

Genetics of common complex psychiatric disorders I

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Part 1: Biometrics

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Genetics and Environmental Influences on Behaviour and Mental Health

What is a “common”, “complex” psychiatric disorder?

Common: Affects 1% or more of the population

Complex: Inheritance cannot be explained by a single gene

Psychiatric disorders are defined by disruption to higher-order brain functions of moods, perceptions, thoughts, beliefs, and behaviours but usually in the absence of major neurological impairments (consciousness, senses, memory). Psychiatric disorders include depressive and anxiety disorders (major depressive disorder, panic disorder), manic and psychotic disorders (bipolar disorder, schizophrenia), obsessive-compulsive disorders, eating disorders (anorexia nervosa, bulimia nervosa), substance-use disorders and personality disorders. Childhood conditions like attention-deficit/hyperactivity and autism can also be included, but only when they lead to clinically-salient impairment or distress. There are also many shades of sadness, hallucinations, eccentricities, mood swings, body-image preoccupations, recreational substances use, personalities, etc that are not psychiatric disorders but may still be informative to study from an aetiological and genetic standpoint.

- Sullivan PF and Geschwind DH (2019) [Defining the Genetic, Genomic, Cellular, and Diagnostic Architectures of Psychiatric Disorders](https://doi.org/10.1016/j.cell.2019.01.015). *Cell* doi:10.1016/j.cell.2019.01.015

Psychiatric diagnoses

- Depressive disorder: Marked and persistent low mood and inability to feel pleasure.
- Anxiety disorder: Extensive, pervasive, unrealistic, and disabling worry.
- Bipolar disorder: Intense mood swings between mania and depression.
- Schizophrenia: Persistent hallucinations and delusions that severely impair functioning, highly disorganised thought and speech.
- Eating disorders: self-starvation or excessive over-eating, debilitating preoccupation with body image.
- Attention-deficit/hyperactivity disorder: Early onset, developmentally inappropriate and persistent hyperactivity and inattention.
- Autism spectrum disorders: Markedly impaired interpersonal interactions, non-goal-directed behaviours, and language development.

Sullivan, P. F., Daly, M. J. & O'Donovan, M. Genetic architectures of psychiatric disorders: the emerging picture and its implications. *Nat Rev Genet* 13, 537 551 (2012).

Frances, A. *Essential of Psychiatric Diagnosis*. Guilford Press (2013).

- Depression: 3% in a week
- Schizophrenia: 1% in lifetime
- Bipolar disorder: 2% in lifetime
- Anxiety disorder: 6% in a week

Psychiatric disorders have many causes, correlates, and consequences (genetics, environment, family life, substance use, relationships)

Incidence of psychiatric disorders range from the common (depression, anxiety) to the rare (bipolar disorder, schizophrenia).

Why genetics?

Why use genetics to study mental health and psychiatric disorders?

- Biological understanding of genes, pathways
- Shared aetiology with other disorders
- Risk prediction
- Drug retargeting
- Causal analysis of environmental risk factors

