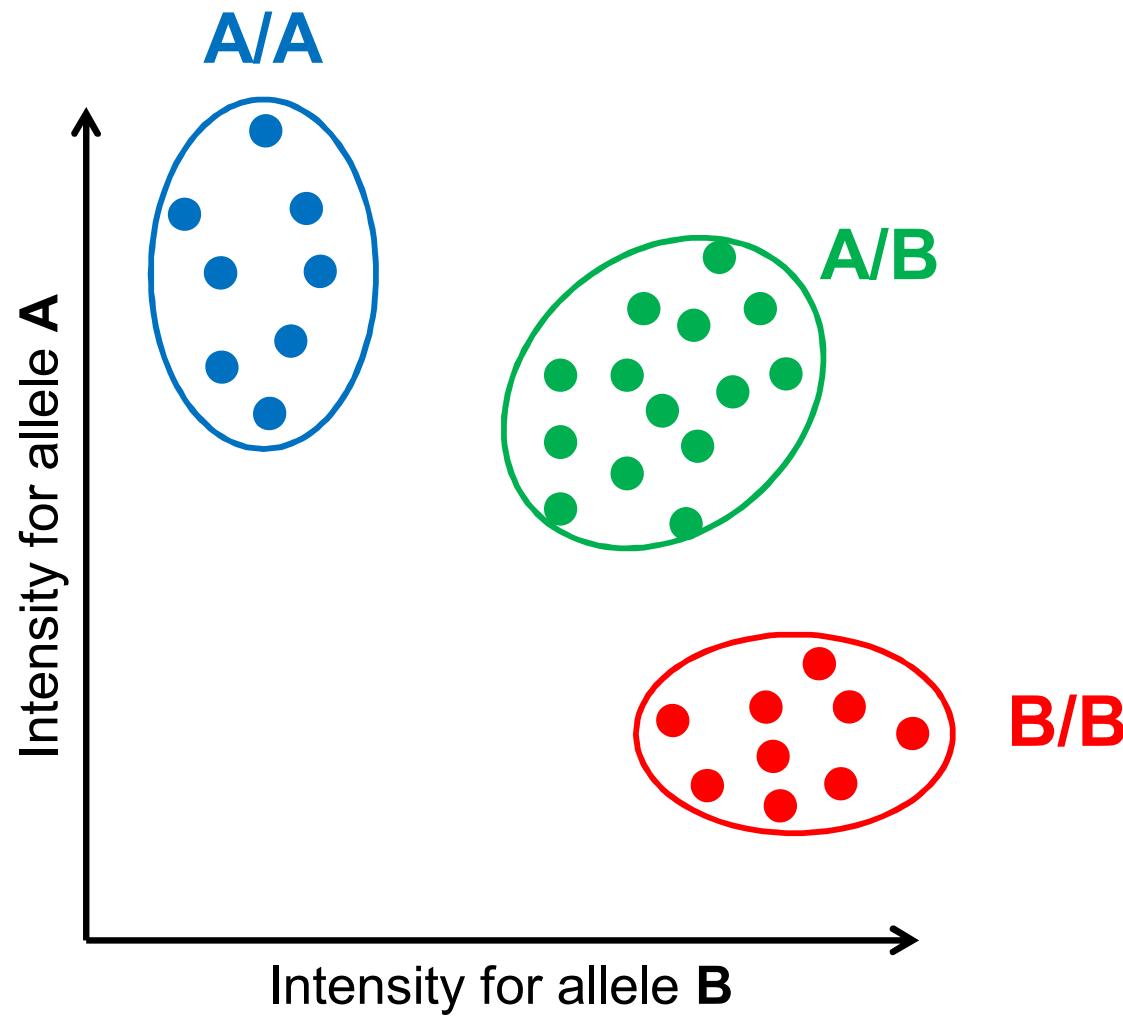
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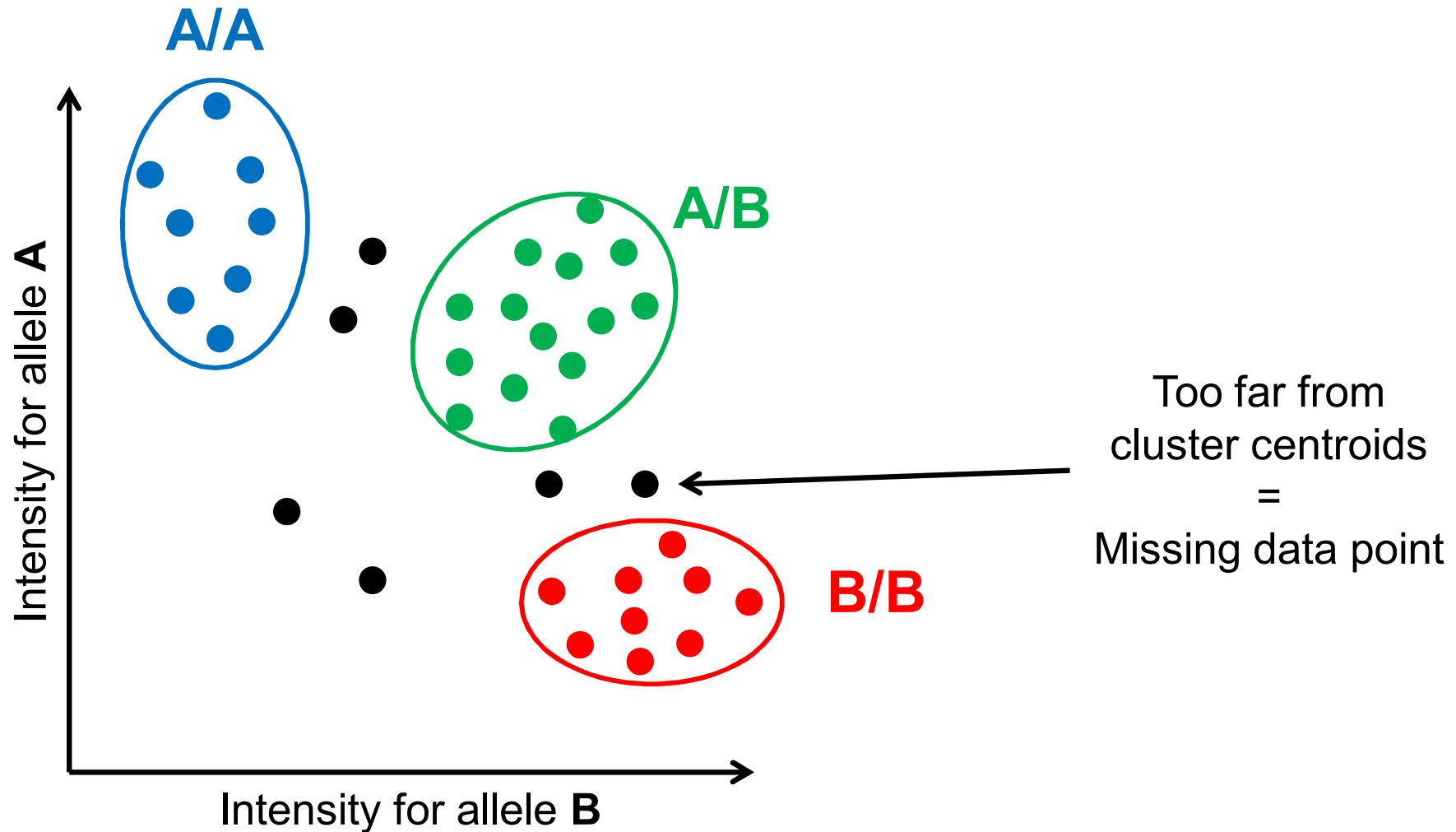
Genotype calling from chips



