

Genome-wide association studies I: Identifying genetic associations with complex traits

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MSc Global Health Science and Epidemiology

Genetic Epidemiology Module

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Learning objectives

Understand a genome-wide association study (GWAS) and the concept of a hypothesis-free approach to studying genetic associations.

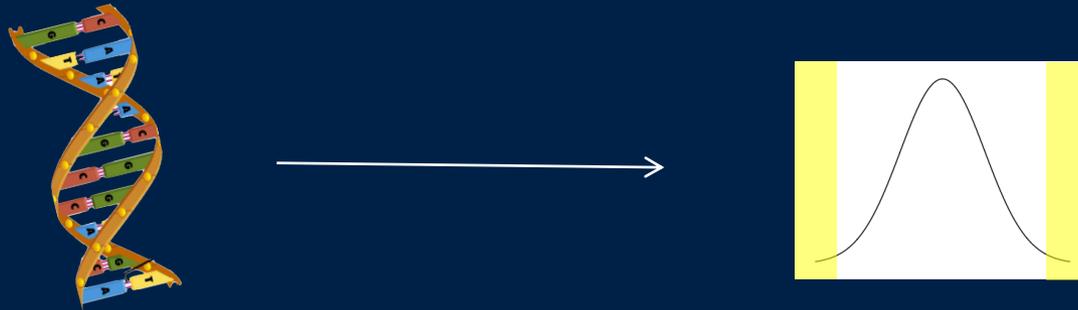
Have a working knowledge of the different steps involved in the conduct of GWAS, including study design, quality control and basic analyses.

Be able to interpret and critically appraise evidence from genome-wide association studies.

Understand the relevance of replication, meta-analysis and consortia, and multi-ancestry approaches, in genome-wide association studies.

Appreciate the use of post-GWAS analyses including fine mapping, gene and pathway analyses, and the concept of causal variants.

Learning objectives



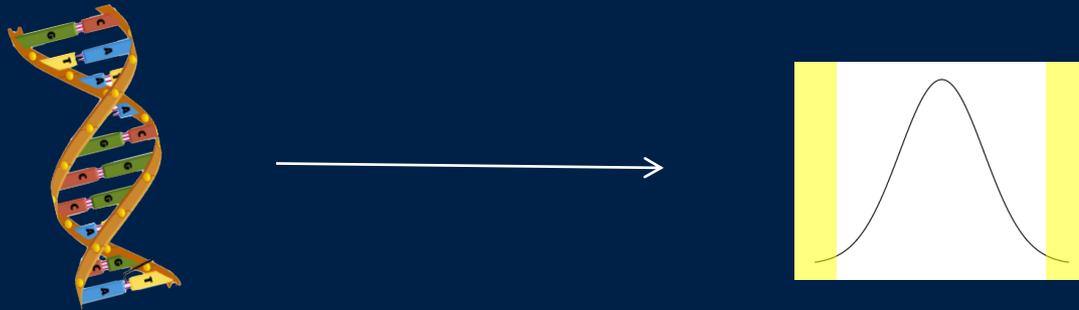
Describe studies aiming to find genetic differences between individuals that influence susceptibility to complex diseases (or other traits).

Lecture outline

- • Background
- Searching for a needle in a haystack
- Genome-wide association studies in theory
- GWAS in practice
- The challenge of understanding biology

The human genome is ~3.2 billion base pairs long.

About 1 in 100 – 1000 of those bases vary between people.



What proportion of phenotypic variation is due to genetic variation?

Human traits are highly heritable

Idea: if genetics determines a trait, then *more genetically similar individuals should have more similar phenotypes*. Can estimate how much genetics determines trait variation by comparing trait similarity in monozygotic (identical) and dizygotic twins.

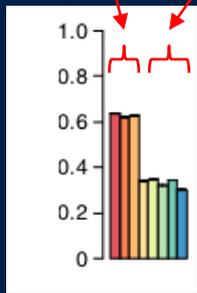
Meta-analysis of the heritability of human traits based on fifty years of twin studies

Tinca J C Polderman^{1,10}, Beben Benyamin^{2,10}, Christiaan A de Leeuw^{1,3}, Patrick F Sullivan⁴⁻⁶, Arjen van Bochoven⁷, Peter M Visscher^{2,8,11} & Danielle Posthuma^{1,9,11}

(2015)

MZ
Twins
 $r \sim 0.64$

DZ
Twins
 $r \sim 0.34$



All studied
traits

Across all traits, phenotypes are much more highly correlated between monozygotics than dizygotic twins. Heritability (averaged across traits) is about 60%.

Definition: Heritability is the proportion of trait variation explained by inherited factors (including genetics). Can be estimated as $h^2 \approx 2 \times (r_{MZ} - r_{DZ})$

Human traits are highly heritable

If genetics determines a trait, then *more genetically similar individuals should have more similar phenotypes.*

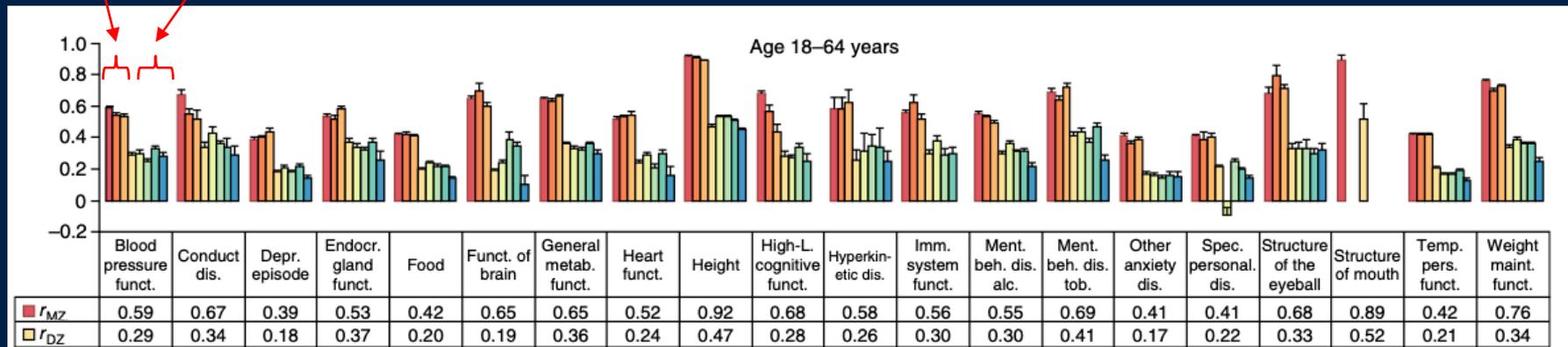
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(2015)

Monozygotic

Dizygotic



Blood pressure
 $h^2 \approx 60\%$

Depression
 $h^2 \approx 42\%$

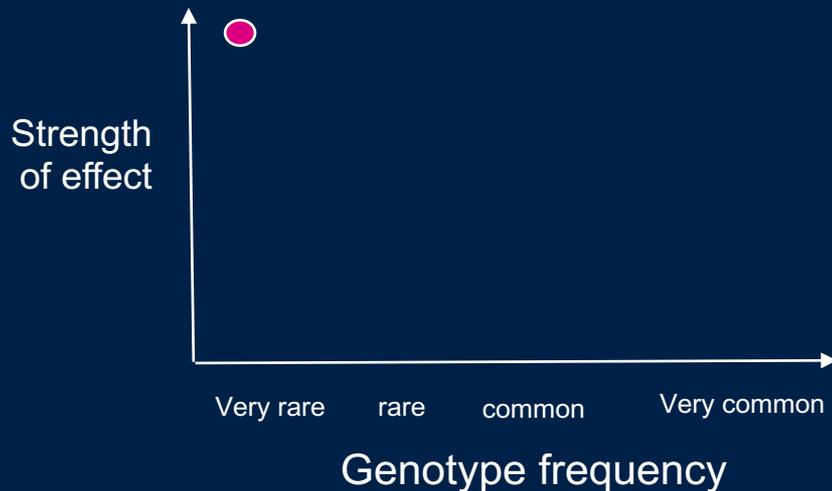
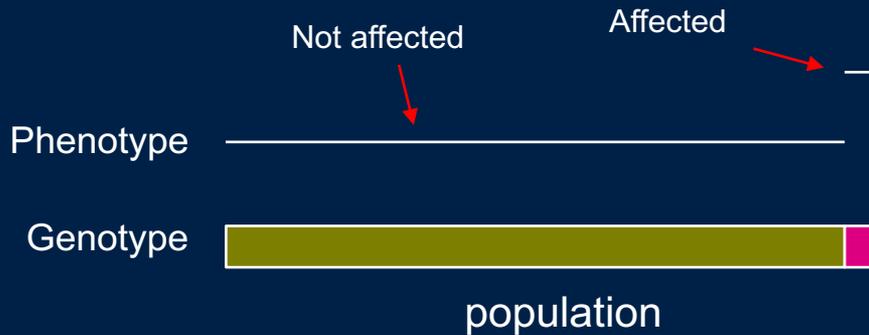
Adult height
 $h^2 \approx 90\%$

“Higher level cognitive function”
 $h^2 \approx 80\%$

Structure of the eyeball
 $h^2 \approx 70\%$

Lots of theoretical caveats might apply here – see Lecture 1. These estimates are also a little higher than some from other methods. But in general it is true that *a large proportion of variation in most human phenotypes is caused by genetics.*

Two possible extreme genetic architectures



Example: Huntington's

Cell, Vol. 72, 971-983, March 26, 1993, Copyright © 1993 by Cell Press

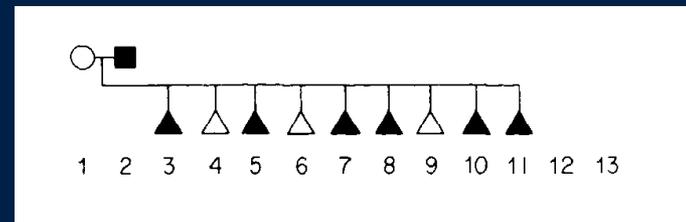
A Novel Gene Containing a Trinucleotide Repeat That Is Expanded and Unstable on Huntington's Disease Chromosomes

The Huntington's Disease Collaborative Research Group*

Introduction

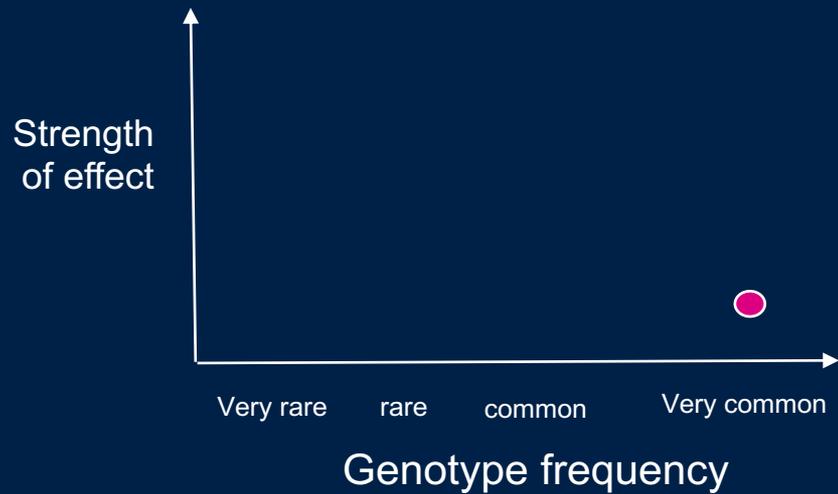
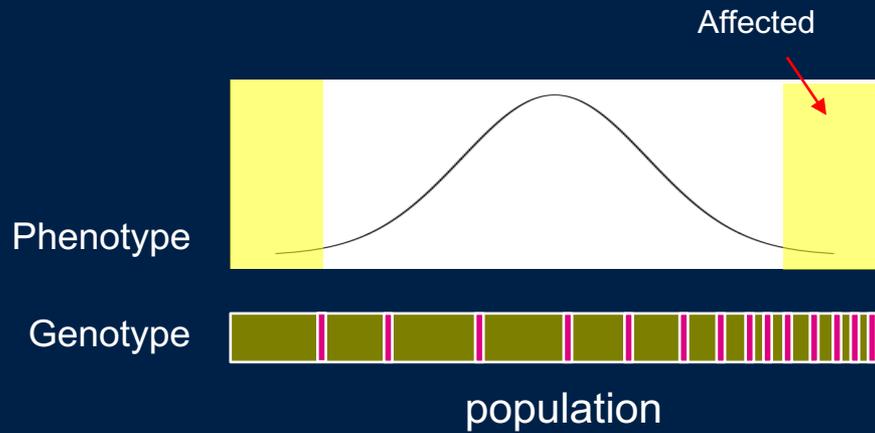
Affects ~1 in 20,000 people of European ancestry (less in Africa and Asia)

Discovered by looking in families

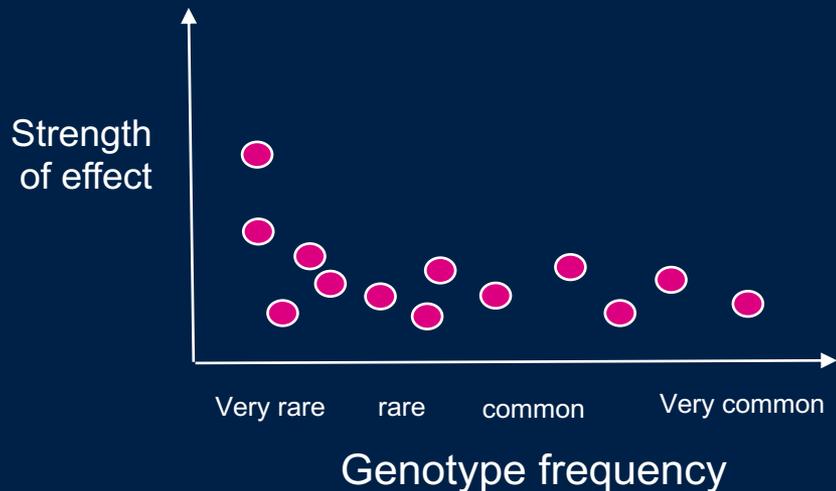
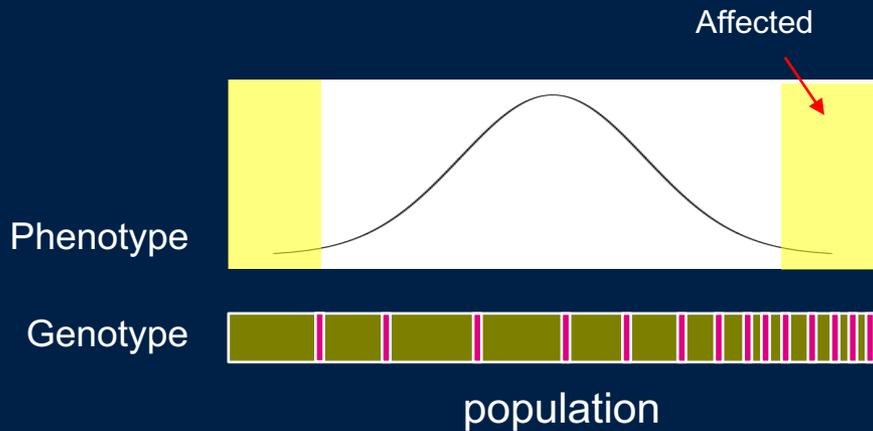


A "Mendelian" trait

Two possible extreme genetic architectures



Two possible extreme genetic architectures



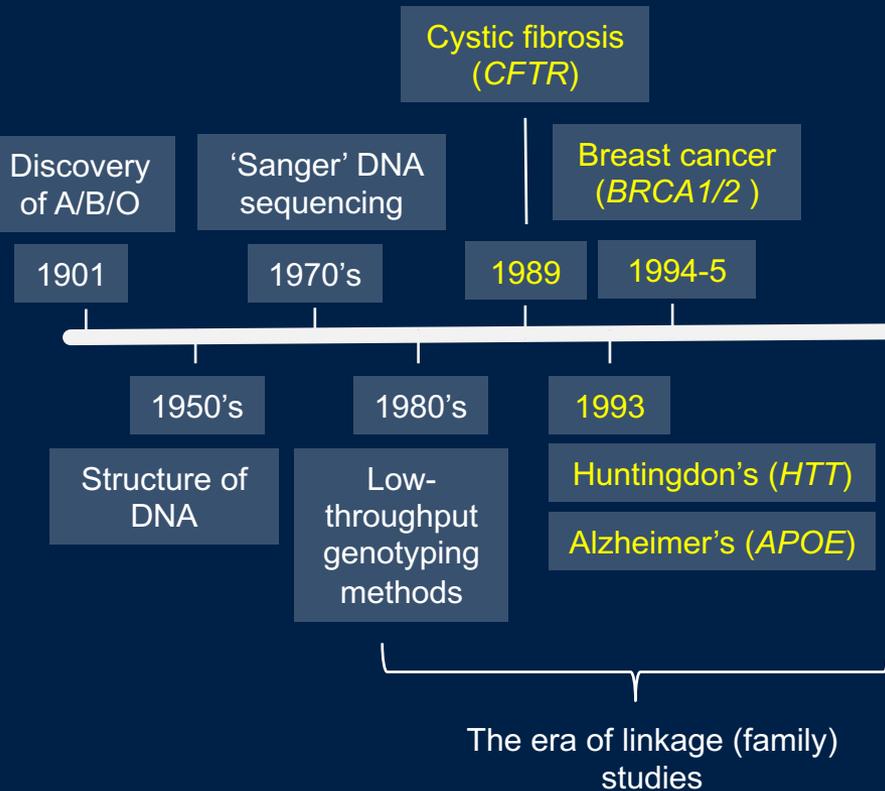
A complex trait.