Genetics of categorical traits















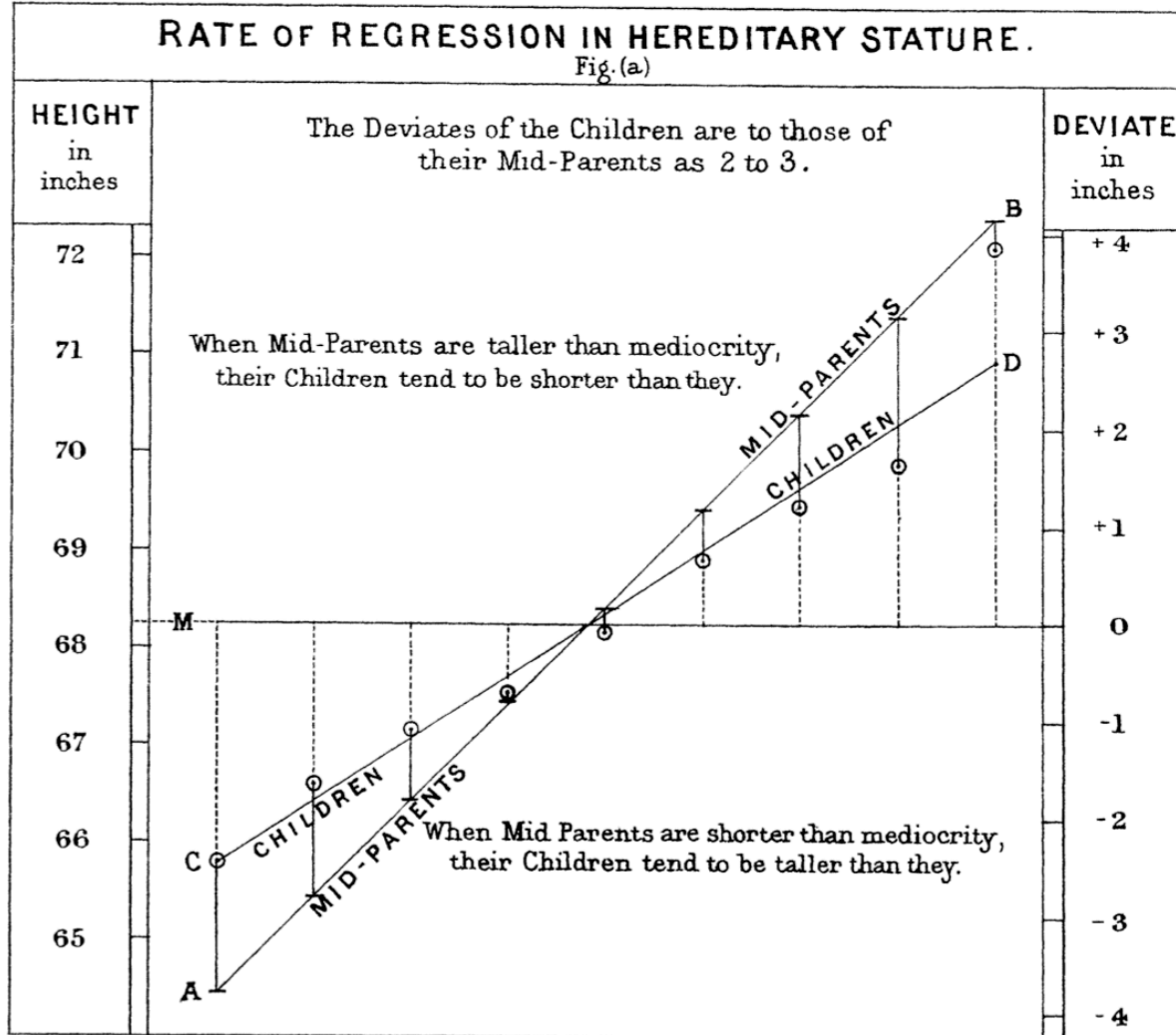
Seed		Flower	Pod		Stem	
Form	Cotyledons	Color	Form	Color	Place	Size
						
Grey & Round	Yellow	White	Full	Yellow	Axial pods, Flowers along	Long (6-7ft)
						
White & Wrinkled	Green	Violet	Constricted	Green	Terminal pods, Flowers top	Short (1/2 - 1ft)
1	2	3	4	5	6	7

Figure 1: Diagram showing the seven “characters” observed by Mendel

Gregor Mendel (1822–1884), working in what is now Czechia, discovered the transmission of traits from parents to offspring could be explained by the inheritance of two “elements”, which we now call alleles. Mendel was concerned with discrete or categorical phenotypes.