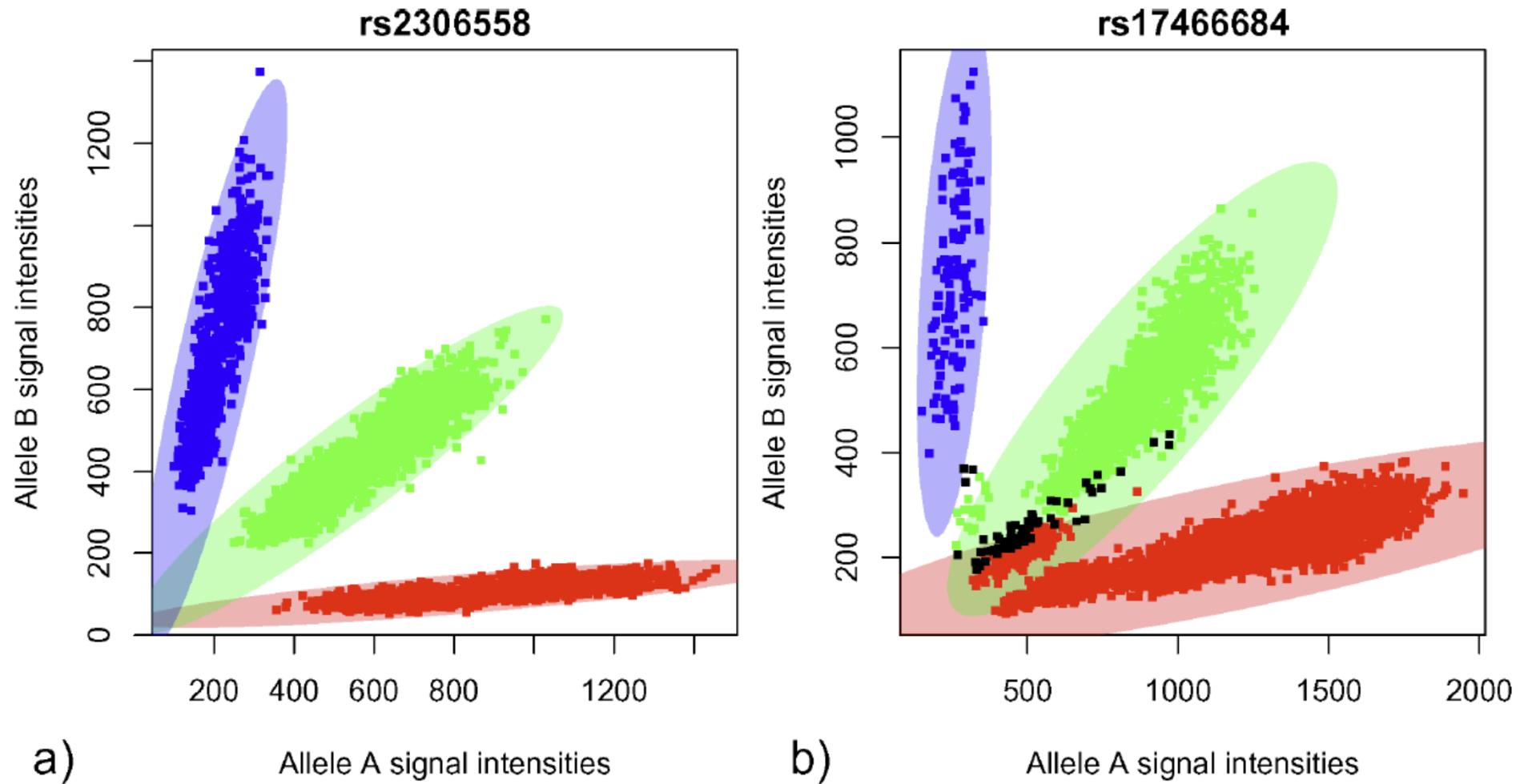
Genotype calling from chips



Genotype calling from chips

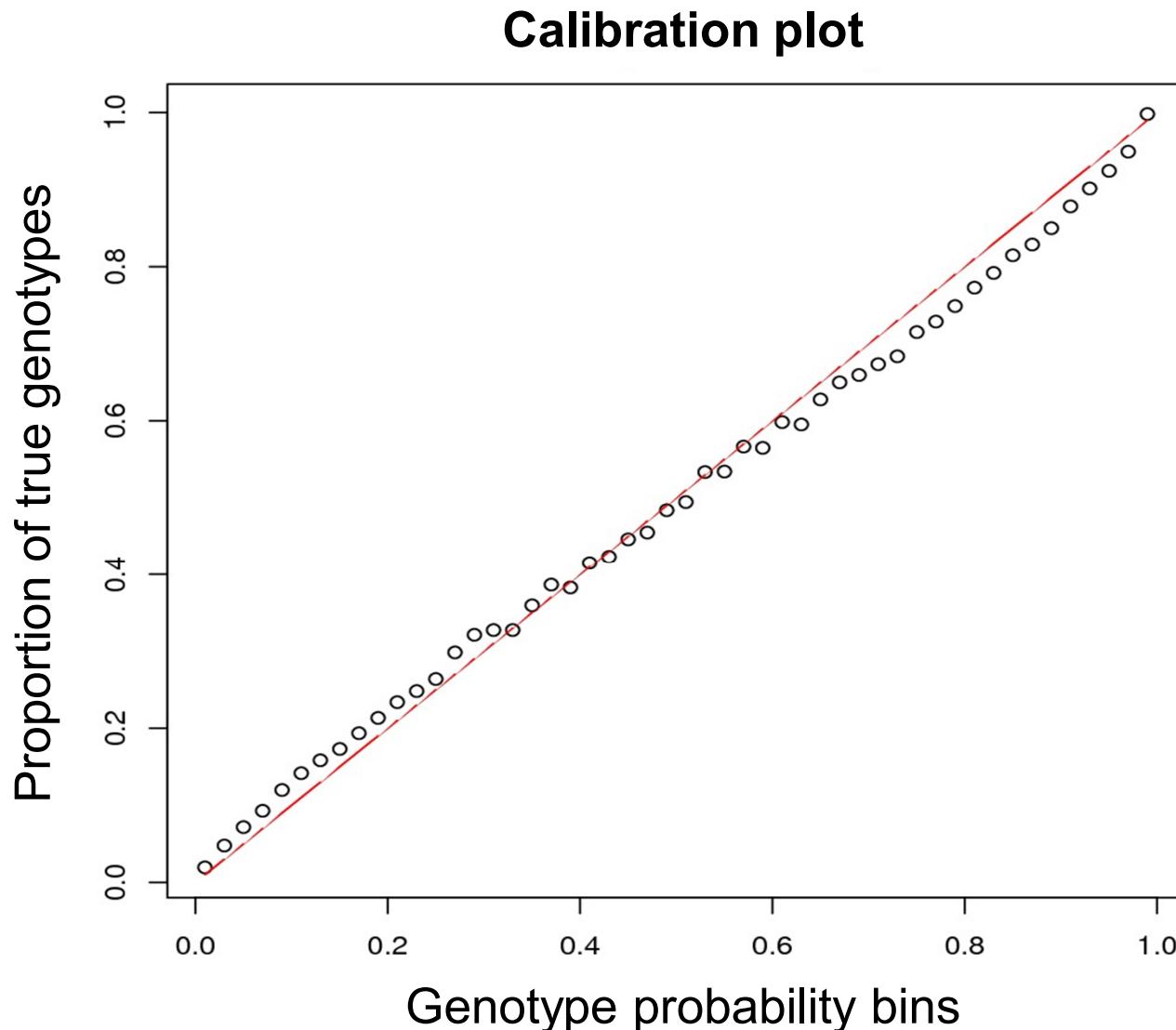


Genotype calling from chips



Approaches to call genotypes from intensities (GenomeStudio, APCA or Chiamo)
Automated approaches to call genotypes at millions of SNPs prior to the analysis.
Any finding need to be checked manually and carefully.