Caused by many factors, each having a small overall effect. including

- Many genetic variants, including common ones
- Environmental factors
- Gene-environment or gene-gene interactions
- ...

End of an era

“Linkage Mapping was successful in identifying the genetic basis of many human diseases in which the disease penetrance resembles a simple Mendelian model e.g. Huntington’s disease, Cystic Fibrosis, some forms of breast cancer, Alzheimers, ...”



“...but the literature is now replete with linkage screens for an array of common ‘complex’ disorders such as schizophrenia, manic depression, autism, asthma, type I and type II diabetes, Multiple Sclerosis, Lupus. Although many of these studies have reported significant linkage findings, none has lead to convincing replication”

– Risch “*Searching for genetic determinants in the new millennium*” Nature (2000)

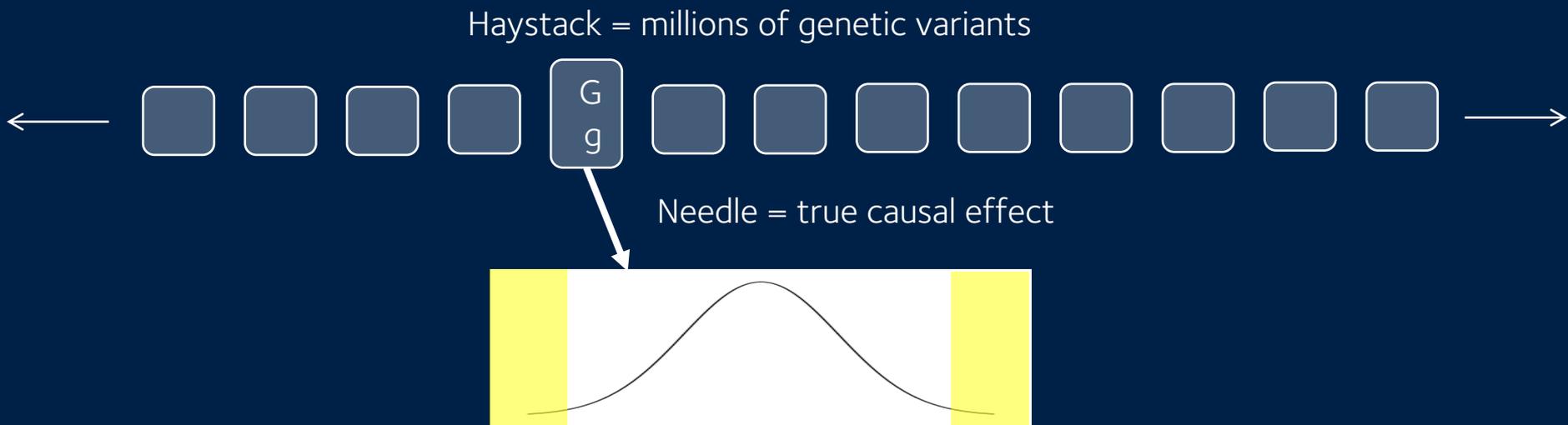
Summary

- With few exceptions, human phenotypes are highly heritable – a large proportion of phenotype variation seems to be caused by genetics.
- In principle this heritability could occur in different ways – for example through single variants with strong effects, or through multiple variants with small effects.
- By the 2,000s family studies had identified the causes of several mendelian traits, but had failed to solve the genetics of multiple complex diseases.

Lecture outline

- Background
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- GWAS in practice
- The challenge of understanding biology

Causal effects generate relative risk $\neq 1$



If genotype G causes disease, then carrying G will make you more likely to have disease. i.e.:

$$\text{Relative risk} = \frac{P(\text{disease}|\text{genotype } G)}{P(\text{disease}|\text{genotype } g)} > 1$$

“Probability (or frequency) of disease given genotype G ”

(ditto for genotype g)

A causal effect will generate a relative risk $\neq 1$.

How to estimate relative risk?

$$RR = \frac{P(\text{disease} | G)}{P(\text{disease} | g)}$$

Disease frequencies

(in population)

How to estimate relative risk?

$$RR = \frac{P(\text{disease} | G)}{P(\text{disease} | g)} = \frac{P(G | \text{disease})}{P(g | \text{disease})} \times \frac{P(g)}{P(G)} \quad (\text{in population})$$

Disease frequencies

Genotype frequencies
in cases and controls

How to estimate relative risk?

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Disease frequencies

Genotype frequencies
in cases and controls

Suppose we sample some disease cases and population* controls:

	G	g
Disease cases:	a	b
Controls:	c	d

$$OR = \frac{a}{b} \times \frac{d}{c}$$

(in sample)

⇒ The sample *odds ratio* estimates the population relative risk.

* If the controls are 'true' (unaffected) controls, this is still approximately true provided the disease is rare.

How to estimate relative risk?

$$RR = \frac{P(\text{disease} | G)}{P(\text{disease} | g)} = \frac{P(G | \text{disease})}{P(g | \text{disease})} \times \frac{P(g)}{P(G)} \quad (\text{in population})$$

Disease frequencies

Genotype frequencies
in cases and controls

Suppose we sample some disease cases and population* controls:

	<i>G</i>	<i>g</i>
Disease cases:	a	b
Controls:	c	d

$$OR = \frac{a}{b} \times \frac{d}{c}$$

(in sample)

⇒ The sample *odds ratio* estimates the population relative risk. How accurately?

$$\text{Standard error}(\log OR) \approx \frac{1}{\sqrt{N \times f(1-f) \times \phi(1-\phi)}}$$

Sample size

Genotype frequency

Ratio of cases to
controls in study

Example: O blood group is associated with malaria protection

	O	non-O
Severe malaria cases	686	843
Controls:	839	700

$$OR = \frac{686}{843} \times \frac{700}{839} = 0.68$$

Data from $N=3,068$ samples from Kilifi, Kenya
MalariaGEN 2019 doi: 10.1038/s41467-019-13480-z

O blood group is associated with a
~30% lower chance of severe malaria
(all else being equal).

Example: O blood group is associated with malaria protection

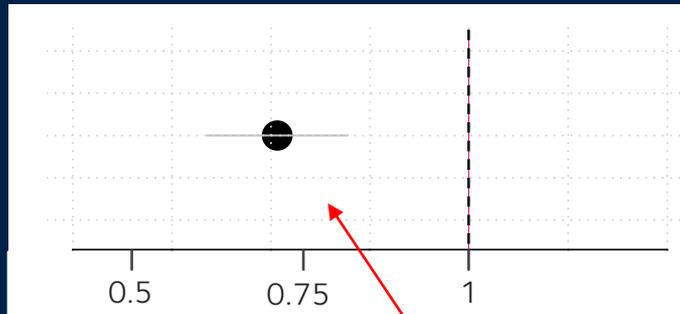
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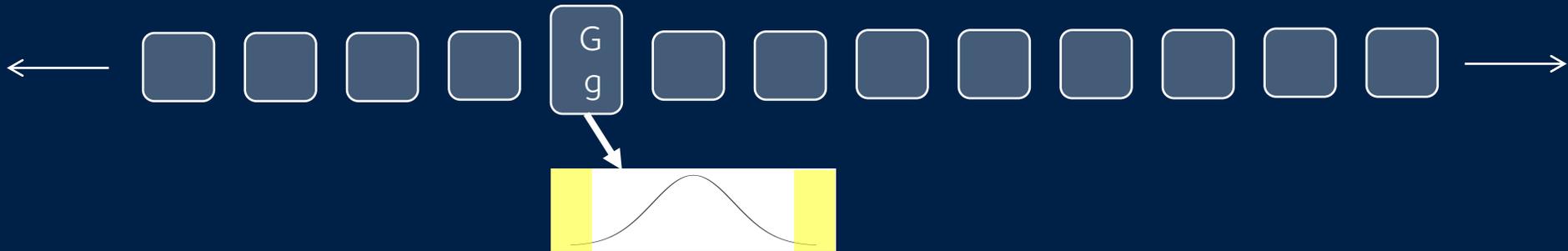
$$\text{Standard error}(\log OR) \approx \frac{1}{\sqrt{3068 \times 0.45 \times 0.55 \times 0.25}} \approx 0.073 \quad (\text{on log scale})$$



Estimated relative risk = 0.68
95% CI = 0.59-0.78
(estimate +/- 1.96 standard errors)

Estimate is > 5 standard errors from zero
 $P = 9.6 \times 10^{-8}$

Summary



- If a genotype G causes disease, then it will have $RR \neq 1$
- Relative risk can be estimated from a case-control sample.

$$se_{\text{(of log OR)}} \approx \frac{1}{\sqrt{N \times f(1-f) \times \phi(1-\phi)}}$$

Larger samples, and higher-frequency variants, lead to more accurate estimates.

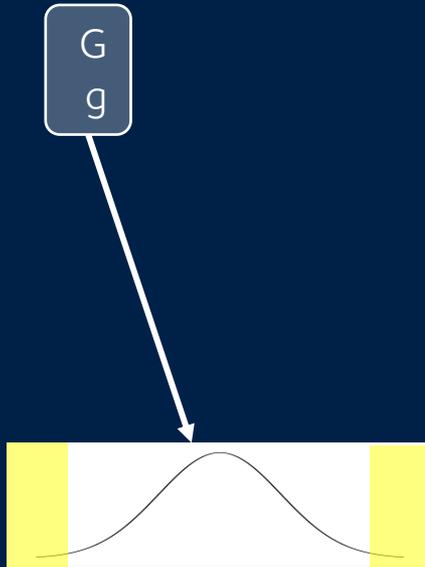
- If the estimate is sufficiently far from one (how far?) this suggests we have found a true causal effect.
- But must take care of confounders!

Anatomy of a GWAS, version 1

1. Collect as many cases and controls as possible
2. Genotype them at as many variants across the genome as possible
3. Estimate relative risks, and look for statistical evidence that of $RR \neq 1$
4. If estimate is many standard deviations from zero, bingo! We may have found a true causal effect.
5. (Now try to understand the underlying biology.)

Three potential confounders

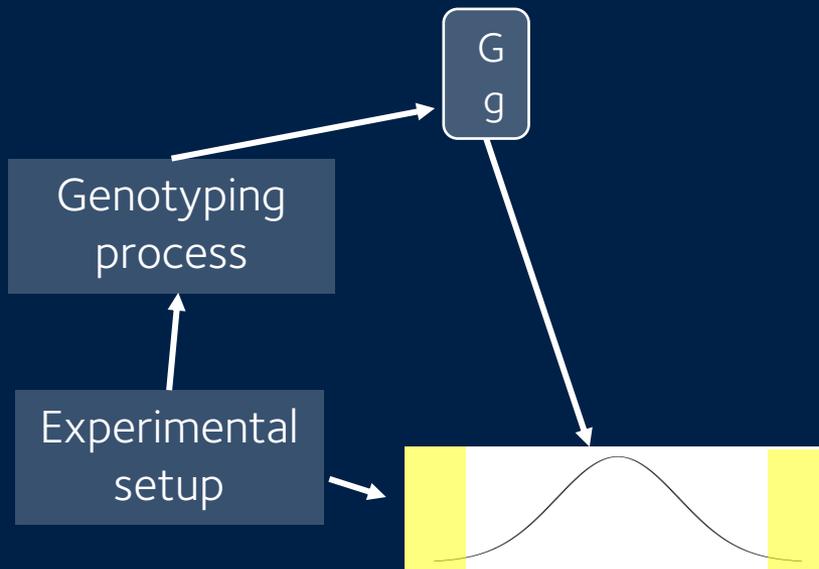
Association tests capture all causal paths from genotype to phenotype – even those that have nothing to do with biology.



Three potential confounders

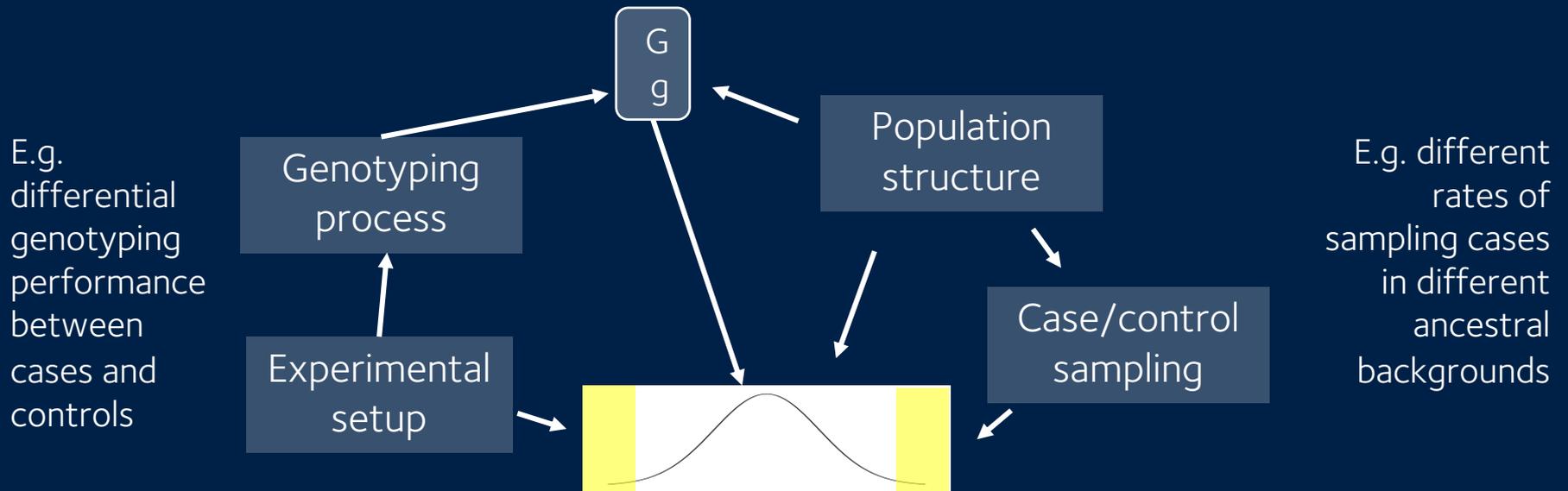
Association tests capture all causal paths from genotype to phenotype – even those that have nothing to do with biology.

E.g.
differential
genotyping
performance
between
cases and
controls



Three potential confounders

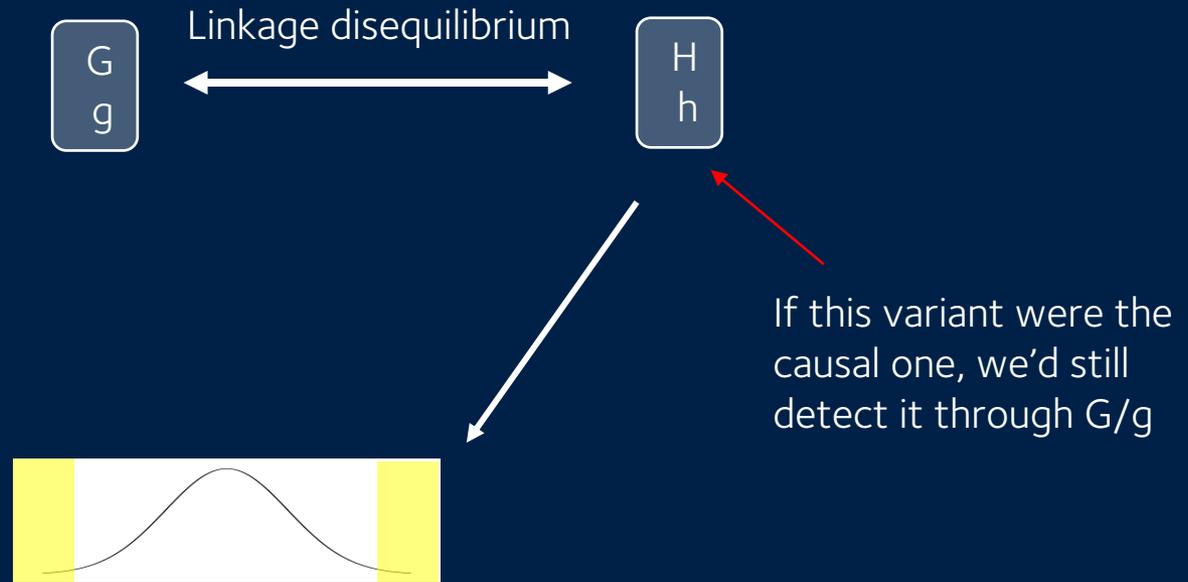
Association tests capture all causal paths from genotype to phenotype – even those that have nothing to do with biology.