[Mendel pea plant figure](#) by Mariana Ruiz (LadyofHats) [public domain]

Genetics of continuous traits

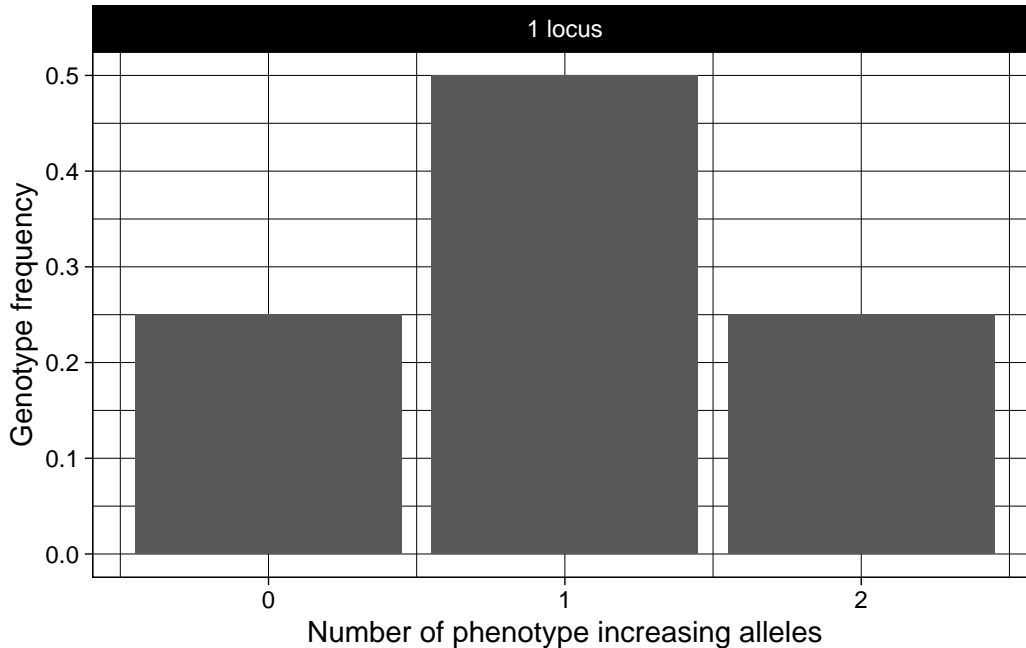


Separately, Francis Galton (1822–1911), was studying the inheritance of continuous or metric phenotypes. He noticed the parents who were tall tended to have children that were slightly shorter than themselves (and vice versa). This was termed “regression to the mean” from which the name of the statistical method “regression” is derived.

“In their search for universal hereditary laws, Galton and Pearson were driven by the linear model and the normal distribution because the associated parameters had scientific meaning for them that went beyond mere description.” - Wachsmuth, A., Wilkinson, L., & Dallal, G. E. (2003). [Galton's Bend](#). *The American Statistician*, 57(3), 190–192. doi:10.1198/0003130031874