What does genotype probability mean?



We usually set as missing all genotypes with low prob. The threshold we use depends on the study design.

Can we afford some incorrect calls? Two scenarios:

1. Interest in global patterns, a value of 0.95 is okay,
2. Interest in some particular variants, a value 0.999 is more adapted.

The goal of Quality Control

- Errors in genotype calling can introduce systematic bias in downstream analysis and can increase the chance of false discoveries
- Assess data quality to remove sub-standard genotypes, individuals and variants from subsequent analysis
- To do so, there are standardized QC procedures implemented in well-established software packages

Variant Call Format

Variants

##fileformat=VCFv4.2
##FILTER=<ID=PASS,Description="All filters passed">
##simulateGenotypeData=1.1
##source=simulateGenotypeDataFrom1000G
##FORMAT=<ID=GT,Number=1,Type=String,Description="Genotype">
##contig=<ID=20>

#CHROM	POS	ID	REF	ALT	QUAL	FILTER	INFO	FORMAT	HG01124	HG01363	HG01500	HG01501	HG01503
20	61651	rs76553454	C	A	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	63231	rs6076506	T	G	100	PASS	.	GT	0/1	0/0	0/0	0/0	0/0
20	63244	rs6139074	A	C	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	63799	rs1418258	C	T	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	68749	rs6086616	T	C	100	PASS	.	GT	0/1	1/1	0/1	1/1	0/0
20	69094	rs6039403	G	A	100	PASS	.	GT	0/0	0/0	1/1	0/0	0/0
20	69408	rs17685809	C	T	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	71093	rs6040395	G	A	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	74347	rs6135141	G	A	100	PASS	.	GT	0/0	0/0	0/1	0/1	0/0
20	75254	rs892665	C	A	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	80655	rs2196239	A	G	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	83611	rs114000219	C	A	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	87112	rs34383360	G	A	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	87416	rs1935386	A	C	100	PASS	.	GT	0/1	1/1	1/1	0/1	0/0
20	88155	rs78720032	T	C	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	90814	rs75507632	T	C	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0
20	90984	rs1077784	A	G	100	PASS	.	GT	0/0	0/1	0/0	0/0	0/0
20	91088	rs2002370	C	T	100	PASS	.	GT	0/1	1/1	1/1	0/1	0/0
20	91346	rs11698455	G	A	100	PASS	.	GT	0/0	0/1	0/0	0/0	0/0
20	91508	rs6051659	G	A	100	PASS	.	GT	0/1	1/1	1/1	1/1	0/0
20	92366	rs13039134	A	G	100	PASS	.	GT	0/1	0/1	1/1	0/1	0/0
20	96931	rs6052070	A	G	100	PASS	.	GT	0/0	0/1	0/0	0/0	0/0
20	97122	rs6515824	C	T	100	PASS	.	GT	0/0	0/1	0/0	0/0	0/0
20	98930	rs6116135	G	A	100	PASS	.	GT	0/0	0/1	0/0	0/0	0/0
20	100505	rs6037772	T	C	100	PASS	.	GT	0/0	0/0	0/0	0/0	0/0

Position Reference Alternative Quality Annotation Genotypes

Chromosome Variant Genotype field

Sample IDs

HEADER

HG01124 has genotype TC at rs6086616

Missing genotype

Basic operations on VCF

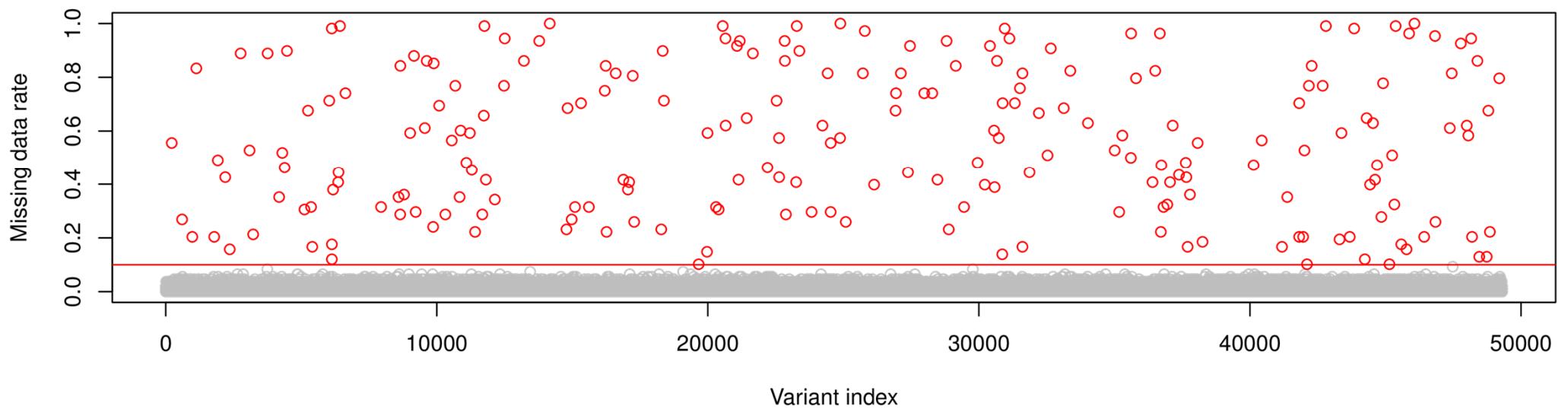
- BCFtools (www.htslib.org)
 - Efficient data management
 - Excellent API to develop new tools
- VCFtools (vcftools.sourceforge.net)
 - Standard data analysis (frequency, LD, etc...)
- PLINK1.9 (www.cog-genomics.org/plink2)
 - Efficient implementation of all PLINK functionalities (e.g. association testing)

Filter variants with low call rates

Apply threshold to genotype probabilities in order to call genotype, otherwise treated as missing. The choice of calling threshold will impact results:

- Too low: include poor quality genotypes.
- Too high: unnecessarily remove high quality genotypes, or may introduce bias by preferentially calling specific genotypes.

Variants with poor call rates (i.e. high missing data rates) likely result from poor genotype calling. They need to be removed from the data.