Three potential confounders

Association tests capture all causal paths from genotype to phenotype – even those that have nothing to do with biology.

Will also pick up effects from all nearby causal variants that are in LD



If this variant were the causal one, we'd still detect it through G/g

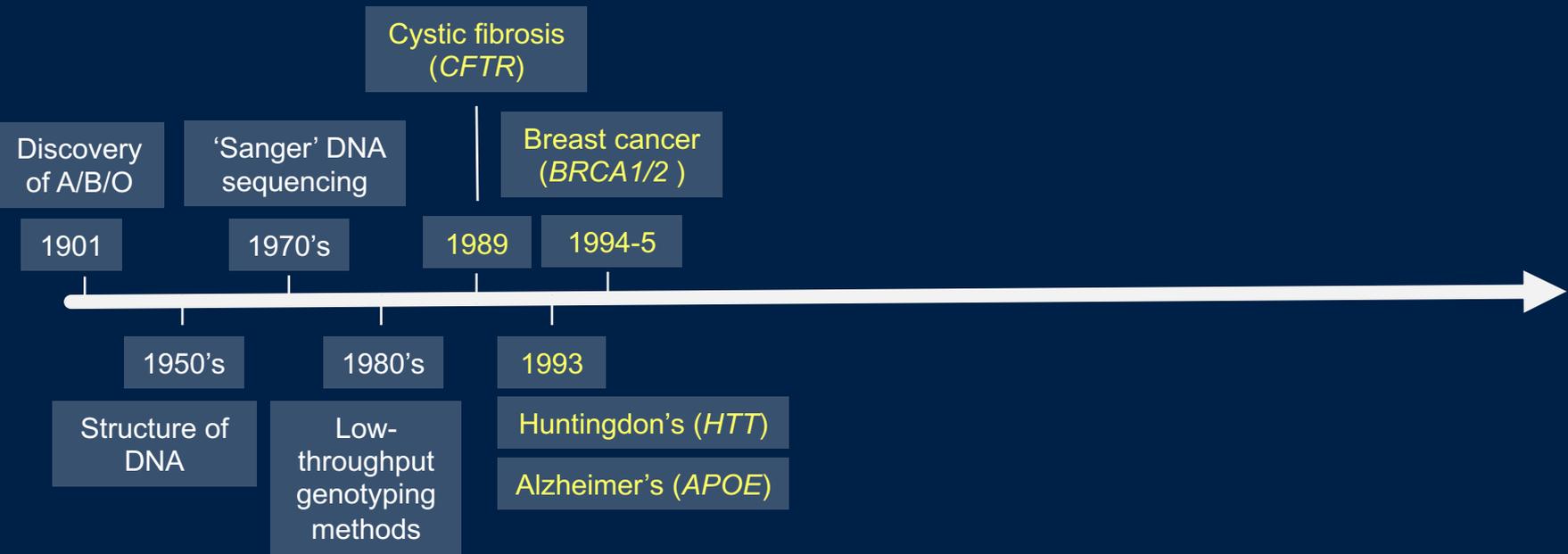
Anatomy of a GWAS, version 2

1. Collect as many cases and controls as possible
2. Genotype or impute them at as many variants across the genome as possible. Rely on LD to capture the others.
3. Deal with potential confounders – careful data quality control and handle population structure. 
4. Estimate relative risks, and look for statistical evidence that of $RR \neq 1$
5. If estimate is many standard deviations from zero, bingo! We may have found a true causal effect.
6. Does it replicate in other studies, or have other corroborating evidence? 
7. (Now try to understand the underlying biology.)

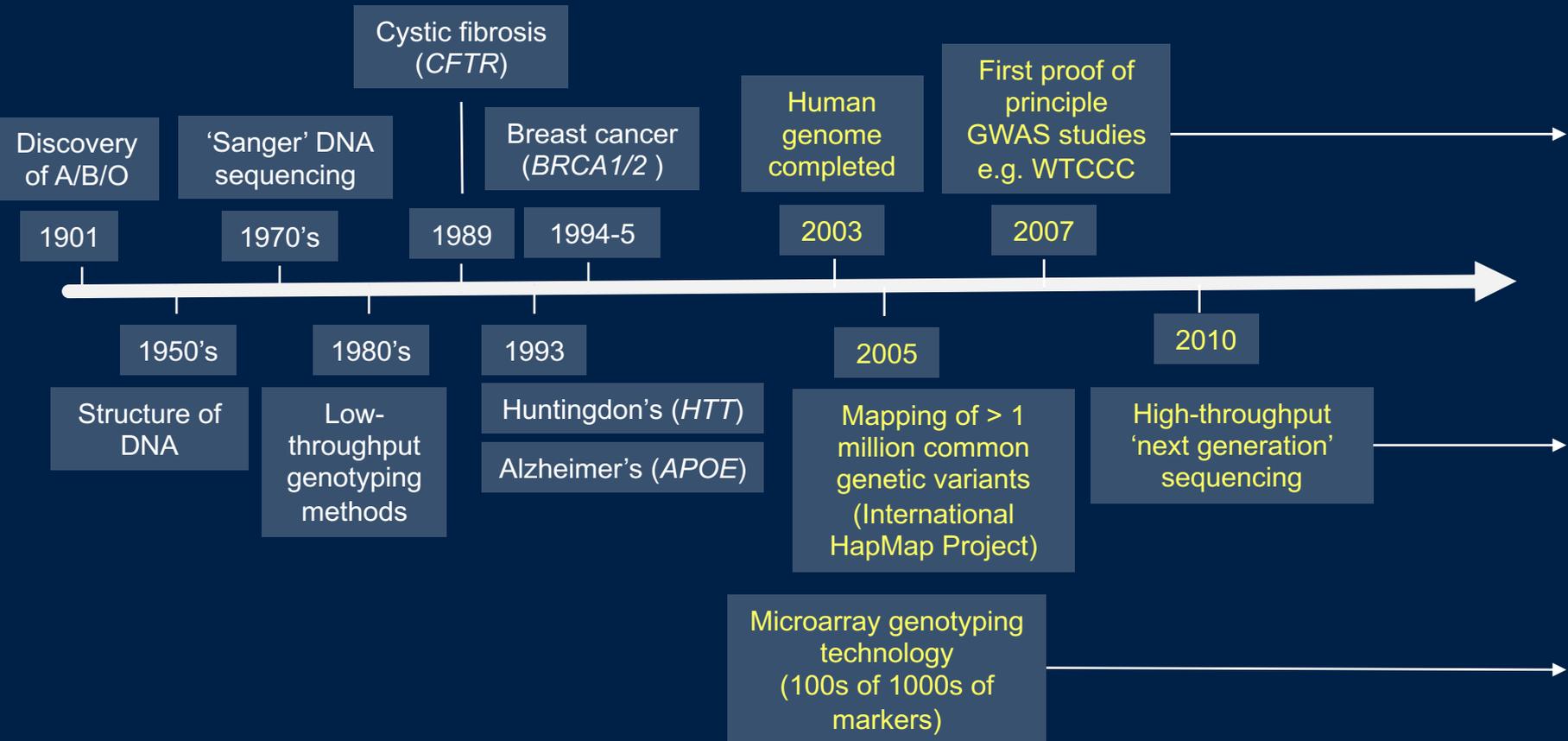
Lecture outline

- Background
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- GWAS in practice
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End of the linkage era



The birth of GWAS



Understanding human genetic diversity

A haplotype map of the human genome

The International HapMap Consortium*

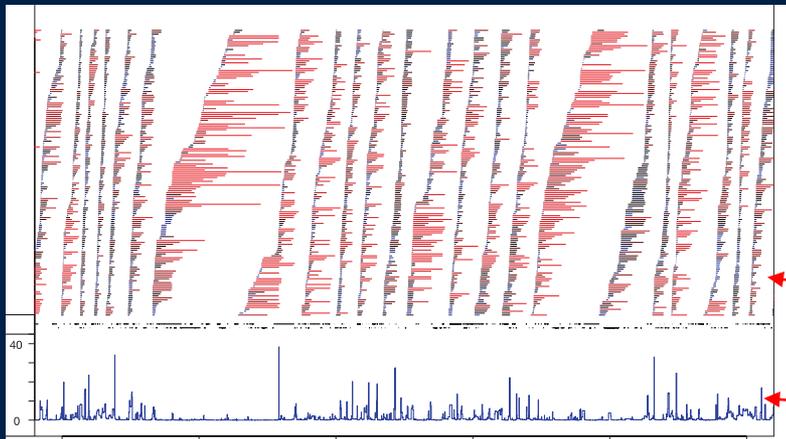
Inherited genetic variation has a critical but as yet largely uncharacterized role in human disease. Here we report a public database of common variation in the human genome: more than one million single nucleotide polymorphisms (SNPs) for which accurate and complete genotypes have been obtained in 269 DNA samples from four populations, including ten 500-kilobase regions in which essentially all information about common DNA variation has been extracted. These data document the generality of recombination hotspots, a block-like structure of linkage disequilibrium and low haplotype diversity, leading to substantial correlations of SNPs with many of their neighbours. We show how the HapMap resource can guide the design and analysis of genetic association studies, shed light on structural variation and recombination, and identify loci that may have been subject to natural selection during human evolution.

International HapMap Project

doi:10.1038/nature0422 (2005)

A database of > 1 M SNPs found in European, African, and Asian ancestry individuals

(A subset of the samples later used in the 1000 Genomes Project)



Recombination turns out to be highly nonuniform. It is concentrated in *recombination hotspots*. So mutations are carried on longer haplotypes than had been expected.

Shared haplotype lengths

Map of recombination rate

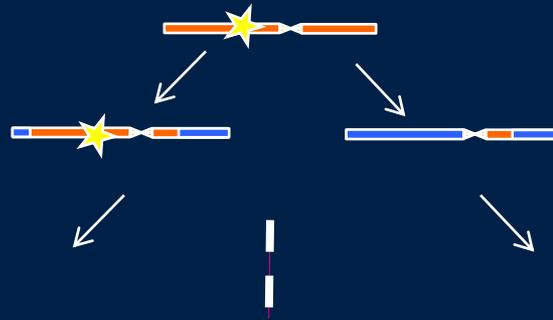
Tag SNP set size	Common SNPs captured (%)		
	YRI	CEU	CHB + JPT
10,000	12.3	20.4	21.9
20,000	19.1	30.9	33.2
50,000	32.7	50.4	53.6
100,000	47.2	68.5	72.2
250,000	70.1	94.1	98.5

As in Table 7, tag SNPs were picked to capture common SNPs in HapMap release 16c1 using Haploview, selecting SNPs in order of the fraction of sites captured. Common SNPs were captured by fixed-size sets of pairwise tags at $r^2 \geq 0.8$.

Estimated how many SNPs genome-wide would need to be typed to capture (by LD) most common genetic variants. E.g. 250,000 would capture ~95% of SNPs in European populations.

What does human genetic variation look like anyway?

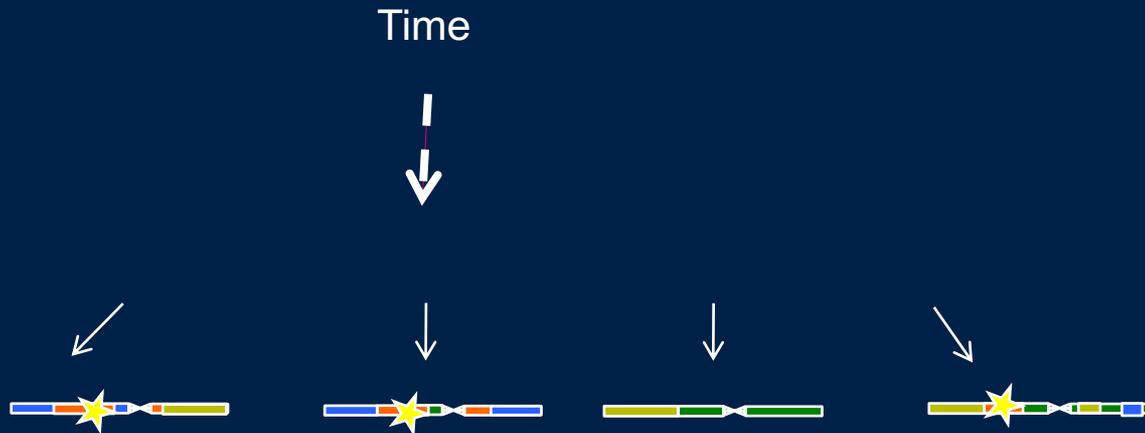
Mutation arises



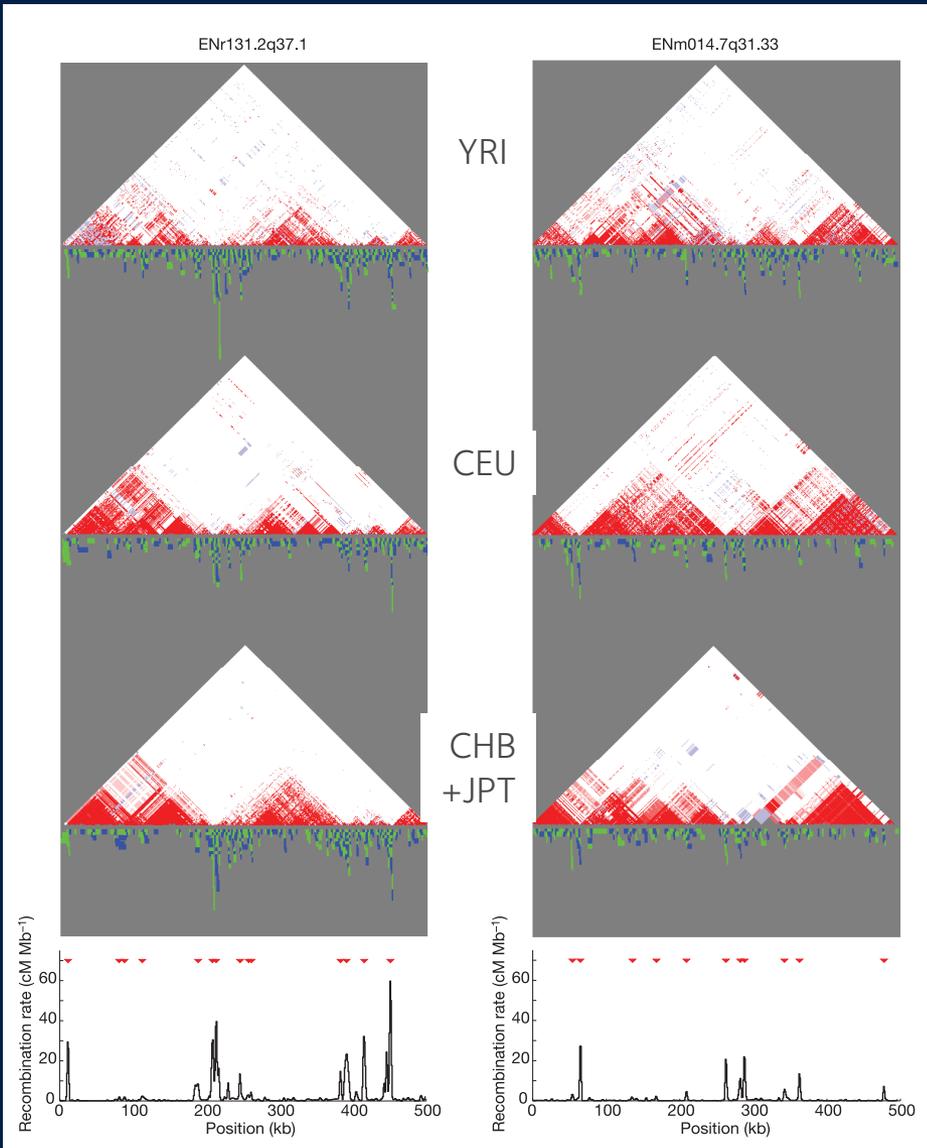
Gets passed on through many generations

Changes in frequency cause variants to become correlated (LD)