For more on Galton's legacy, see <https://adelphigenetics.org/history/>

Reconciling categorical + continuous genetics = quantitative genetics

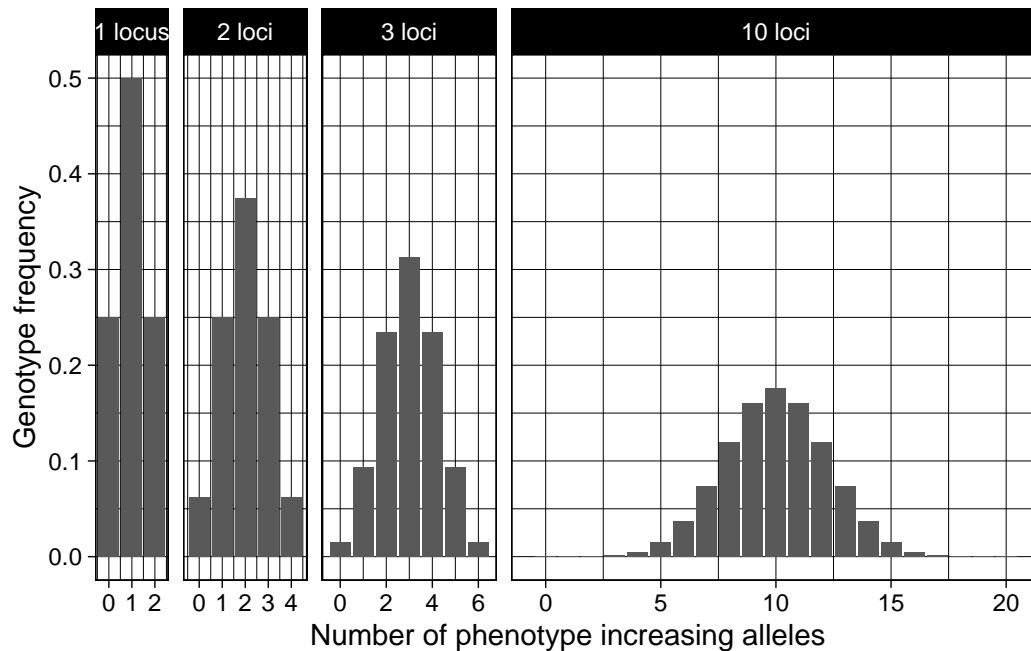


Ronald Fisher reconciled the inheritance of continuous and categorical phenotypes by showing that a continuous phenotype could be made from the inheritance of a large number (dozens, hundreds, or thousands) categorical genes. The term “variance” comes from Fisher’s discoveries.

- Fisher, R. A. (1918). [XV.—The correlation between relatives on the supposition of Mendelian inheritance](#). *Transactions of the Royal Society of Edinburgh*. doi:10.1017/S0080456800012163
- Charlesworth and Edwards (2018). [A century of variance](#). *Significance* 15(4). doi:10.1111/j.1740-9713.2018.01170.x
- Bodmer et al (2021) [The outstanding scientist, R.A. Fisher: his views on eugenics and race](#). *Heredity* doi:10.1038/s41437-020-00394-6

Polygenic traits are quantitative traits

Adding up effects from a large number of genetic effects to make a continuous phenotype is related to the Central Limit Theorem.



“R. A. Fisher’s 1918 paper, ‘The correlation between relatives on the supposition of Mendelian inheritance’, resolved the often bitter conflict between biometricians and Mendelians, which raged for a decade following the rediscovery of Mendel’s work. Fisher showed that a complex quantitative trait could be explained by Mendelian inheritance if several genes affect the trait.” Because he crossed true-breeding plants, Mendel’s experiments showed that a single locus with two alleles of equal frequency results in three genotypes (see the figure, part a). If the allelic effects are additive, the three genotypes produce three phenotypes; in the case of Mendel’s qualitative traits, the allelic effects showed complete dominance, so only two phenotypes were observed. However, assuming equal and additive effects, 2 genes yield 9 genotypes and 5 phenotypes (part b) and 3 genes yield 27 genotypes and 7 phenotypes (part c). With unequal and non-additive allelic effects and some environmental influence, three genes would result in a normal bell-shaped curve of continuous variation (part d). This logic assumes common alleles; rare alleles will skew the distribution. Genome-wide association research suggests that many more than three genes affect most traits, which underscores the expectation that polygenic traits are quantitative traits.”

- Plomin, R., Haworth, C. & Davis, O. [Common disorders are quantitative traits](#). *Nat Rev Genet* **10**, 872-878 (2009). doi:10.1038/nrg2670

Biometrics

What are the sources of family resemblance? How do we quantify them numerically?

Heritability

Proportion of similarity in phenotypes that can be attributed to similarity in genotypes.