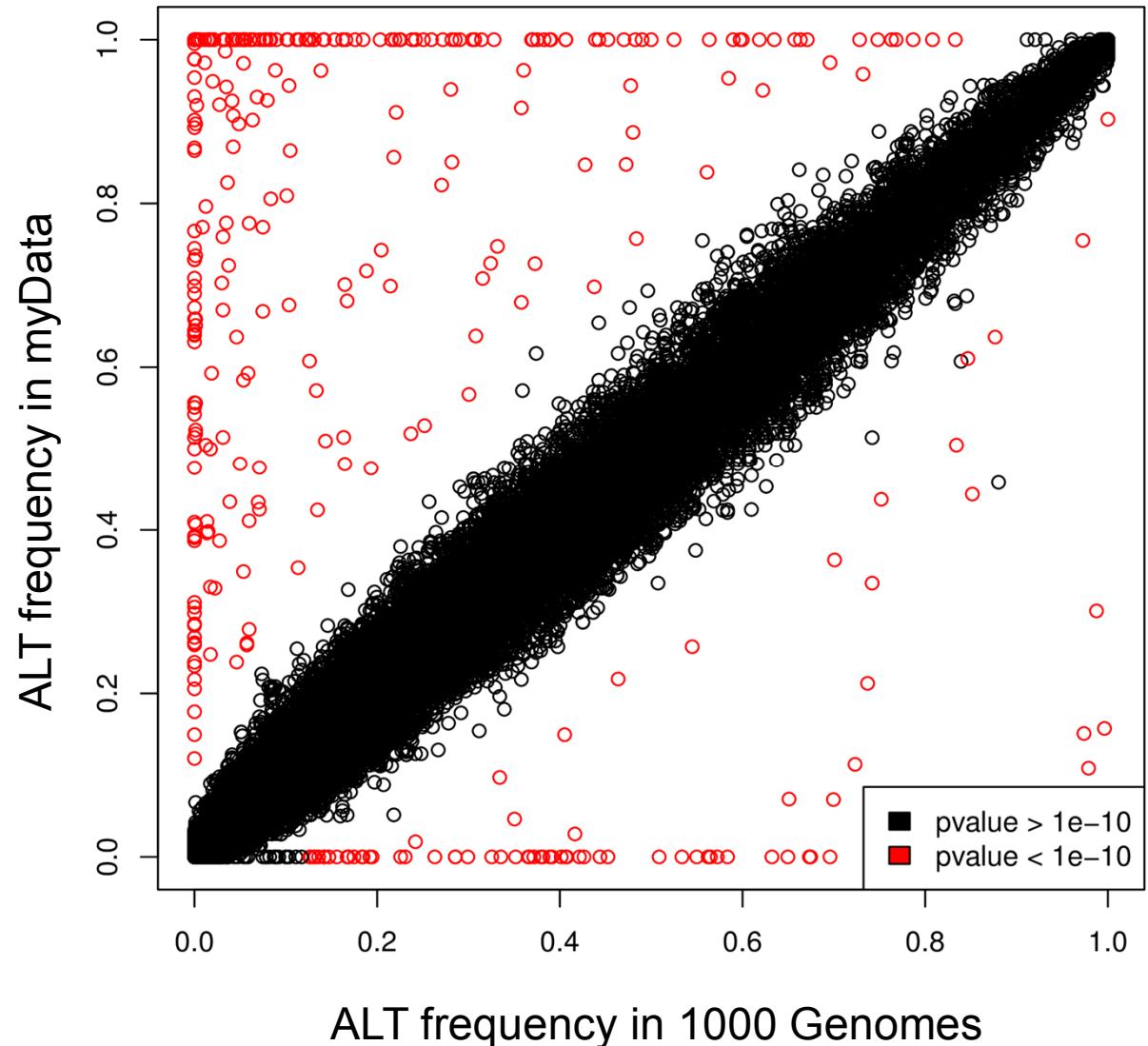


Check variant frequencies

To detect genotyping errors, one can also use large and high quality reference panels such as 1000 Genomes.

We compare the variant frequencies in our data to the frequencies in 1000 Genomes from a relevant population.

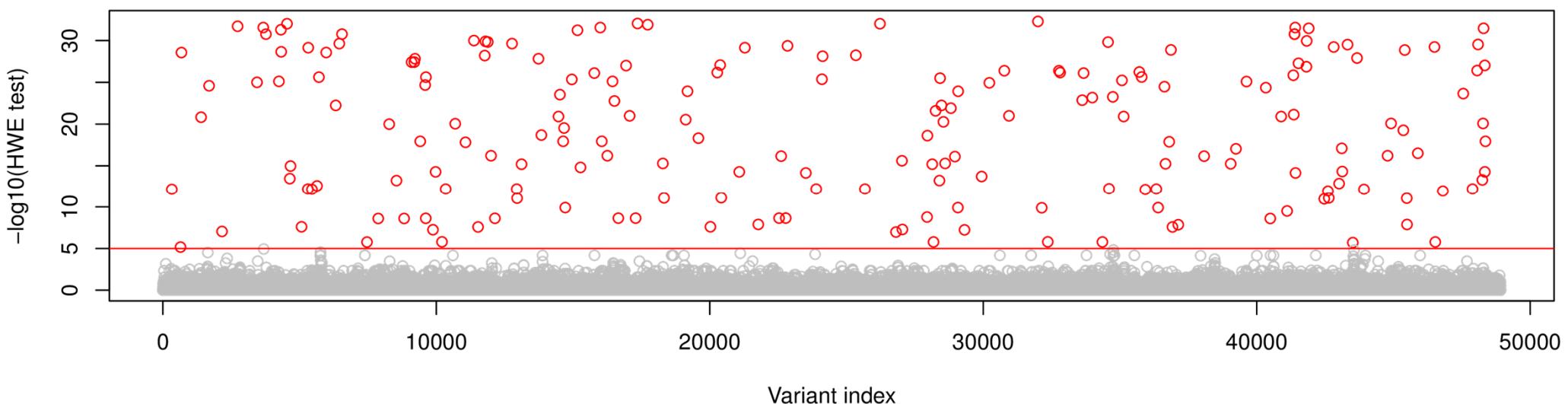
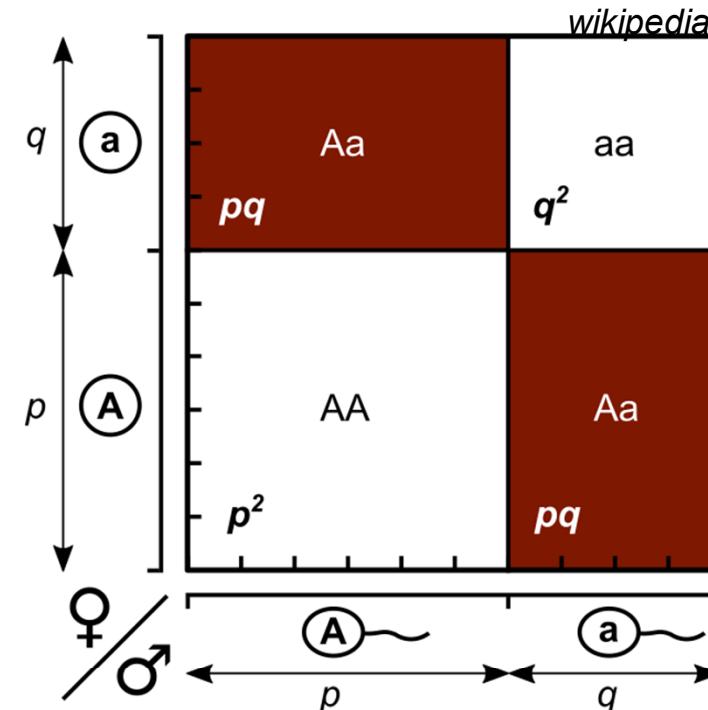
Then, one can test for massive differences and remove the corresponding variants.