Recombination breaks this down leading to local patterns

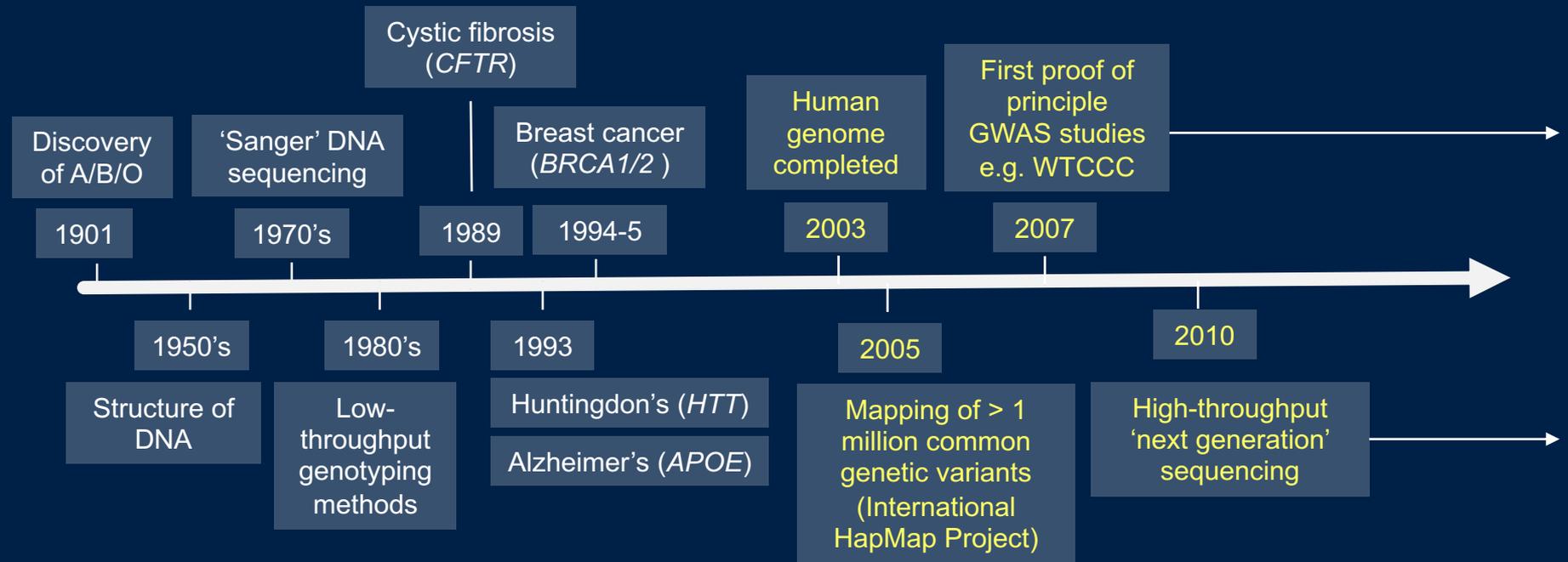


Patterns of LD depend on overall population size.
There are higher levels of LD in smaller populations.

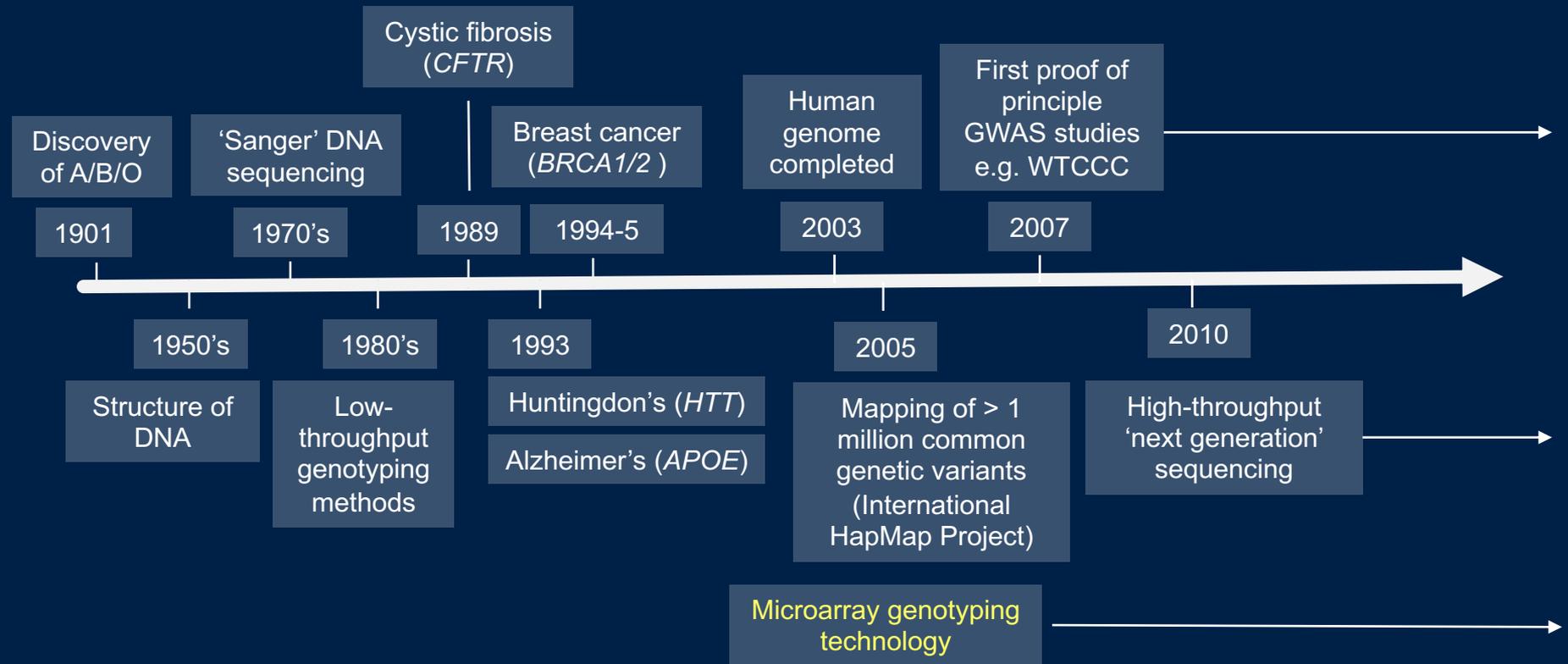


Block-like structure of LD
(correlations between SNPs
in two different regions)

The birth of GWAS



The birth of GWAS



Microarrays developed in the late 90's / early 2000's.
For the first time was possible to rapidly type hundreds of thousands or millions of SNPs

Lecture outline

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A real GWAS study – WTCCC

Genome-wide association study of 14,000 cases of seven common diseases and 3,000 shared controls

The Wellcome Trust Case Control Consortium*

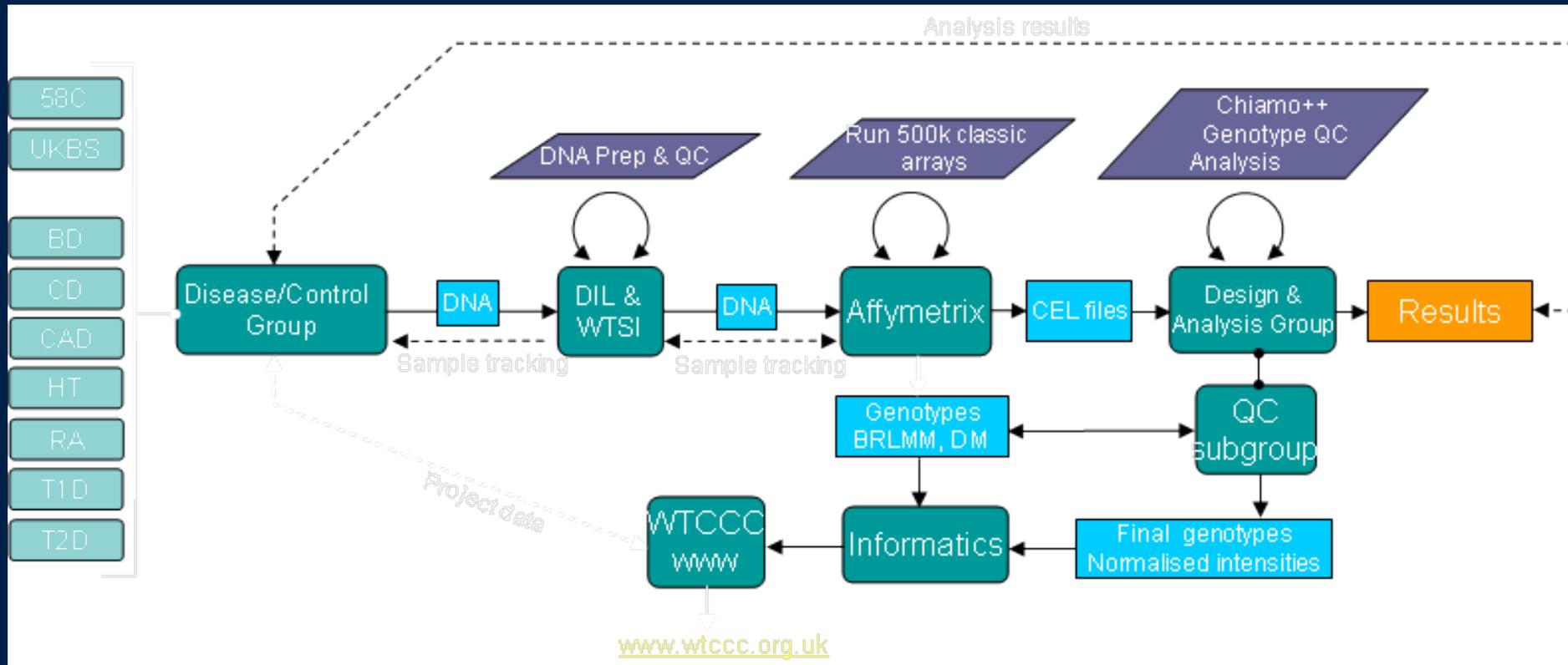
Nature (2007)

Studied seven common diseases in the UK

Bipolar disorder, Coronary Artery Disease, Crohn's disease, Hypertension, Rheumatoid arthritis, Type 1 and Type 2 Diabetes

Genotyped at 500,000 SNPs across the genome

A real study - WTCCC



Anatomy of a GWAS – what to look for

1. Collect as many cases and controls as possible

N=2,000 cases and
3,000 controls

2. Genotype (or impute) them at as many variants across the genome as possible

Genotyped at 500k
SNPs

3. Deal with potential confounders – careful data quality control and handle population structure.

Have they done adequate
data quality control?
Have they dealt with
possible confounders?

4. Estimate relative risks, and look for statistical evidence that of $RR \neq 1$

5. If estimate is many standard deviations from zero, bingo! We may have found a true causal effect.

Did they find anything
with strong evidence?

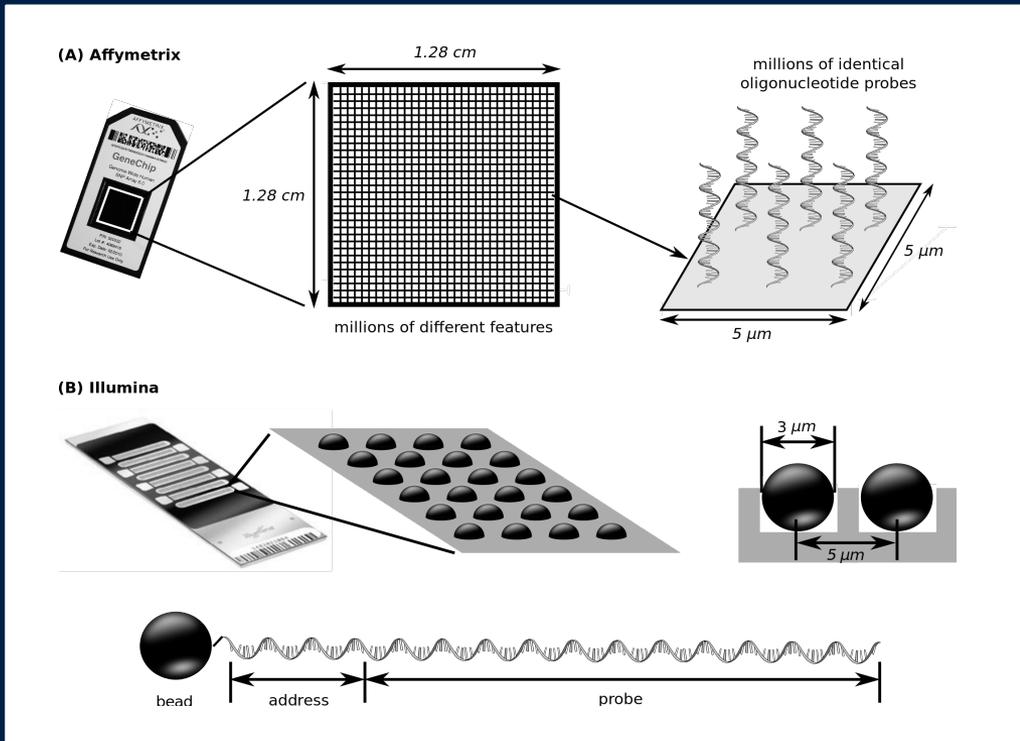
6. Does it replicate in other studies, or have other corroborating evidence?

Is it convincing?

7. (Now try to understand the underlying biology.)

What about biology?

How a microarray works

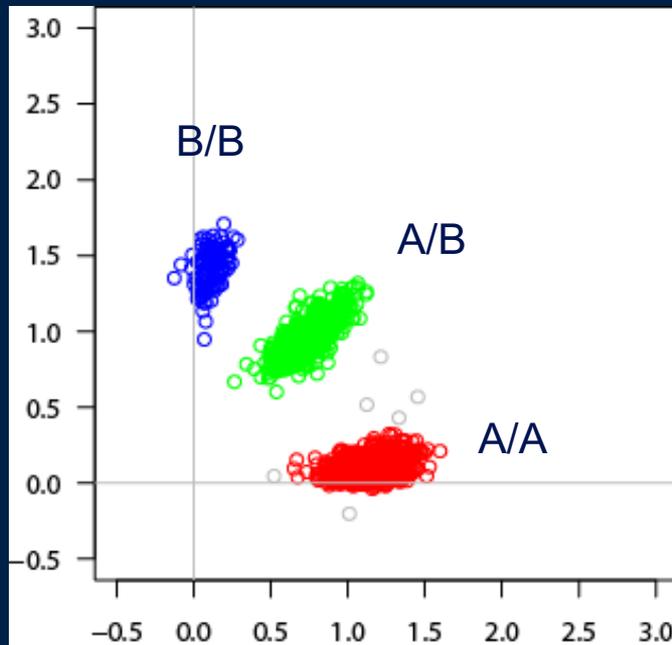


Wash the DNA over and let it hybridise to millions of probes – one for each SNP

Flourescent markers are then attached. A picture is taken of the array.

A microarray gives you intensities, not genotypes

For each SNP, you get back this:

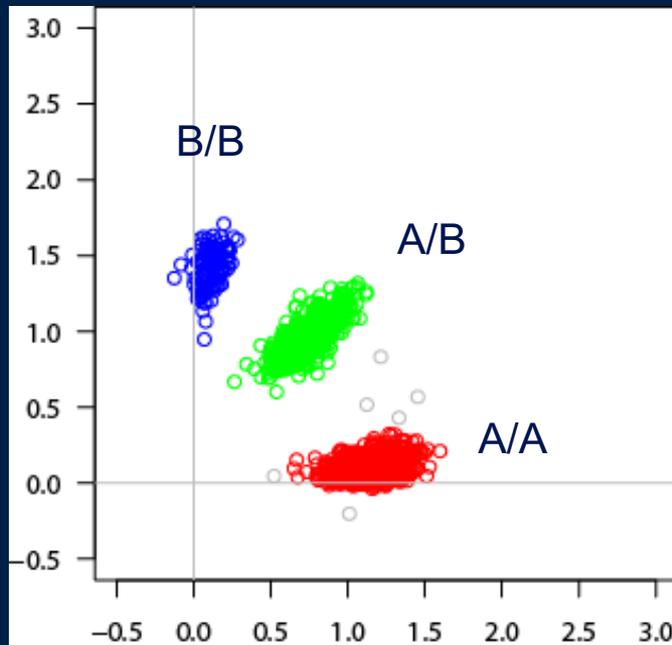


An algorithm is needed to turn the intensity values (x/y axis values) into genotype calls (colours).