Model: Phenotype (P) = Genotype (G) + Environment (E)

Variance decomposition

$$\text{var}(P) = \text{var}(G) + \text{var}(E)$$

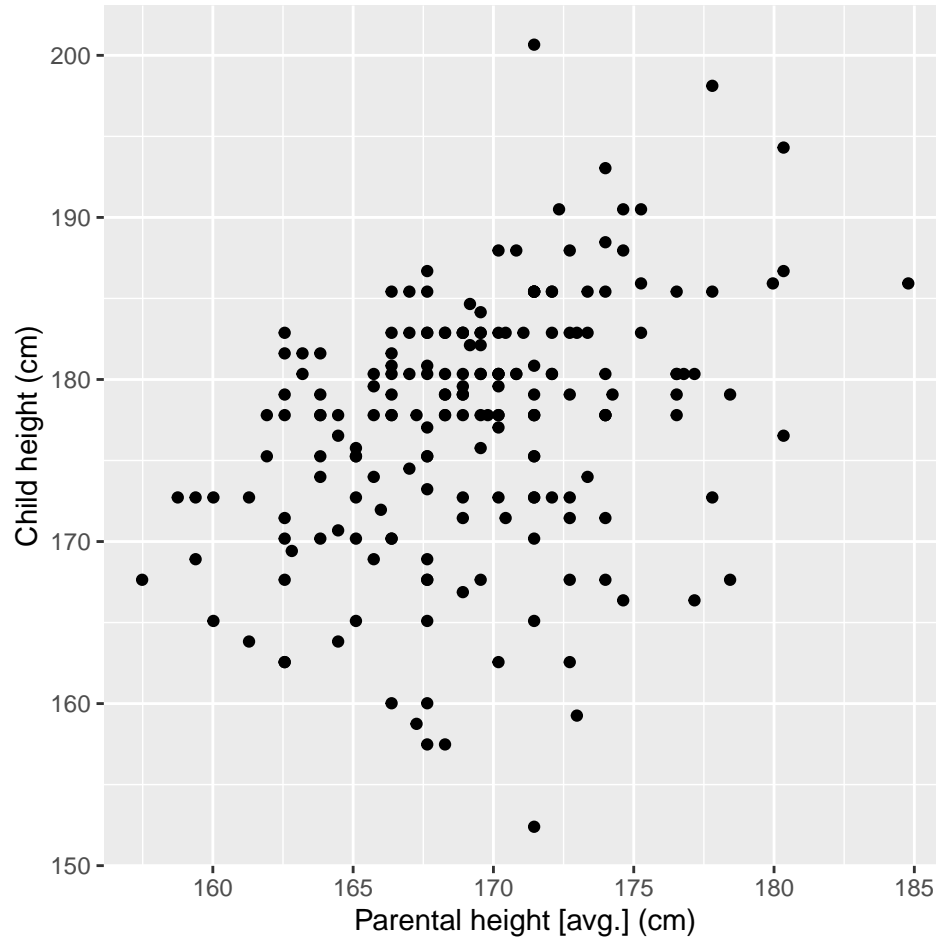
Proportion of variance

$$H^2 = \frac{\text{var}(G)}{\text{var}(P)}, e^2 = \frac{\text{var}(E)}{\text{var}(P)}, H^2 + e^2 = 1$$

The effect G denotes all genetic effects ($G = A + D + I$ for additive, dominance, and epistatic variance). In the more limited case where it is assumed genetic effects act only additively (they don't interact with each other), then only additive genetic variance $\text{var}(A)$ is used with (narrow-sense) heritability defined as $h^2 = \frac{\text{var}(A)}{\text{var}(P)}$.