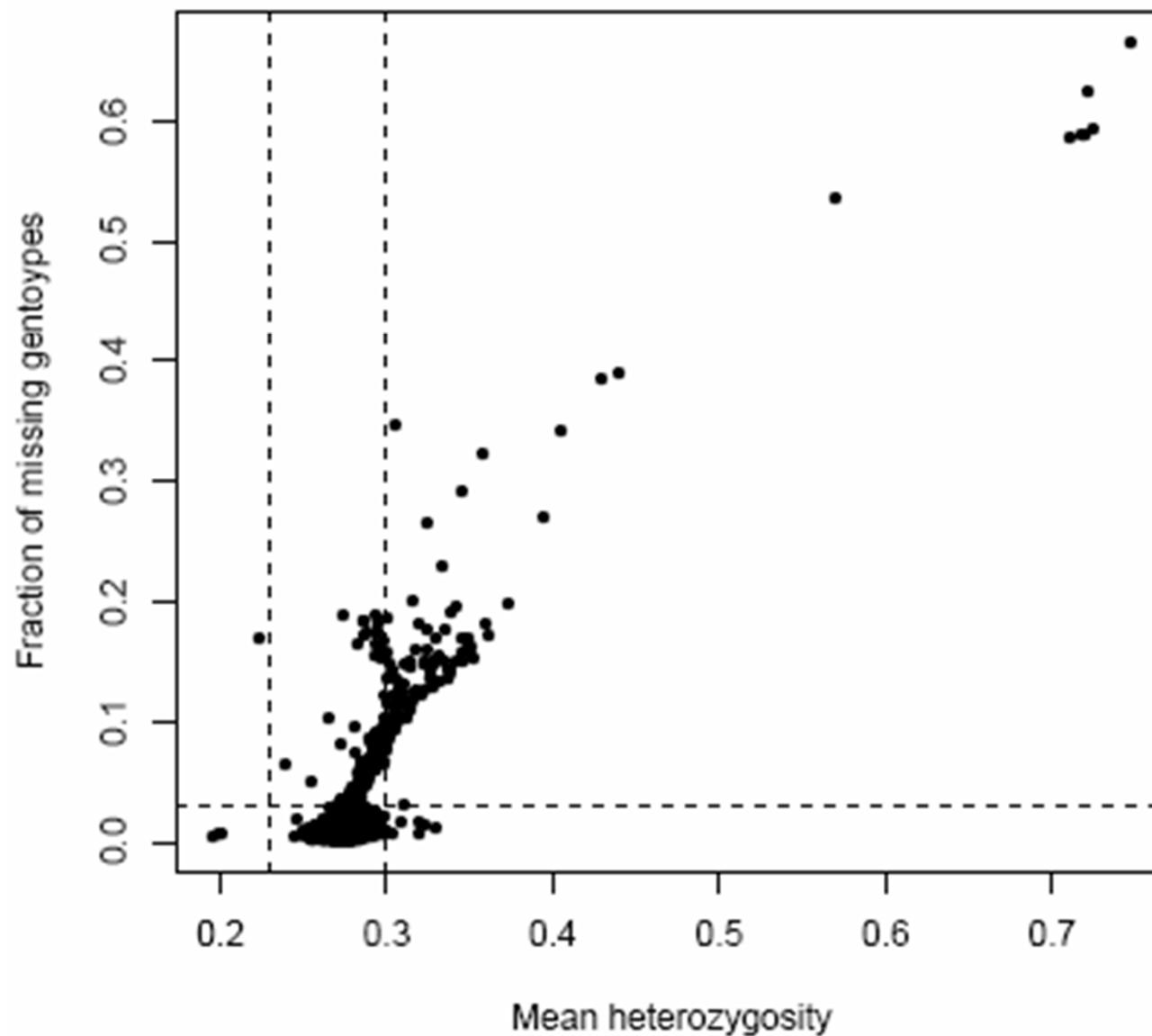
Check Hardy Weinberg Equilibrium

Under the assumption of random mating between individuals in a population, we have the following relationships between allele frequencies and genotype frequencies.

This can also be used to assess the quality of the genotype data by statistical testing (*Wigginton, Cutler and Abecasis, 2005*).