Each dot represents DNA from one individual.
X axis = image intensity for 1st SNP allele
Y axis = image intensity for 2nd SNP allele

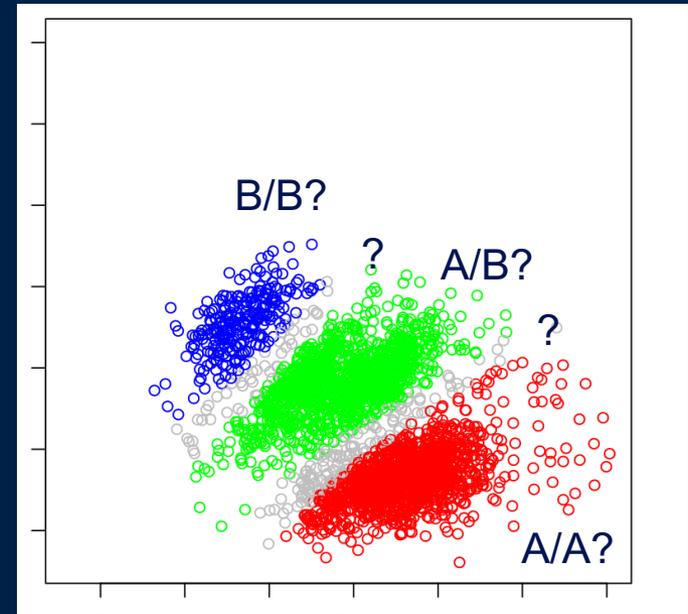
A microarray gives you intensities, not genotypes

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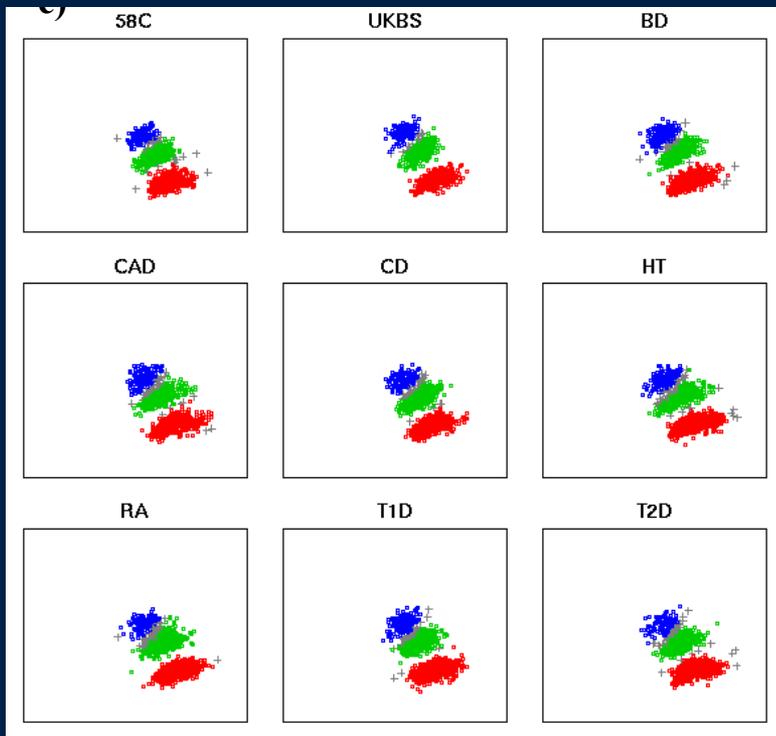
Each dot represents DNA from one individual.
X axis = image intensity for 1st allele probe
Y axis = image intensity for 2nd allele probe

Or this if you're less lucky:



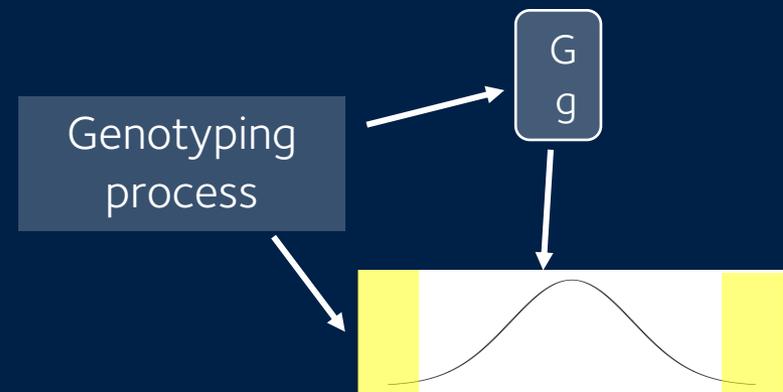
Small genotyping errors in cases or controls could easily confound the study

An algorithm is needed to call genotypes



The authors developed a genotype calling algorithm to turn these data (intensities, X and Y axis) into genotype calls (colours). Samples lying outside clusters, or in overlapping clusters, would be called as missing. (NB. Nowadays most studies use off-the-shelf algorithms for this.)

In particular cases and controls were jointly called.



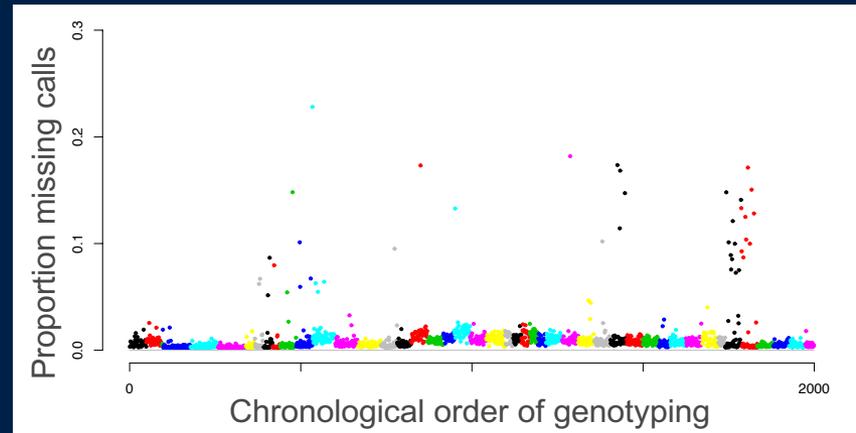
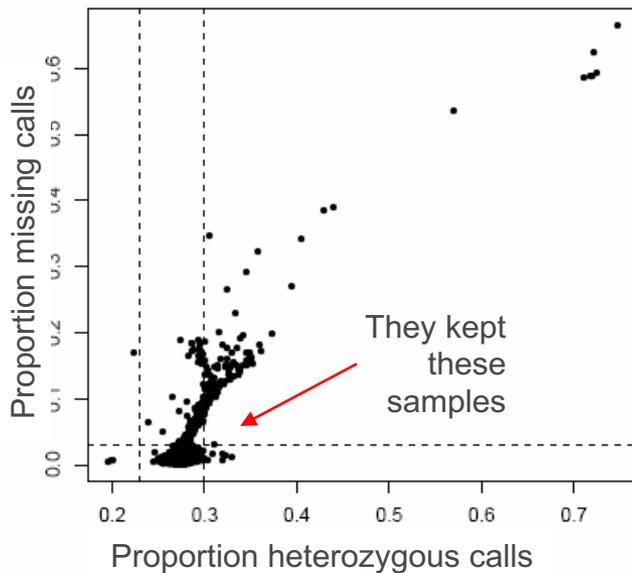
Collection	Missingness	Heterozygosity	External discordance	Non-European ancestry	Duplicate	Relative	Total
58C	9	0	4	6	4	1	24
UKBS	8	0	5	14	0	15	42
BD	30	0	0	9	77	13	129
CAD	41	1	0	13	2	5	62
CD	43	4	6	54	131	18	256
HT	29	0	0	2	6	11	48
RA	47	1	0	26	53	9	136
T1D	7	2	1	18	6	3	37
T2D	36	1	0	11	16	11	75
Total	250	9	16	153	295	86	809

Supplementary Table 4 | Exclusion summary by collection. Six filters were applied for sample exclusion: 1. SNP call rate < 97% (missingness). 2. Heterozygosity > 30% or < 23% across all SNPs. 3. External discordance with genotype or phenotype data. 4. Individuals identified as having recent non-European ancestry by the Multidimensional Scaling analysis (see Methods). 5. Duplicates (the copy with more missing data was removed) 6. Individuals with too much IBS sharing (>86%); likely relatives. Where individuals could be excluded for more than one reason, they appear in the leftmost such column.

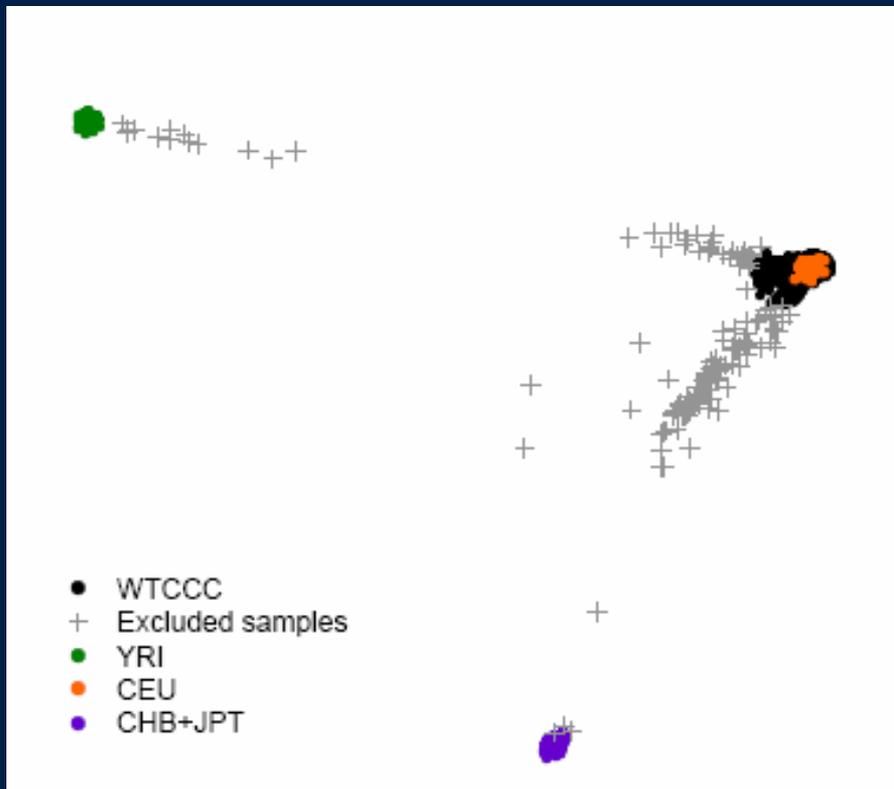
They then threw away 809 samples!

Due to:

- Poor genotyping rates
- Evidence of contamination (too many heterozygous genotypes)
- Evidence of being not of European ancestry
- A duplicate, or close relative of another sample



Some of the poor quality data was apparently due to batch effects.

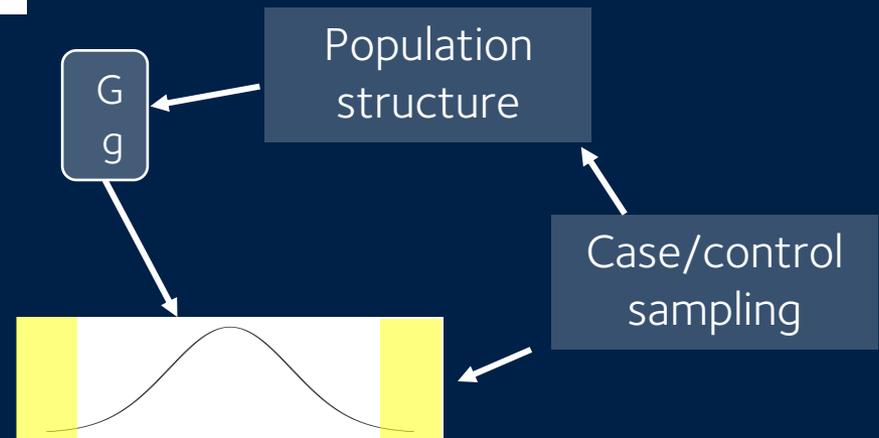


To avoid confounding by population structure, the samples were all supposed to be from the United Kingdom, and with European ancestry.

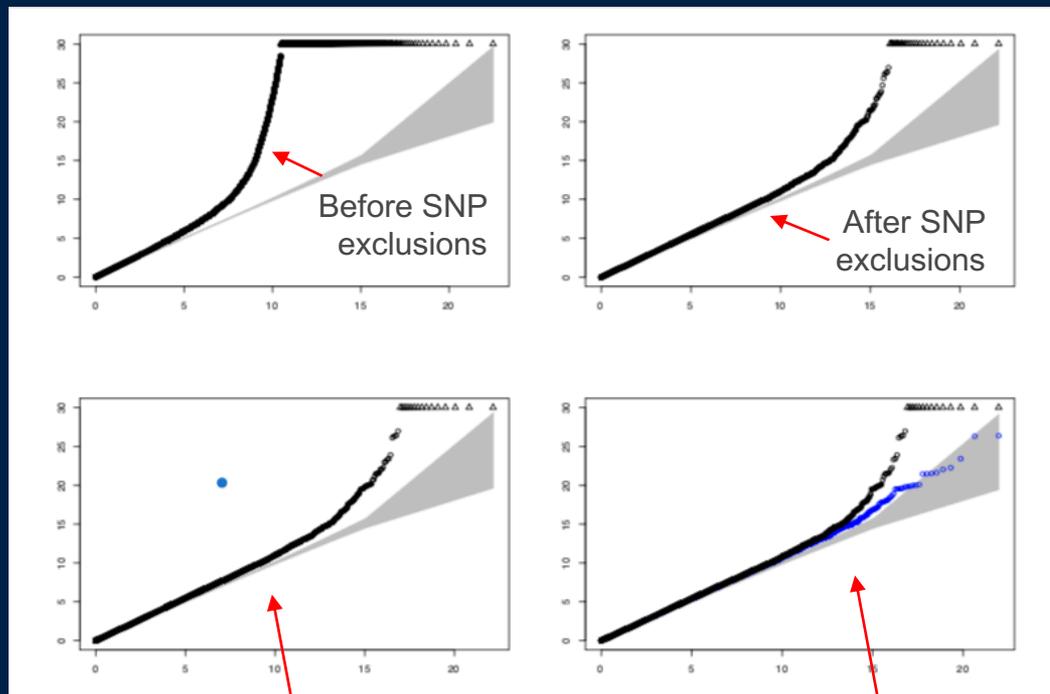
They used a method called *principal components analysis* to detect ancestry against the HapMap project samples. Some non-European ancestry individuals had been typed.

153 individuals were excluded on this basis.

PCA computes genome-wide relationships between samples and then looks for directions of greatest variation. Since relatedness typically decreases with geographic distance, principal components typically reflect geography.



Using quantile-quantile plots to assess residual confounding



After visually inspecting cluster plots for remaining associated SNPs

(Blue dots)... and after removing remaining strongly-associated regions that they claim to be real

- They also excluded 25,567 SNPs from the study for
- High missing data rates
 - Deviation from Hardy-Weinberg equilibrium (lecture 1) in controls
 - Frequency differences between the two control groups
 - And they visually inspected cluster plots for remaining SNPs

If there are few true signals, and if we have removed confounders – then P-values should largely come from a uniform distribution – they should lie on the diagonal.

Anatomy of a GWAS – what to look for

1. Collect as many cases and controls as possible

N=2,000 cases and
3,000 controls

2. Genotype (or impute) them at as many variants across the genome as possible

Genotyped at 500k
SNPs

3. Deal with potential confounders – careful data quality control and handle population structure.

Have they done adequate
data quality control?
Have they dealt with
possible confounders?