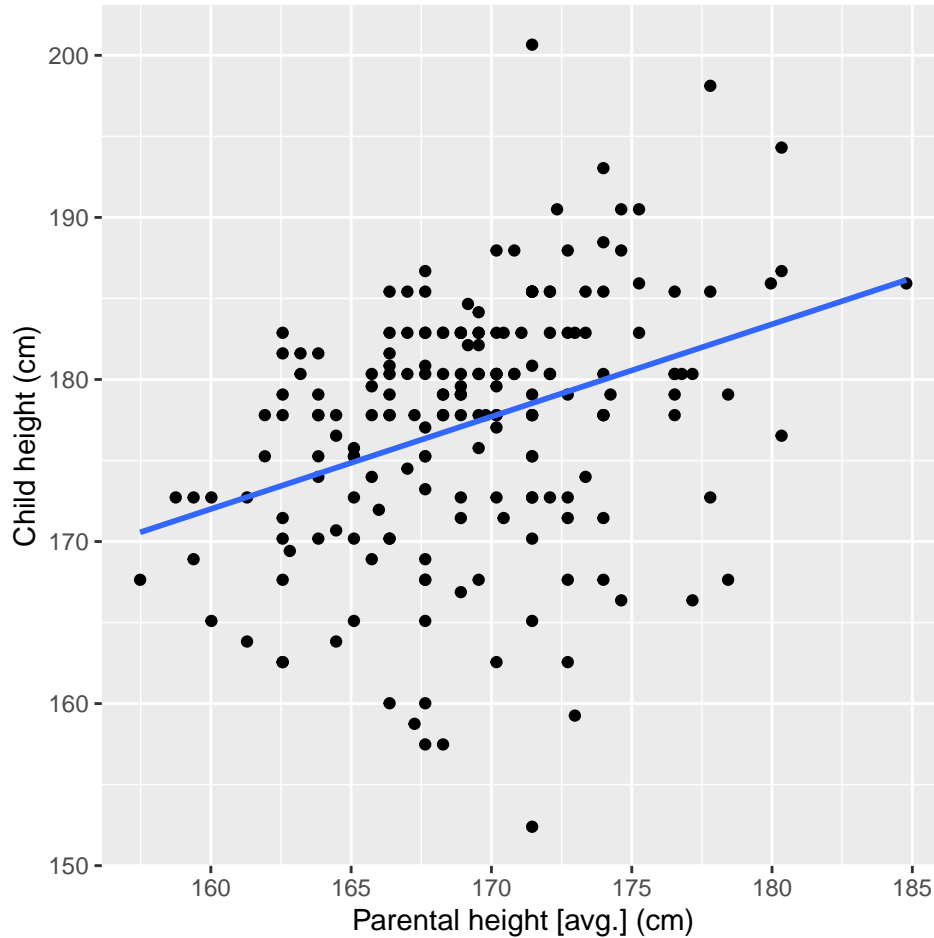
- Tenesa, A., Haley, C. [The heritability of human disease: estimation, uses and abuses](#). *Nat Rev Genet* **14**, 139–149 (2013). doi:10.1038/nrg3377
- Visscher, P., Hill, W. & Wray, N. [Heritability in the genomics era — concepts and misconceptions](#). *Nat Rev Genet* **9**, 255–266 (2008). doi:10.1038/nrg2322

How to estimate heritability from data



Plot of child (offspring) height versus the average of their parents' heights. What is a statistic that can be used to summarise the relationship between these two variables?

How to estimate heritability from data



$$\beta = \frac{\text{cov}(X,Y)}{\text{var}(X)}$$

Estimate the beta coefficient (slope) for a simple regression from the covariance between predictor (X) and outcome (Y) variable divided by the variance of the predictor (X).

Simple model of genetic and environmental effects

$$P = A + E$$

The phenotype value P is influenced by an additive genetic effect A and an environmental effect E .

For simplicity assume that P is an individual's deviation from the average phenotype in the population.