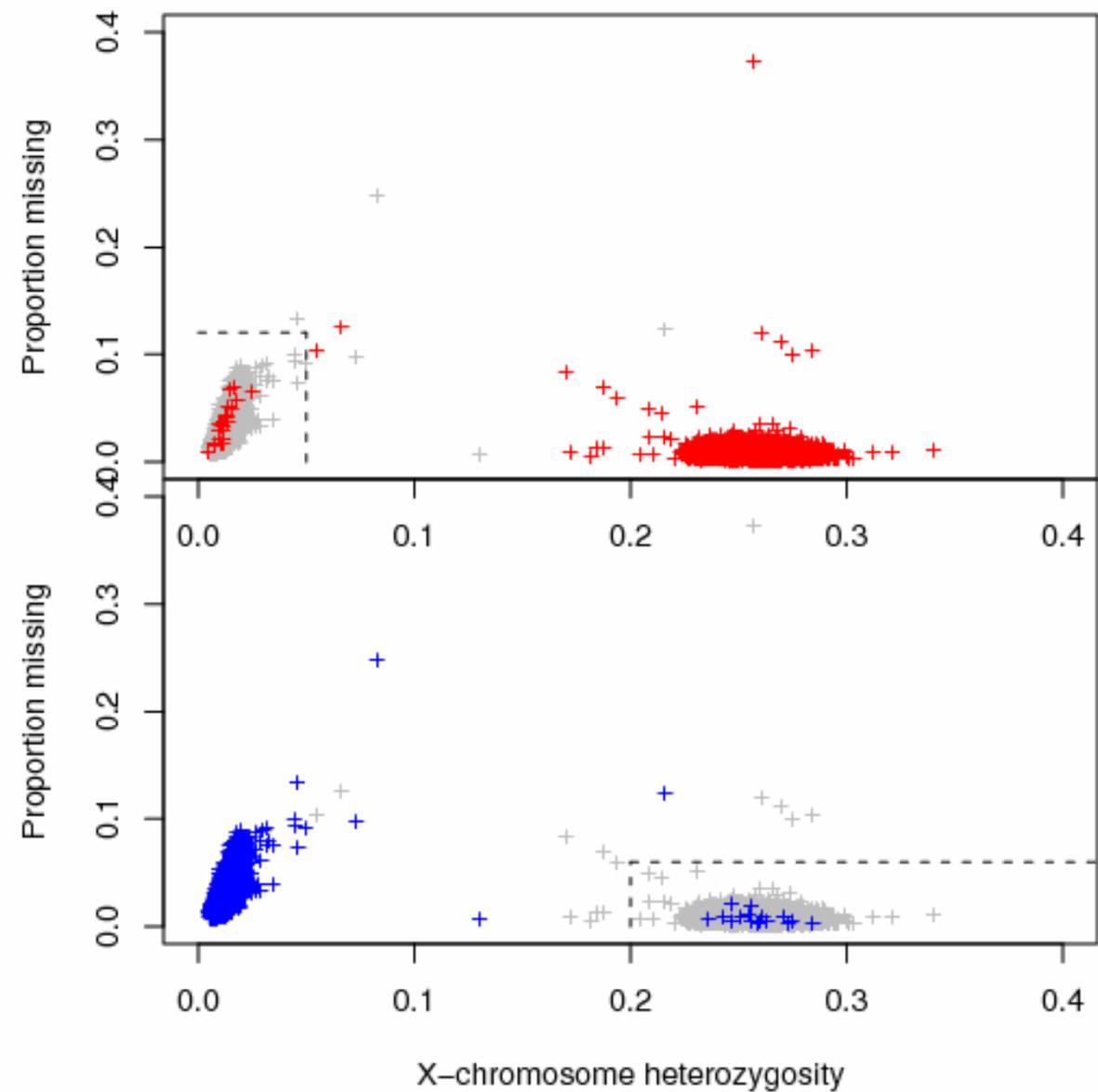
Filter samples with low call rate



Gender check with chromosome X

Each individual plotted twice according to reported gender:
females in red and **males in blue**.

Should these samples be removed from the study or the sex corrected based on heterozygosity? May impact on results if sex is adjusted for in the analysis or if sex specific analyses are to be undertaken.



Check relatedness

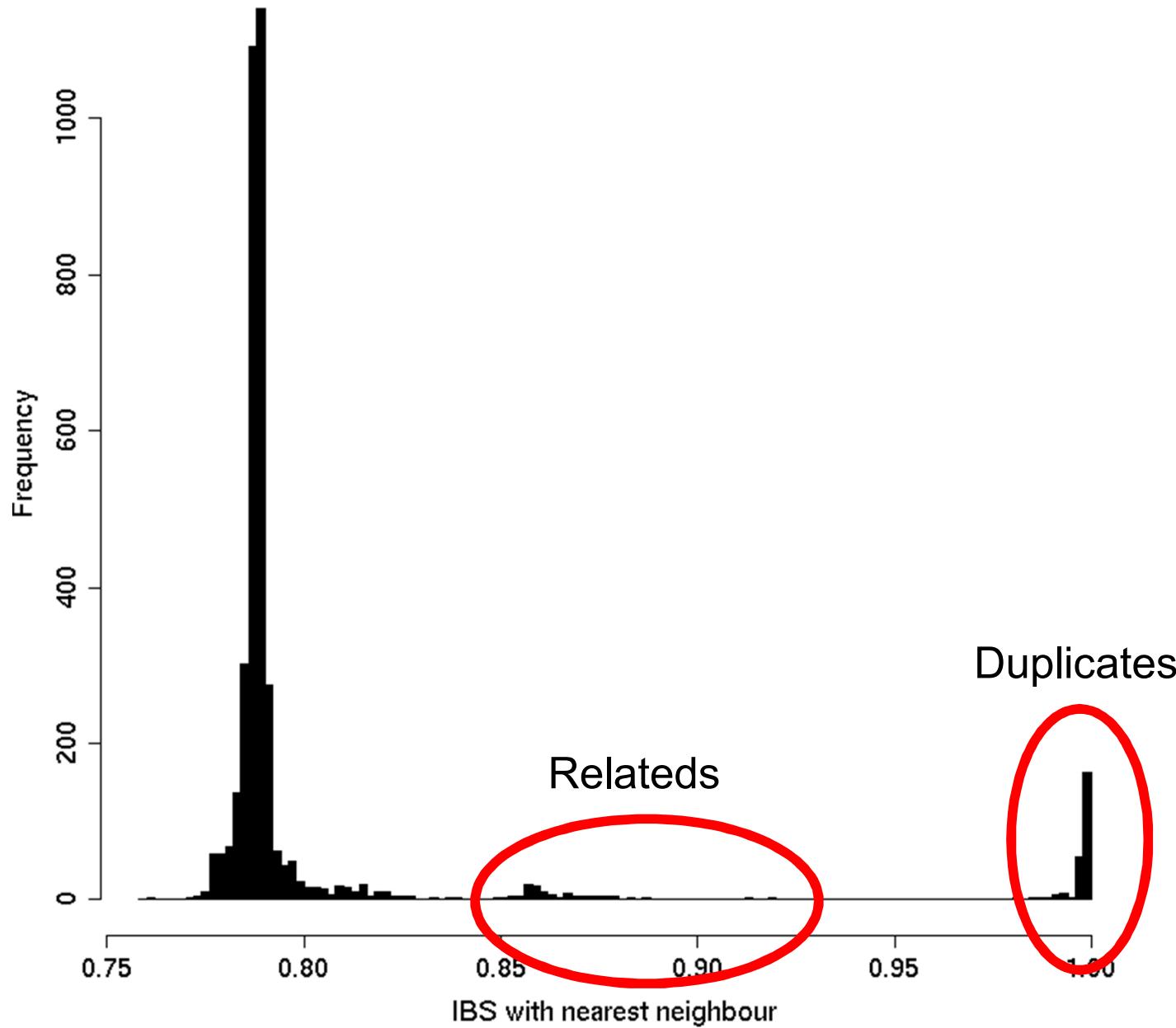
- Over M markers, the IBS between the i th and j th individuals is given by

$$IBS_{ij} = 1 - \frac{1}{2M} \sum_k |G_{ik} - G_{jk}|$$

where G_{ik} and G_{jk} denote the number of minor alleles (0, 1 or 2) carried by the i th and j th individuals at variant k .

- Identical samples will share IBS near to 100% (allowing for genotyping errors).
- Related individuals will share higher IBS than unrelated individuals.