4. Estimate relative risks, and look for statistical evidence that of $RR \neq 1$

5. If estimate is many standard deviations from zero, bingo! We may have found a true causal effect.

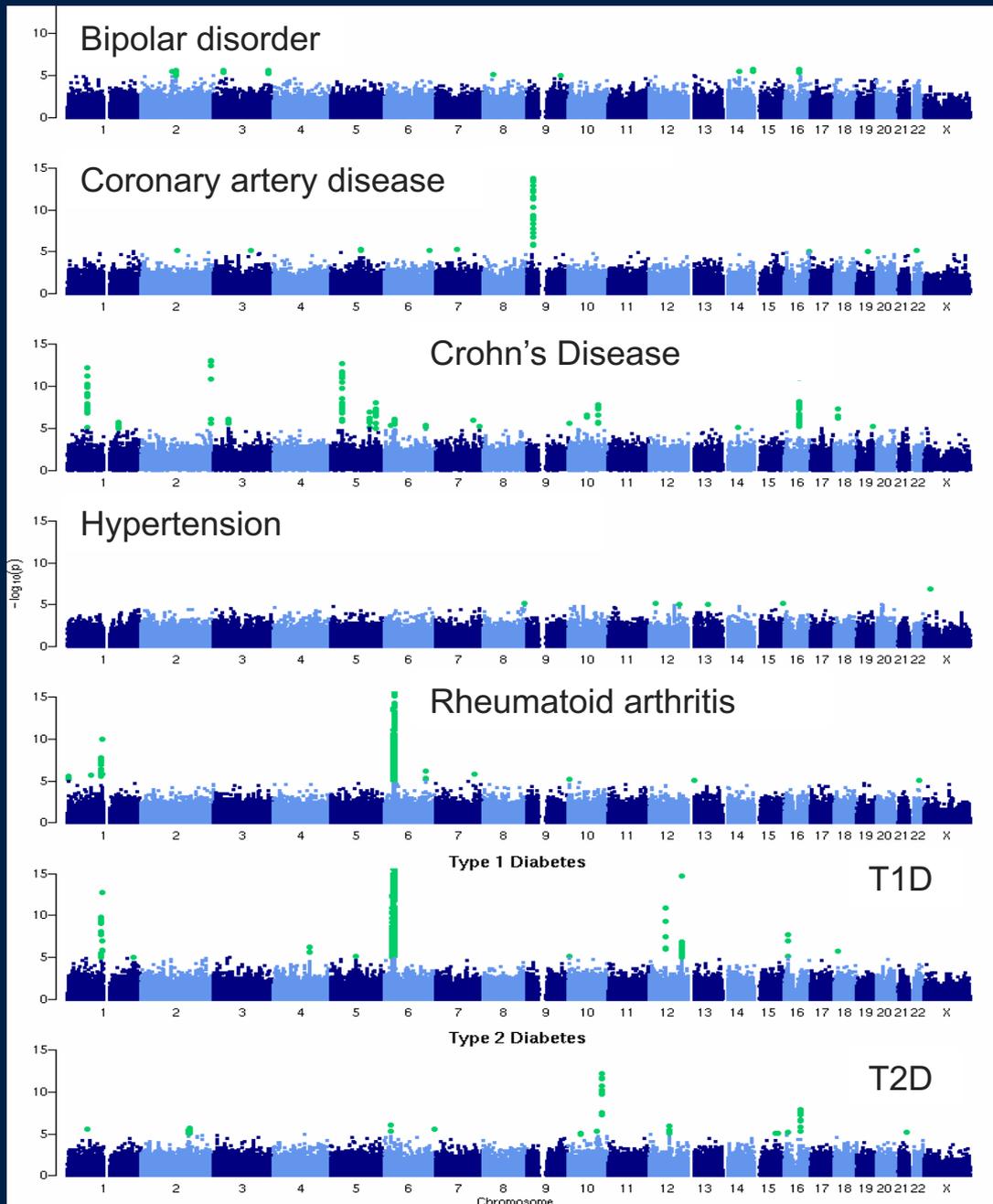
Did they find anything
with strong evidence?

6. Does it replicate in other studies, or have other corroborating evidence?

Is it convincing?

7. (Now try to understand the underlying biology.)

What about biology?



Number of associations with strong evidence

1

1

9

0

3

7

3

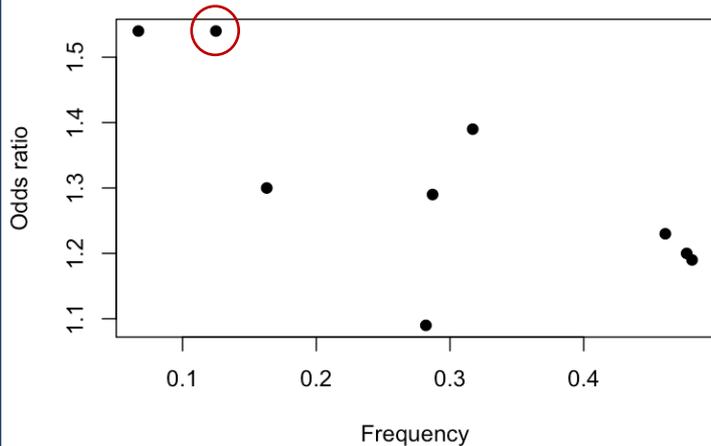
The study found 25 associations at their nominal P-value threshold.

Twelve of these provided replication of previously implicated variants. Thirteen were new associations.

The traits clearly differ in their genetic architecture

Some SNPs were associated with some evidence with multiple traits (mainly for the autoimmune diseases).

Frequency vs. effect size, WTCCC Crohn's disease

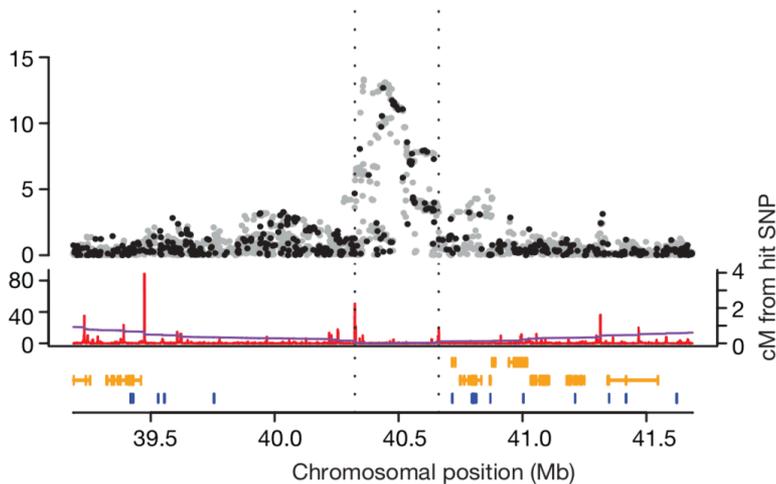


Effect sizes were generally modest

E.g. across the 9 associations with Crohn's disease, the maximum estimated odds ratio was 1.54, (similar to the O blood group example)

(A strong effect with Type 1 Diabetes was observed in the MHC locus)

CD hit region, chromosome 5



Zooming into these associations gives us a more detailed picture of the regional association – here shown for the strong association on chromosome 5.

Zooming in to a GWAS 'hit' plot

Sometimes called a 'locus zoom' plot. Here are some things to look for:

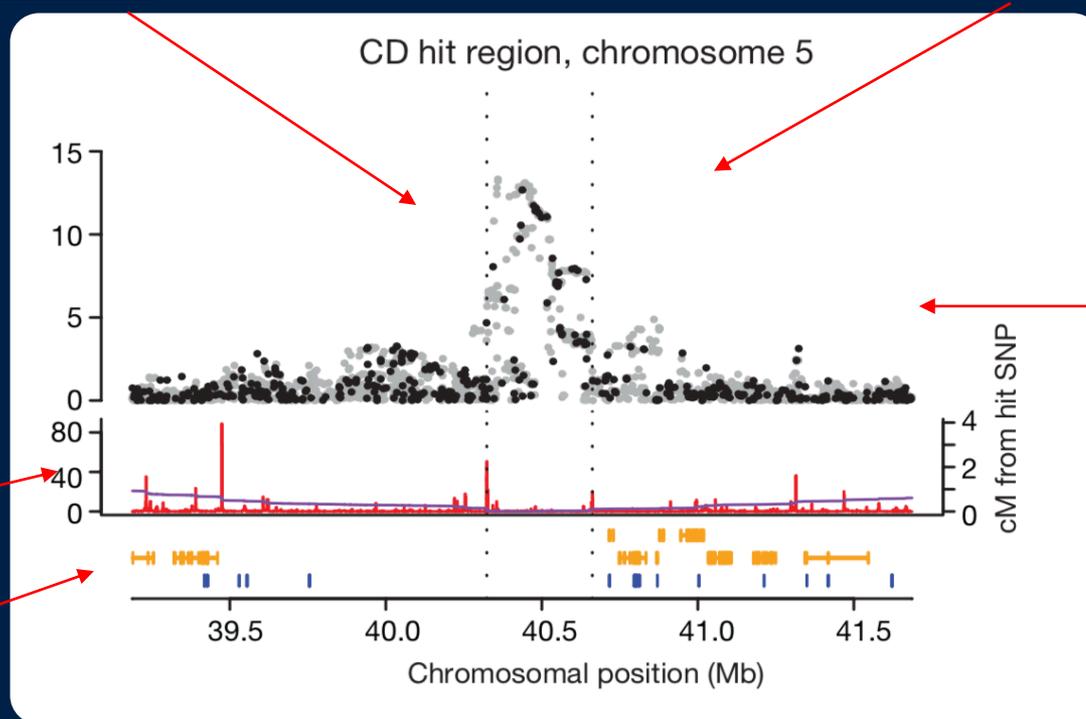
Evidence for association with each SNP
($-\log_{10}$ P-value or \log_{10} Bayes factor)

Delineation of association region boundaries (usually based on heuristics)

Black points were typed, grey points were imputed from HapMap

The recombination rate, here as estimated by HapMap

Regional genes



Signal ought to follow LD patterns. In particular ought to drop off near recombination hotspots