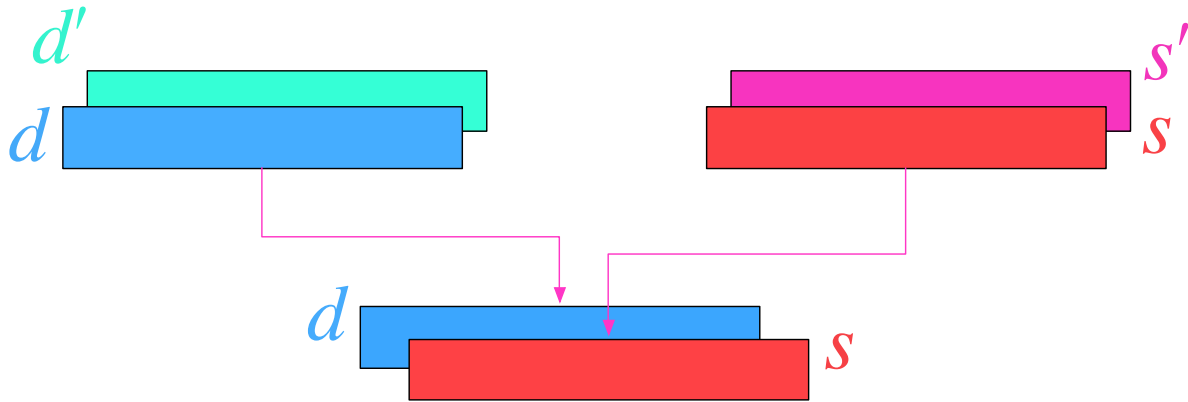
Simple model of genomics

$$A = d + s$$

Each individual has two copies of the genome, one inherited from each parent.

Here we label the genome from the mother d and the genome from the father s . These are the transmitted genetic effects.

Simple model of inheritance



Each parent also has a copy of the genome which they do not pass on to their child, labelled d' for the mother and s' for the father. These are the non-transmitted genetic effects.

Simple model of genetics, environment, and inheritance

Phenotype (P) value is the sum of the two genetic values plus an environmental value (e).

- Mother's phenotype: $P_d = d + d' + e_d$
- Father's phenotype: $P_s = s + s' + e_s$
- Child's phenotype: $P_o = d + s + e_o$

Regression equation

$$\beta = \frac{\text{cov}(X, Y)}{\text{var}(X)}$$

- X = average of parents' phenotypes
- Y = offspring phenotype

$$\text{Therefore, } \beta = \frac{\text{cov}(\frac{P_d + P_s}{2}, P_o)}{\text{var}(\frac{P_d + P_s}{2})}$$

Parent–offspring covariance

$$\begin{aligned}\text{cov}\left(\frac{P_d + P_s}{2}, P_o\right) \\ = \text{cov}\left(\frac{d + d' + e_d + s + s' + e_s}{2}, d + s + e_o\right)\end{aligned}$$

Parent-offspring covariance

Expand the terms. Recall that:

$$\text{cov}(A + X, B + Y) = \text{cov}(A, B) + \text{cov}(A, Y) + \text{cov}(X, B) + \text{cov}(X, Y)$$

Thus we can do a pairwise expansion to:

$$\begin{aligned}&= \text{cov}\left(\frac{d}{2} + \frac{d'}{2} + \frac{e_d}{2} + \frac{s}{2} + \frac{s'}{2} + \frac{e_s}{2}, d + s + e_o\right) \\ &= \text{cov}\left(\frac{d}{2}, d\right) + \text{cov}\left(\frac{d'}{2}, d\right) + \dots + \text{cov}\left(\frac{e_s}{2}, e_o\right)\end{aligned}$$

Simplifications

Some terms can be simplified.

Covariance between a genetic effect and itself

$$\text{cov}\left(\frac{d}{2}, d\right), \text{cov}\left(\frac{s}{2}, s\right)$$

Simplifies to:

$$\begin{aligned}\text{cov}\left(\frac{d}{2}, d\right) &= \frac{1}{2}\text{cov}(d, d) = \frac{1}{2}\text{var}(d) \\ \text{cov}\left(\frac{s}{2}, s\right) &= \frac{1}{2}\text{cov}(s, s) = \frac{1}{2}\text{var}(s)\end{aligned}$$

Assumptions

For some terms we might make an assumption that they are equal to 0.