Check relatedness

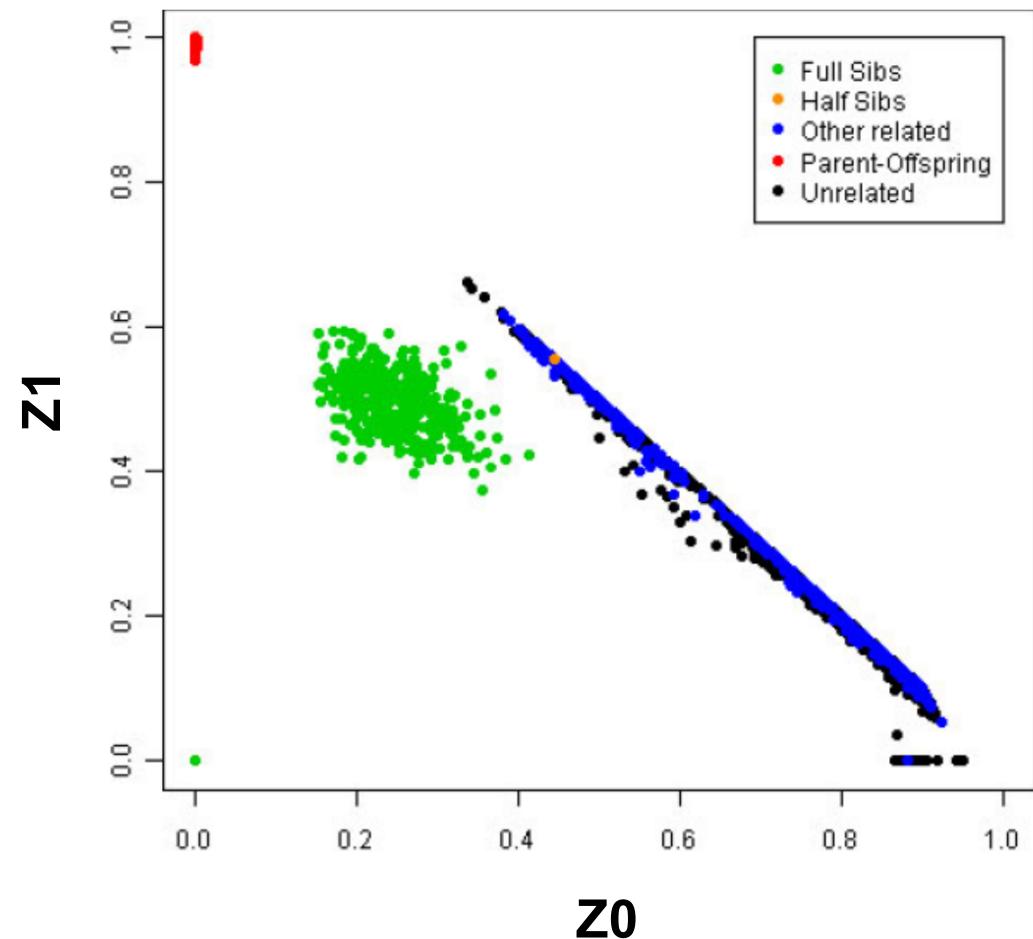


Alternative checks for relatedness

IBS can be used to estimate the proportion of the genome at which a pair of individuals share 0, 1 or 2 chromosomes IBD “identical by descent” (denoted z_0 , z_1 , or z_2).

Once relative spotted, two options:

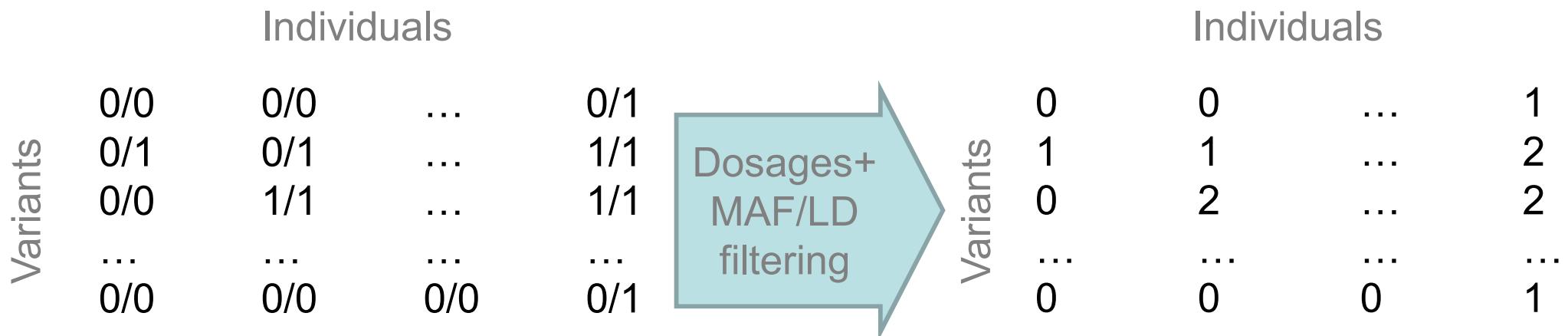
1. Accounting for relatedness in downstream analysis,
2. Remove close relatives.



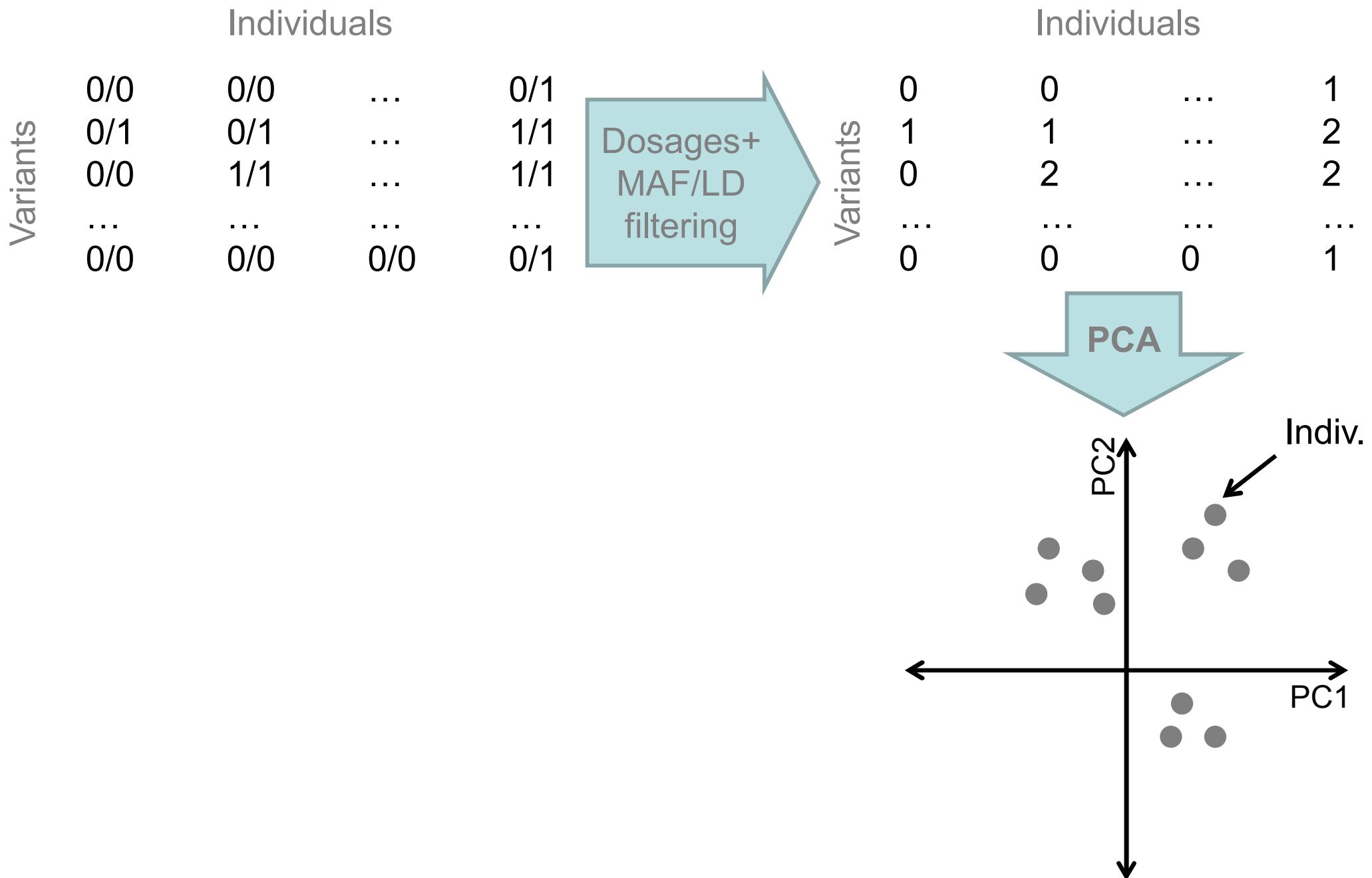
Population structure

- PCA calculate axes of genetic variation that maximise the variability between individuals.
 - Plotting axes of genetic variation from PCA including 1000 genomes samples can be used to identify population outliers.
 - Axes of genetic variation can be used as covariates in downstream analysis to adjust for the effects of population structure.
- Alternative methods exist such as MDS for instance as implemented in PLINK.