Position of SNPs in the reference genome assembly

Table 4 | Regions of the genome showing moderate evidence of association

Collection	Chromosome	Region (Mb)	SNP	Trend P-value	Genotypic P-value	log ₁₀ (BP) _{additive}	log ₁₀ (BP) _{general}	Rise allele	Minor allele	Heterozygote odds ratio	Homozygote odds ratio	Control MAF	Case MAF
BD	2p25	11.94-12.00	rs4027132	1.31 × 10 ⁻⁰⁵	9.68 × 10 ⁻⁰⁶	3.07	2.84	A	G	1.39 (1.19-1.64)	1.51 (1.27-1.79)	0.459	0.414
BD	2q12	10.41-104.58	rs7570682	3.11 × 10 ⁻⁰⁶	1.64 × 10 ⁻⁰⁶	3.68	3.23	A	A	1.23 (1.09-1.40)	1.64 (1.28-2.12)	0.214	0.255
BD	2q14	115.63-116.11	rs1375144	2.43 × 10 ⁻⁰⁵	1.31 × 10 ⁻⁰⁵	3.80	2.92	A	G	1.32 (1.07-1.63)	1.59 (1.29-1.96)	0.337	0.291
BD	2q37	241.23-241.28	rs2953145	1.11 × 10 ⁻⁰⁵	6.57 × 10 ⁻⁰⁶	3.22	3.50	A	G	1.84 (1.31-2.58)	2.14 (1.53-2.98)	0.226	0.189
BD	3p23	32.26-32.33	rs4276227	4.57 × 10 ⁻⁰⁶	2.62 × 10 ⁻⁰⁵	3.52	3.04	C	T	1.20 (0.99-1.46)	1.49 (1.23-1.81)	0.371	0.326
BD	3q27	184.29-184.40	rs683395	2.30 × 10 ⁻⁰⁶	5.11 × 10 ⁻⁰⁶	3.87	3.73	G	G	1.47 (1.26-1.71)	1.30 (0.69-2.46)	0.080	0.109
BD	6p21	42.82-42.86	rs6458307	3.43 × 10 ⁻⁰⁶	4.35 × 10 ⁻⁰⁶	-0.80	2.84	T	T	0.84 (0.75-0.96)	1.39 (1.13-1.69)	0.312	0.321
BD	8q12	34.22-34.61	rs2609653	6.06 × 10 ⁻⁰⁶	-	3.44	3.21	C	C	1.43 (1.19-1.71)	3.62 (1.26-10.44)	0.052	0.074
BD	9q32	114.31-114.39	rs10982256	8.80 × 10 ⁻⁰⁶	4.41 × 10 ⁻⁰⁵	3.23	2.37	T	T	1.26 (1.08-1.47)	1.47 (1.24-1.74)	0.471	0.425
BD	14q22	57.17-57.24	rs10134944	3.21 × 10 ⁻⁰⁶	6.89 × 10 ⁻⁰⁶	3.73	3.59	T	T	1.45 (1.24-1.68)	1.32 (0.74-2.33)	0.086	0.115
BD	14q32	103.43-103.62	rs11622475	2.10 × 10 ⁻⁰⁶	8.14 × 10 ⁻⁰⁶	3.87	3.24	C	T	1.13 (0.89-1.44)	1.47 (1.17-1.86)	0.300	0.256
BD	16q12	51.36-51.50	rs1344484	1.64 × 10 ⁻⁰⁶	1.03 × 10 ⁻⁰⁶	3.94	3.41	T	C	1.24 (1.03-1.48)	1.52 (1.27-1.82)	0.402	0.353
BD	20p13	3.70-3.73	rs3761218	4.43 × 10 ⁻⁰⁵	6.71 × 10 ⁻⁰⁶	2.58	3.18	T	C	0.97 (0.81-1.15)	1.31 (1.09-1.57)	0.397	0.356
CAD	14q3	236.77-236.85	rs17672135	1.04 × 10 ⁻⁰⁴	2.35 × 10 ⁻⁰⁶	2.36	3.88	T	C	0.70 (0.61-0.81)	1.32 (0.79-2.22)	0.134	0.108
CAD	5q21	99.98-100.11	rs383830	5.72 × 10 ⁻⁰⁶	1.34 × 10 ⁻⁰⁵	3.49	3.22	A	A	1.60 (1.16-2.21)	1.92 (1.40-2.63)	0.220	0.182
CAD	6q25	151.34-151.42	rs6922269	6.33 × 10 ⁻⁰⁶	1.50 × 10 ⁻⁰⁶	3.38	3.14	A	A	1.17 (1.04-1.32)	1.65 (1.32-2.06)	0.253	0.294
CAD	16q23	81.72-81.79	rs8055236	9.73 × 10 ⁻⁰⁶	5.60 × 10 ⁻⁰⁶	3.28	3.59	G	T	1.91 (1.33-2.74)	2.23 (1.56-3.17)	0.198	0.162
CAD	19q12	34.74-34.78	rs7250581	9.12 × 10 ⁻⁰⁶	2.50 × 10 ⁻⁰⁵	3.30	2.87	G	A	1.06 (0.79-1.43)	1.40 (1.05-1.86)	0.220	0.182
CAD	22q12	25.01-25.06	rs688034	6.90 × 10 ⁻⁰⁶	3.75 × 10 ⁻⁰⁶	3.33	3.15	T	T	1.11 (0.98-1.25)	1.62 (1.34-1.95)	0.310	0.355
CD	1q24	169.53-169.67	rs12037606	1.79 × 10 ⁻⁰⁶	1.09 × 10 ⁻⁰⁵	3.89	3.35	A	A	1.22 (1.07-1.40)	1.52 (1.28-1.82)	0.388	0.438
CD	5q23	131.40-131.90	rs6596075	5.40 × 10 ⁻⁰⁶	3.19 × 10 ⁻⁰⁵	4.54	4.01	C	G	1.55 (1.00-2.39)	2.06 (1.35-3.14)	0.166	0.127
CD	6q22	20.83-20.85	rs6908425	5.13 × 10 ⁻⁰⁶	1.10 × 10 ⁻⁰⁶	3.55	3.38	C	T	1.63 (1.18-2.25)	1.95 (1.43-2.67)	0.230	0.190
CD	6p21	32.79-32.91	rs4969220	8.65 × 10 ⁻⁰⁷	2.28 × 10 ⁻⁰⁶	4.19	3.92	A	A	1.14 (0.98-1.32)	1.52 (1.28-1.79)	0.481	0.534
CD	6q23	138.06-138.17	rs7753394	4.42 × 10 ⁻⁰⁶	2.59 × 10 ⁻⁰⁵	3.52	2.99	C	C	1.21 (1.04-1.40)	1.48 (1.25-1.76)	0.482	0.531
CD	7q36	147.62-147.70	rs7807268	6.89 × 10 ⁻⁰⁶	4.42 × 10 ⁻⁰⁶	3.33	3.58	G	G	1.38 (1.20-1.60)	1.47 (1.24-1.74)	0.462	0.509
CD	10p15	38.52-38.57	rs6601764	2.56 × 10 ⁻⁰⁶	8.95 × 10 ⁻⁰⁶	3.74	3.01	C	C	1.16 (1.01-1.33)	1.52 (1.28-1.80)	0.408	0.458
CD	19q13	50.89-51.07	rs8111071	6.14 × 10 ⁻⁰⁶	1.75 × 10 ⁻⁰⁵	3.48	3.29	G	G	1.47 (1.25-1.73)	1.28 (0.56-2.88)	0.070	0.096
HT	14q3	235.67-235.79	rs2820037	5.76 × 10 ⁻⁰⁵	7.66 × 10 ⁻⁰⁷	2.54	3.99	T	T	1.54 (1.03-2.31)	1.09 (0.74-1.62)	0.141	0.171
HT	8q24	140.17-140.35	rs6997709	7.88 × 10 ⁻⁰⁶	4.36 × 10 ⁻⁰⁵	3.32	2.60	G	T	1.20 (0.94-1.52)	1.49 (1.18-1.89)	0.285	0.244
HT	12p12	24.86-24.95	rs7961152	7.39 × 10 ⁻⁰⁶	3.03 × 10 ⁻⁰⁵	3.29	2.51	A	A	1.16 (1.01-1.32)	1.47 (1.25-1.74)	0.415	0.461
HT	12q23	100.52-100.58	rs1110912	9.18 × 10 ⁻⁰⁶	1.94 × 10 ⁻⁰⁶	3.27	3.11	G	G	1.33 (1.18-1.51)	1.34 (0.96-1.86)	0.165	0.200
HT	15q21	66.90-67.04	rs1937506	9.29 × 10 ⁻⁰⁶	4.53 × 10 ⁻⁰⁵	3.25	2.85	G	A	1.33 (1.04-1.69)	1.60 (1.26-2.02)	0.289	0.248
HT	15q26	94.60-94.67	rs2398162	7.85 × 10 ⁻⁰⁶	5.67 × 10 ⁻⁰⁶	3.33	3.40	A	G	0.97 (0.76-1.25)	1.31 (1.03-1.67)	0.258	0.218
RA	1p36	2.44-2.77	rs6684865	5.37 × 10 ⁻⁰⁶	3.14 × 10 ⁻⁰⁵	3.47	2.97	G	C	1.27 (1.02-1.56)	1.54 (1.25-1.90)	0.338	0.294
RA	1p31	80.16-80.36	rs11162922	1.80 × 10 ⁻⁰⁶	1.80 × 10 ⁻⁰⁶	4.11	3.80	A	G	1.27 (0.41-4.01)	2.00 (0.64-6.20)	0.072	0.048
RA	4q15	24.99-25.13	rs3816587	7.65 × 10 ⁻⁰⁶	9.25 × 10 ⁻⁰⁶	0.50	2.64	C	C	0.91 (0.80-1.04)	1.35 (1.14-1.59)	0.406	0.434
RA	6q23	138.00-138.06	rs6920220	4.99 × 10 ⁻⁰⁶	1.58 × 10 ⁻⁰⁵	3.49	3.17	A	A	1.20 (1.06-1.36)	1.72 (1.33-2.22)	0.223	0.263
RA	7q32	130.80-130.84	rs11761231	1.74 × 10 ⁻⁰⁶	2.65 × 10 ⁻⁰⁶	3.92	3.42	C	T	1.44 (1.19-1.75)	1.64 (1.35-1.99)	0.375	0.327
RA	10p15	6.07-6.16	rs2104286	7.02 × 10 ⁻⁰⁶	2.52 × 10 ⁻⁰⁵	3.37	2.57	T	C	1.41 (1.10-1.81)	1.68 (1.31-2.14)	0.286	0.244
RA	13q12	19.845-19.855	rs9550642	8.44 × 10 ⁻⁰⁶	3.90 × 10 ⁻⁰⁶	3.35	3.02	A	A	1.34 (1.15-1.56)	2.23 (1.21-4.13)	0.084	0.112
RA	21q22	41.430-41.465	rs2837960	3.45 × 10 ⁻⁰²	1.68 × 10 ⁻⁰⁶	0.05	2.70	G	G	0.95 (0.83-1.08)	2.30 (1.64-3.23)	0.171	0.188
RA	22q13	35.870-35.885	rs743777	7.92 × 10 ⁻⁰⁶	1.15 × 10 ⁻⁰⁶	3.29	3.52	G	G	1.09 (0.97-1.24)	1.72 (1.40-2.11)	0.292	0.336
T1D	14q2	221.92-222.17	rs2639703	8.46 × 10 ⁻⁰⁶	1.74 × 10 ⁻⁰⁵	3.25	3.06	C	C	1.15 (1.02-1.30)	1.61 (1.31-1.99)	0.276	0.318
T1D	4q27	123.02-123.92	rs17388568	5.01 × 10 ⁻⁰⁶	3.27 × 10 ⁻⁰⁶	4.42	3.89	A	A	1.26 (1.11-1.42)	1.58 (1.27-1.95)	0.260	0.307
T1D	5q14	86.20-86.50	rs2544677	8.23 × 10 ⁻⁰⁶	4.43 × 10 ⁻⁰⁶	3.32	2.70	C	G	1.34 (1.00-1.79)	1.65 (1.24-2.18)	0.242	0.204
T1D	5q31	132.64-132.67	rs17166496	6.06 × 10 ⁻⁰¹	5.20 × 10 ⁻⁰⁶	-0.97	3.25	G	G	0.77 (0.68-0.87)	1.09 (0.92-1.29)	0.391	0.386
T1D	10p15	6.07-6.18	rs2104286	7.96 × 10 ⁻⁰⁶	4.32 × 10 ⁻⁰⁵	3.31	2.88	T	C	1.30 (1.02-1.65)	1.57 (1.25-1.99)	0.286	0.245
T1D	12p13	9.71-9.80	rs11052552	1.02 × 10 ⁻⁰⁴	7.24 × 10 ⁻⁰⁷	2.22	3.80	G	T	1.49 (1.28-1.73)	1.43 (1.21-1.69)	0.486	0.446
T1D	18p11	12.76-12.91	rs2542151	1.89 × 10 ⁻⁰⁶	1.16 × 10 ⁻⁰⁶	3.91	3.52	G	G	1.30 (1.15-1.47)	1.62 (1.17-2.24)	0.163	0.201
T2D	1q31	66.04-66.36	rs4655995	2.68 × 10 ⁻⁰⁶	1.33 × 10 ⁻⁰⁵	3.81	3.47	G	G	1.37 (1.17-1.59)	2.33 (1.23-4.42)	0.080	0.108
T2D	2q24	160.90-161.17	rs6718526	2.40 × 10 ⁻⁰⁶	1.16 × 10 ⁻⁰⁵	3.86	3.35	C	T	1.49 (1.05-2.11)	1.86 (1.32-2.63)	0.209	0.171
T2D	3p14	55.24-55.32	rs358806	4.77 × 10 ⁻⁰¹	3.05 × 10 ⁻⁰⁶	-0.83	2.72	A	A	0.86 (0.75-0.97)	1.78 (1.34-2.36)	0.198	0.204
T2D	4q27	122.92-123.02	rs7659604	2.1 × 10 ⁻⁰²	9.42 × 10 ⁻⁰⁶	0.13	2.74	T	T	1.35 (1.19-1.54)	1.09 (0.91-1.30)	0.380	0.403
T2D	10q11	43.43-43.63	rs9226506	7.78 × 10 ⁻⁰⁶	2.99 × 10 ⁻⁰⁶	3.27	2.92	C	C	1.28 (1.11-1.48)	1.46 (1.24-1.72)	0.492	0.538
T2D	12q13	49.50-49.87	rs12304921	5.37 × 10 ⁻⁰²	7.07 × 10 ⁻⁰⁶	-0.09	2.68	G	G	2.50 (1.53-4.09)	1.94 (1.20-3.15)	0.145	0.159
T2D	12q15	69.58-69.96	rs1495377	1.31 × 10 ⁻⁰⁶	6.52 × 10 ⁻⁰⁶	4.01	3.15	G	G	1.28 (1.11-1.49)	1.51 (1.28-1.78)	0.497	0.547
T2D	15q24	72.24-72.50	rs2930291	7.72 × 10 ⁻⁰⁶	4.40 × 10 ⁻⁰⁵	3.30	2.42	G	A	1.25 (1.04-1.51)	1.50 (1.24-1.82)	0.377	0.332
T2D	15q25	78.12-78.36	rs2903265	9.57 × 10 ⁻⁰⁶	4.98 × 10 ⁻⁰⁵	3.24	2.53	G	A	1.18 (0.93-1.49)	1.47 (1.17-1.86)	0.284	0.243

Regions with at least one SNP with a P value of greater than 5×10^{-7} and less than 1×10^{-5} for either the trend or the genotypic test. Columns as for Table 3. Cluster plots for each SNP have been inspected visually. Positions are in NCBI build-35 coordinates. Genotypic P values were not calculated for SNPs with the lowest MAFs owing to low numbers of rare-allele homozygotes and sensitivity to genotype calling errors.

The results above used a P-value threshold of $P < 5 \times 10^{-7}$

They also reported a longer list of association at lesser levels of evidence ($P < 5 \times 10^{-7}$). Many of these must be real as well.

How much statistical evidence do we really need? How did they choose a good threshold?

How to choose a P-value threshold

They reasoned like this: Based on what we know from HapMap, there are maybe 1 million 'LD blocks' in the human genome. Suppose maybe 10 of them, or so, are associated with the trait. Then the prior chance of association for a randomly chosen region (i.e. chosen 'hypothesis free') will be 10 in a million, i.e. plausibly

Prior odds = 1×10^{-5} before we see any data.

For a P-value threshold α it works out that:

$$\text{odds(associated} | P < \alpha) = \frac{\text{statistical power}}{\alpha} \times \text{prior odds}$$

=> If the statistical power is 50%, say, then setting $\alpha = 5 \times 10^{-7}$ will give a posterior odds of 10 to 1.

This was a good choice! All of their associations have subsequently replicated in larger studies.

Many GWAS use a more stringent $\alpha = 5 \times 10^{-8}$ threshold, while still others attempt to directly estimate the above (c.f. 'False discovery rate' methods).

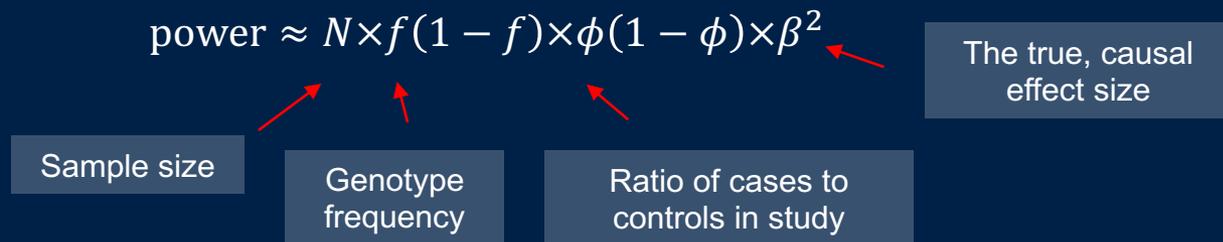
Statistical power

The statistical power says “how likely are we to detect a true effect”. It is essentially determined by:

- The true effect size β (which of course we don't know beforehand)

- The standard error, which we do know approximately $se \approx \frac{1}{\sqrt{N \times f(1-f) \times \phi(1-\phi)}}$

- And also the threshold α , which says ‘how many standard errors away from zero do we need?’



Anatomy of a GWAS – what to look for

1. Collect as many cases and controls as possible

N=2,000 cases and
3,000 controls

2. Genotype (or impute) them at as many variants across the genome as possible

Genotyped at 500k
SNPs

3. Deal with potential confounders – careful data quality control and handle population structure.

Have they done adequate
data quality control?
Have they dealt with
possible confounders?

4. Estimate relative risks, and look for statistical evidence that of $RR \neq 1$

5. If estimate is many standard deviations from zero, bingo! We may have found a true causal effect.

Did they find anything with
strong evidence?

6. Does it replicate in other studies, or have other corroborating evidence?

Is it convincing?

7. (Now try to understand the underlying biology.)

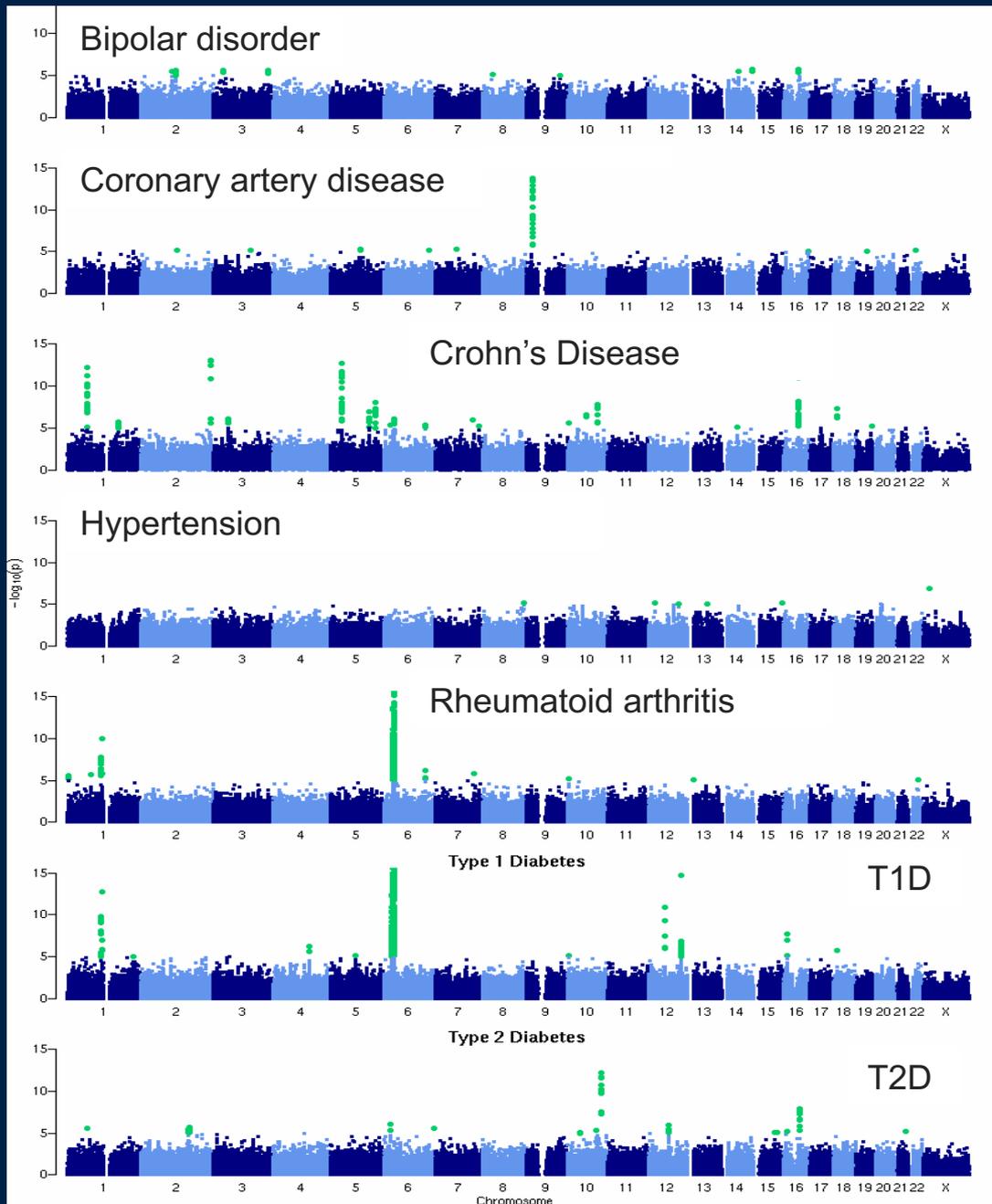
What about biology?

Summary

- In the early 2000's, The HapMap and other projects mapped SNPs genome-wide, and described human haplotype variation and patterns of LD. High-throughput genotyping microarray technology was developed to type these SNPs.
- LD turned out to be more extensive than previously thought – it has a 'blocklike' structure due to recombination happening in hotspots. This enables GWAS even without typing everything in the genome.
- The WTCCC was one of the first large GWAS studies, and demonstrated beyond doubt that GWAS methodology could work and produce new discoveries.
- Although the overall design is simple, we are looking for small differences in risk between cases and controls (often $RR = 1.5$ or smaller). Consequently a lot of careful work is needed to ensure there is no subtle confounding – e.g. from sample collection, genotyping and data quality issues, or environmental covariates.