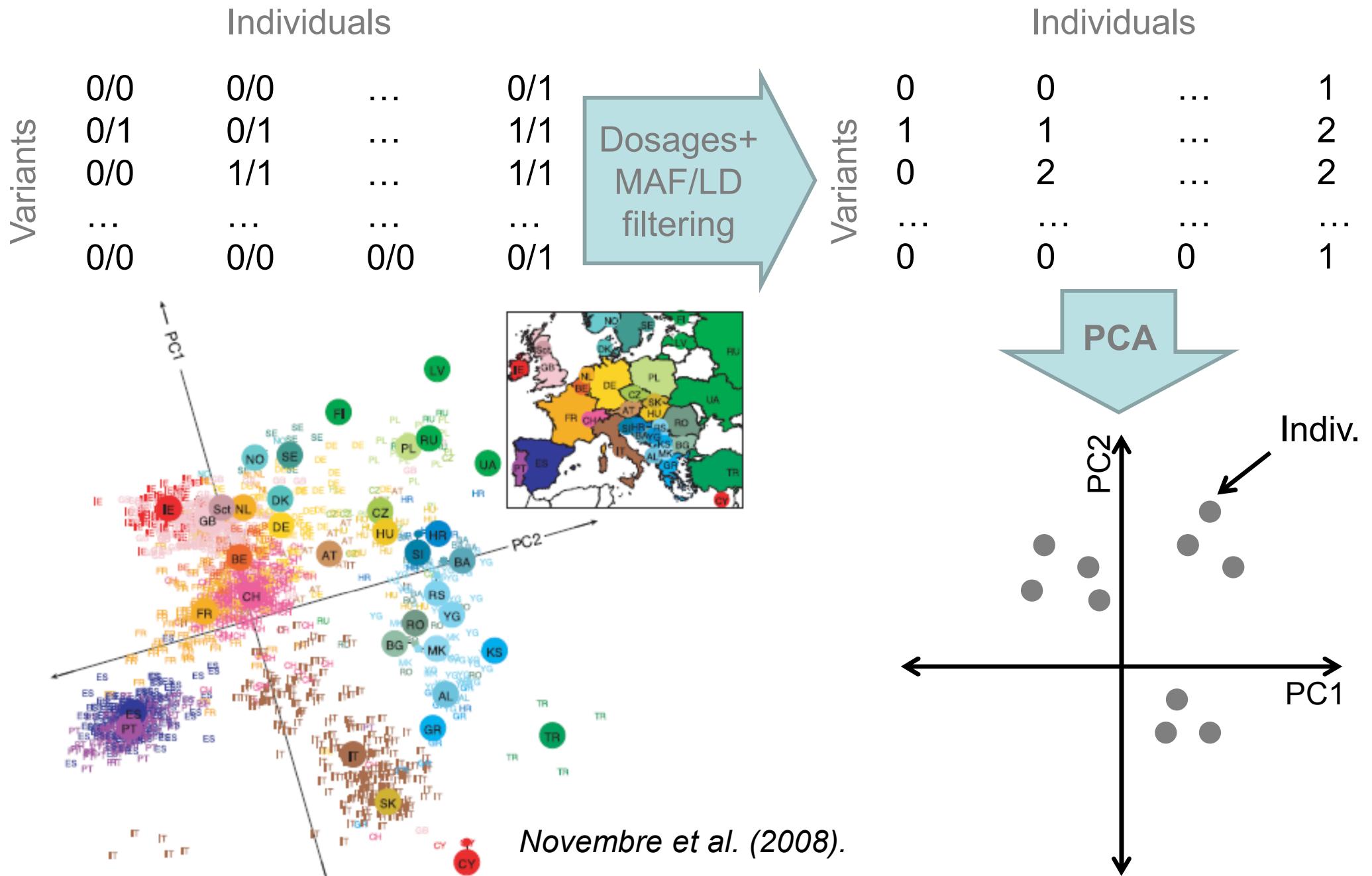
Population stratification



Population stratification

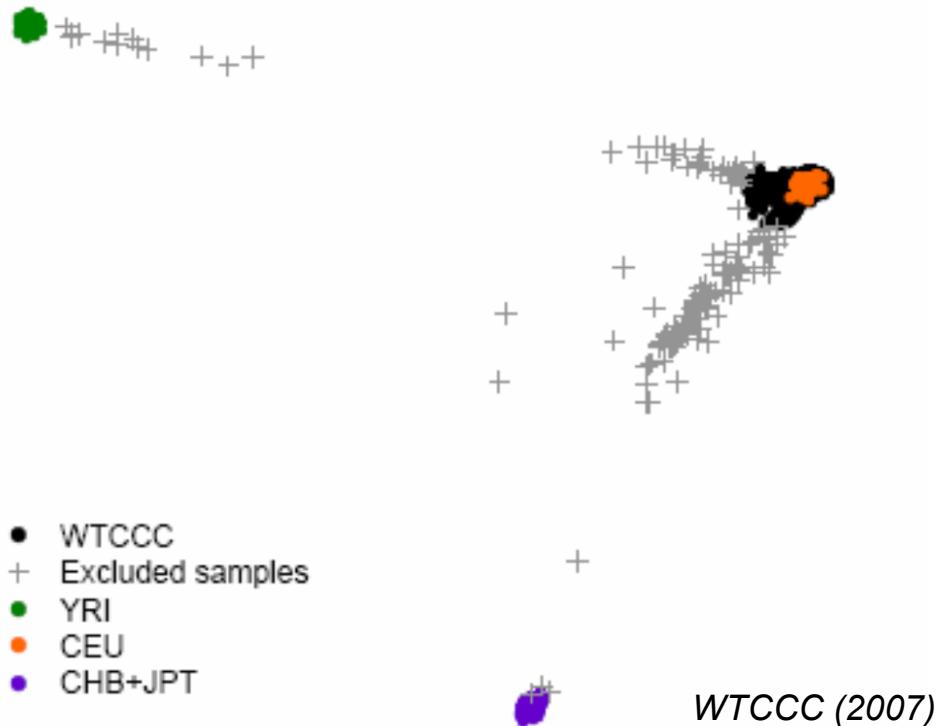


Population stratification



What do I do with my PCs?

1. Remove outliers from analysis



2. Account for structure

- By defining ethnic groups
- By using the PC loadings as covariates

Summary

- QC criteria are subjective and vary from one study to another
- Variant QC filters should eliminate the worst quality markers without “throwing the baby out with the bathwater”
- Sample QC filters should not be so stringent as to remove the majority of the analysis cohort
- Any finding should be followed up with visual inspection of the raw data (i.e. sequencing reads or cluster plots)

Practical

- The goal of the practical of this morning is to go from raw to ready-to-analyze data
- We will use multiple standard QC steps both at the variant and sample levels
- The resulting data set will be used as starting material for the afternoon practical
- Look at **/embo/data/olivier**, everything you need is there.