Lecture outline

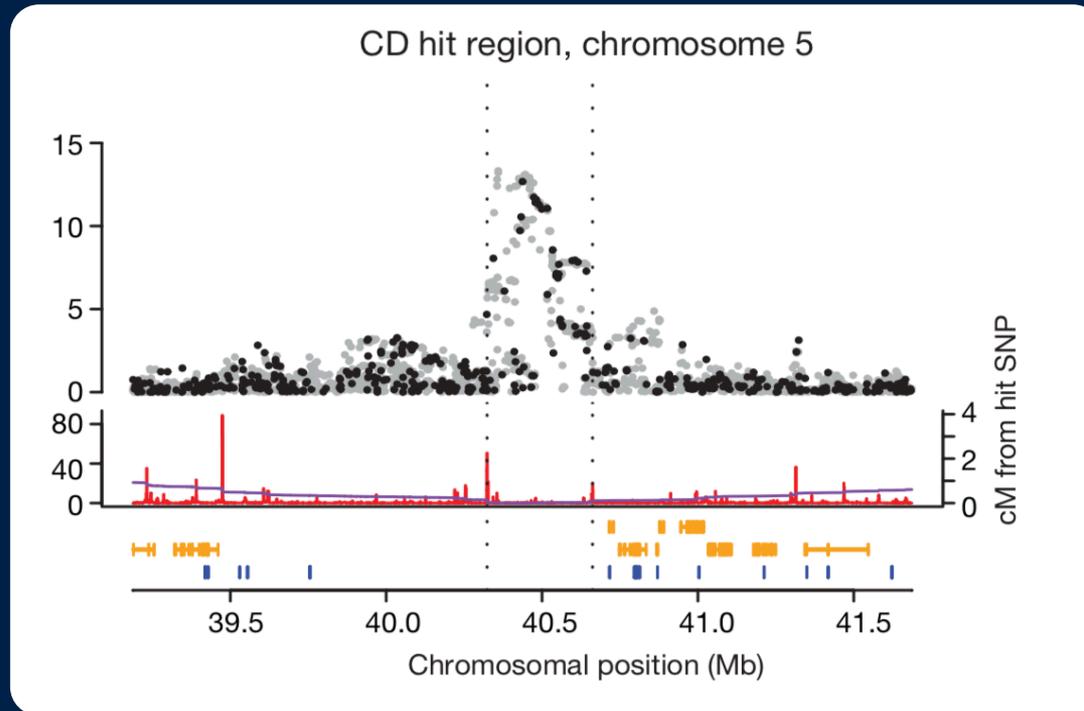
- Background
- Searching for a needle in a haystack
- Genome-wide association studies in theory
- GWAS in practice
- • The challenge of understanding biology



We have clearly learned *something* about the biology of these traits.

What about the underlying causal variants?

The challenge of understanding biology



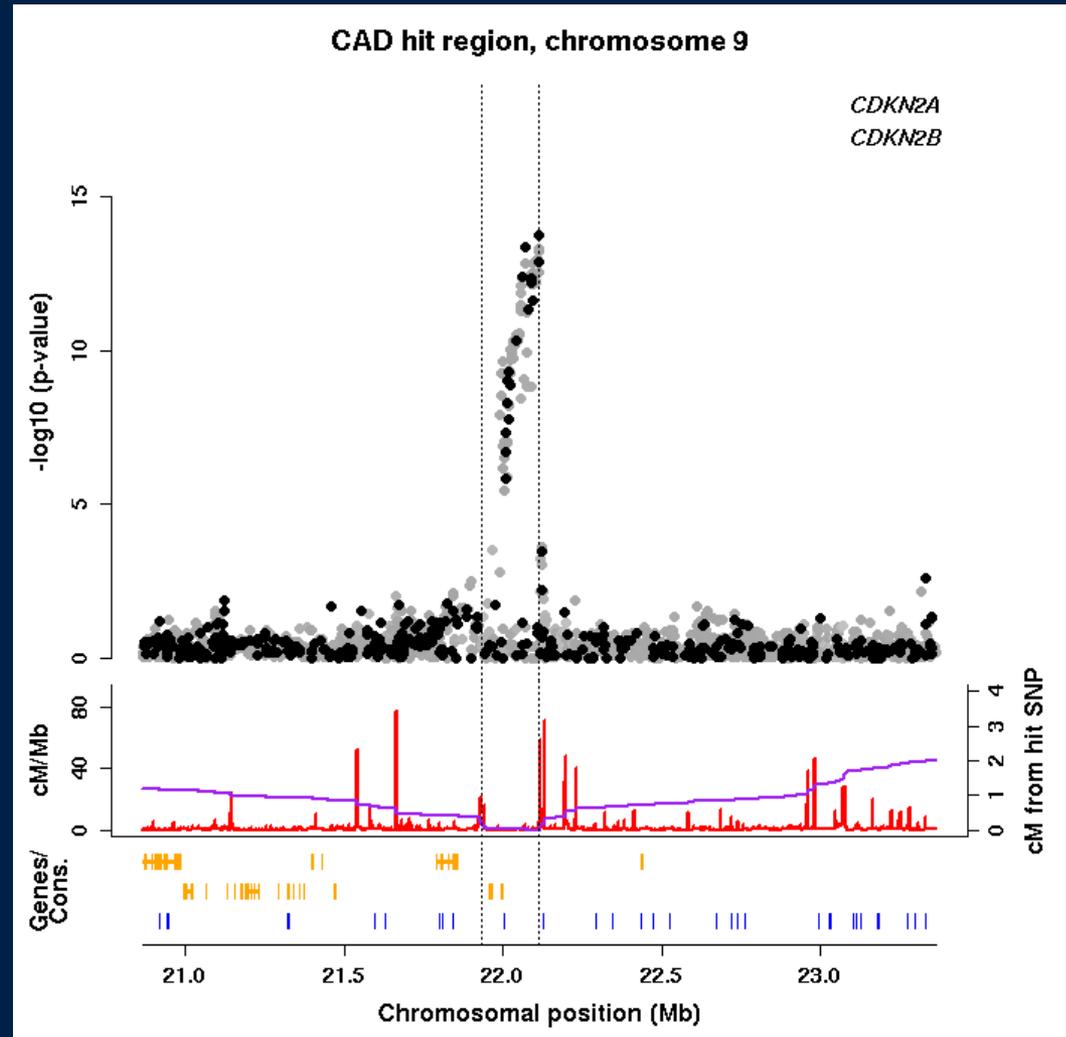
No genes under the main association signal!

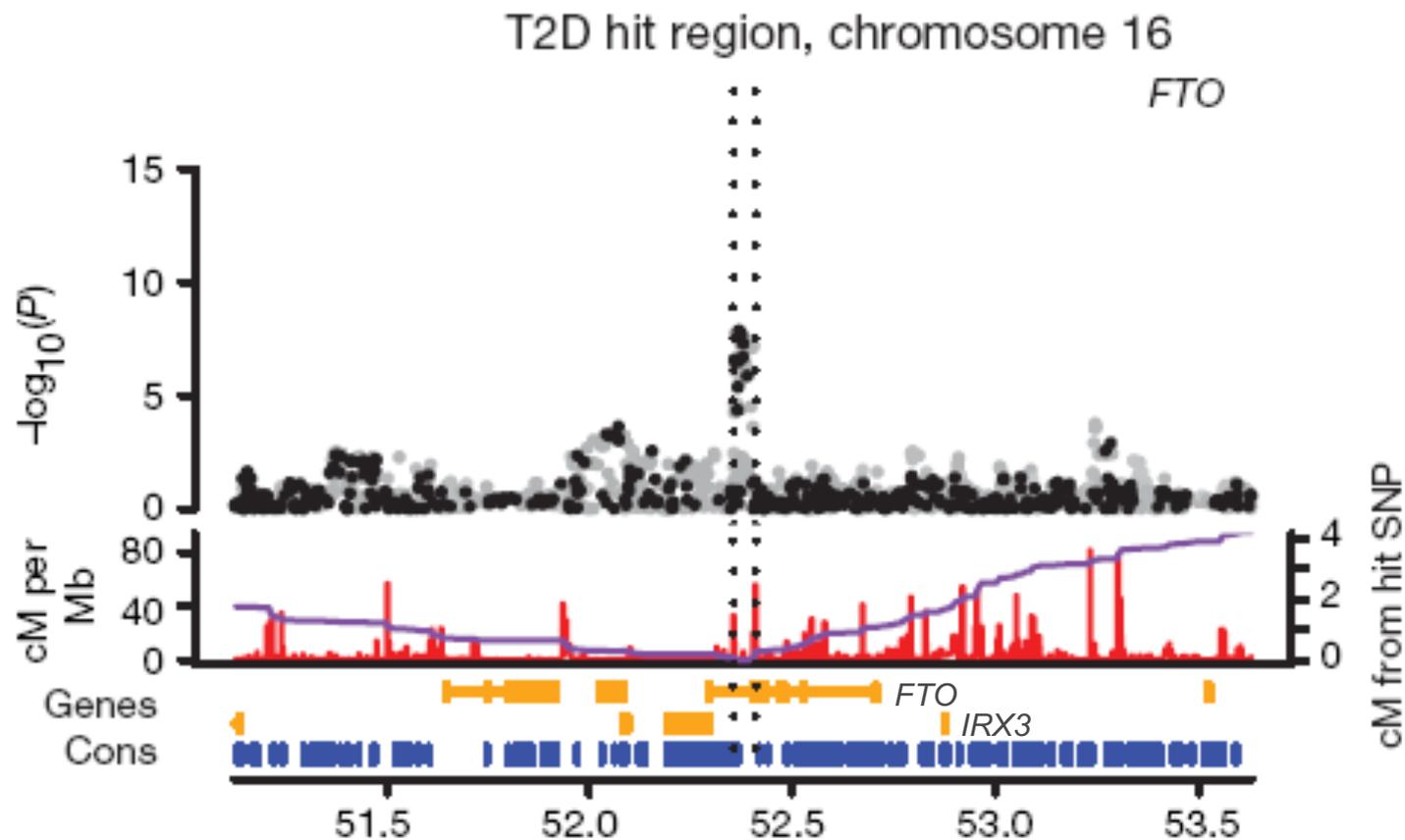
Biology is complicated

Association observed with CAD over a ~100kb region of chromosome 9. This is unquestionably a real association (it has been replicated in several independent studies).

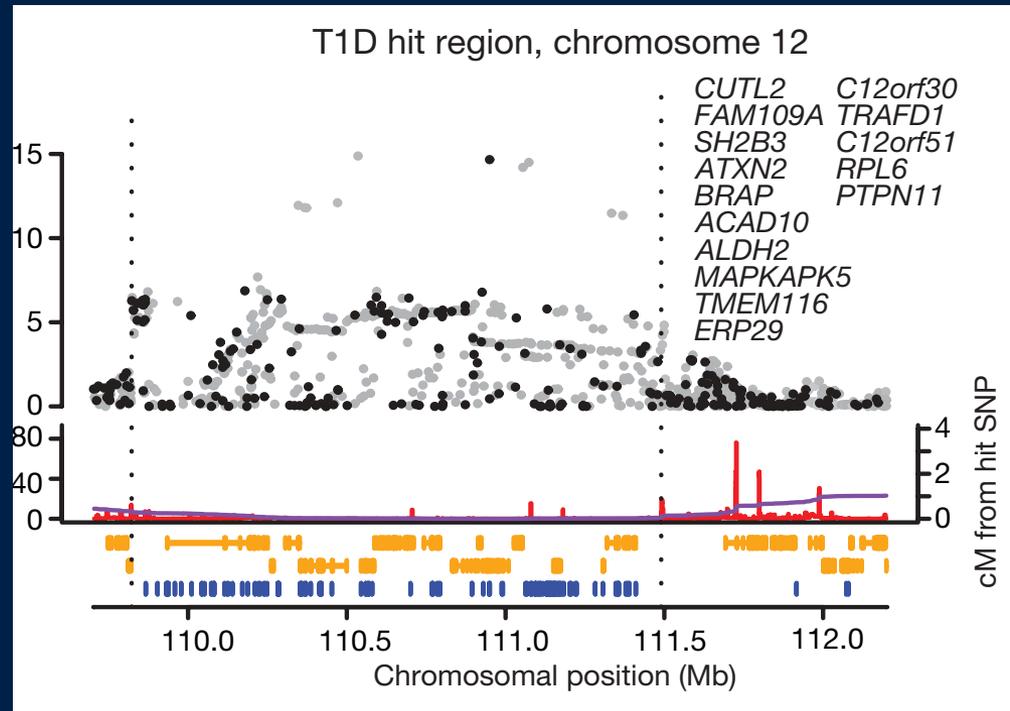
The functional mechanism of this association is not fully solved; it probably involves regulation of expression of the two nearby genes *CDKN2A/B*.

Neither gene was an obvious candidate beforehand - thus, this association does point to novel biology.





This association with Type 2 Diabetes turned out to be through a second, related trait (obesity), again unquestionably a real effect. But as of 2018 the functional mechanism remains unclear. Expression of *FTO* is known to affect obesity, but the SNPs may also affect expression of another gene, *IRX3*, 200kb away.



This pattern has turned out to be typical. It has generally proven extremely hard to narrow down GWAS associations to underlying 'causal' variants.

LD is a double-edged sword.

Next lecture: we will look at this.

How to read a GWAS – checklist

What is the sample size?

How are the samples genotyped? Are cases and controls typed in the same way?

What have the authors done to deal with potential confounders – good data quality control? Population structure? Is it convincing?

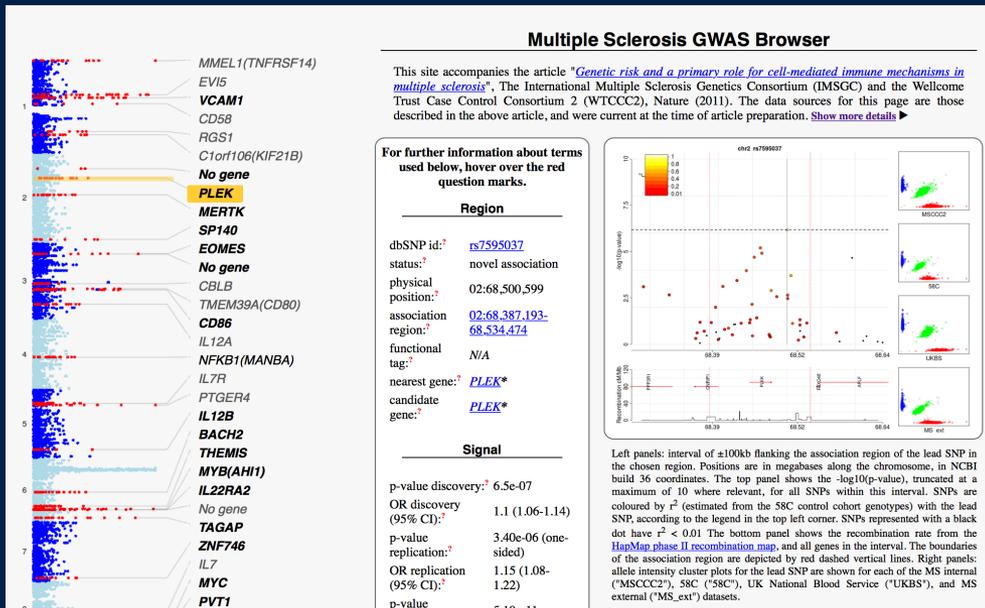
Do the results look sensible? Are the effect sizes reasonable? How strong is the evidence?

Does the signal replicate?

Does the association follow patterns of LD?

If all the above seem fine – what genes are nearby? Can you figure out biology?

Consolidation question



GWAS of multiple sclerosis (2011)

9772 cases, 17,376 controls from across Europe

www.well.ox.ac.uk/wtccc2/ms/
(I think this requires the trailing /)

- Visit the above site and make sure you understand what is shown. Pick a signal and try to work out
- What is the estimated effect size?
 - How strong was the evidence?
 - Did it replicate?
 - Does the association signal look sensible – does it follow LD patterns, and do the cluster plots look sensible?
 - Can you figure out what the nearby genes do? (**Warning:** this can be a time sink!)

Bonus question: read the paper and try to figure out the questions on the checklist.

Next lecture: Feb 24th

Genome-wide association studies II: Identifying genetic associations with complex traits

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