Covariance between genetic effects from the same parent

$$\text{cov}(\frac{d'}{2}, d), \text{cov}(\frac{s'}{2}, s)$$

If there is inbreeding (individual does not have ancestors that are closely related) or assortative mating in the grandparental generation, then genetic effects from the same parent would be expected to have a non-zero covariance.

Covariance between genetic effects from different parents

$$\text{cov}(\frac{d'}{2}, s), \text{cov}(\frac{s'}{2}, d)$$

If there is assortative mating between the parents, then these effects would be expected to covary

Covariance between parent and offspring environment effects

$$\text{cov}(\frac{e_d}{2}, e_o), \text{cov}(\frac{e_s}{2}, e_o)$$

If aspects of the parental environment are transmitted as well or if parents and offspring tend to encounter similar environments, then these effects are expected to covary.

Covariance between parental genetic and offspring environmental effects

$$\text{cov}(\frac{d}{2}, e_o), \text{cov}(\frac{s}{2}, e_o)$$

If parental genetic effects shape the environment of the offspring (referred to as indirect genetic effects), then these effects are expected to covary as well.

Using those assumptions the parent-offspring covariance simplifies to

$$\text{cov}(\frac{P_d + P_s}{2}, P_o) = \frac{\text{var}(d) + \text{var}(s)}{2}$$

Parent variance

The denominator in the regression equation was

$$\text{var}\left(\frac{P_d + P_s}{2}\right)$$

Using the identity

$$\text{var}(aX + bY) = a^2\text{var}(X) + b^2\text{var}(Y) + 2ab\text{cov}(X, Y)$$

the variance of the average parental phenotypes is:

$$\begin{aligned}\text{var}\left(\frac{P_d + P_s}{2}\right) &= \text{var}\left(\frac{1}{2}P_d + \frac{1}{2}P_s\right) \\ &= \left(\frac{1}{2}\right)^2 \text{var}(P_d) + \left(\frac{1}{2}\right)^2 \text{var}(P_s) + 2 \cdot \frac{1}{2} \cdot \frac{1}{2} \text{cov}(P_d, P_s) \\ &= \frac{1}{4}\text{var}(P_d) + \frac{1}{4}\text{var}(P_s) + \frac{1}{2}\text{cov}(P_d, P_s)\end{aligned}$$

If we assume as above that there is no covariation between parental effects ($\text{cov}(P_d, P_s) = 0$), this simplifies to

$$= \frac{\text{var}(P_d) + \text{var}(P_s)}{4}$$

Thus the regression equation is:

$$\beta = \frac{\text{cov}\left(\frac{P_d + P_s}{2}, P_o\right)}{\text{var}\left(\frac{P_d + P_s}{2}\right)} = \frac{\frac{\text{var}(d) + \text{var}(s)}{2}}{\frac{\text{var}(P_d) + \text{var}(P_s)}{4}} = 2 \frac{\text{var}(d) + \text{var}(s)}{\text{var}(P_d) + \text{var}(P_s)}$$

Previously we defined

$$A = d + s$$

thus

$$\text{var}(A) = \text{var}(d) + \text{var}(s)$$

and assume variances in parental phenotypes are equal

$$\text{var}(P_d) = \text{var}(P_s) = \text{var}(P)$$

Then substitute into the regression equation

$$\beta = 2 \frac{\text{var}(d) + \text{var}(s)}{\text{var}(P_d) + \text{var}(P_s)} = 2 \frac{\text{var}(A)}{\text{var}(P) + \text{var}(P)} = 2 \frac{\text{var}(A)}{2\text{var}(P)} = \frac{\text{var}(A)}{\text{var}(P)} = h^2$$

In other words, the beta coefficient from the regression of offspring phenotype on average parent phenotype yields an estimate of the heritability!

Height data

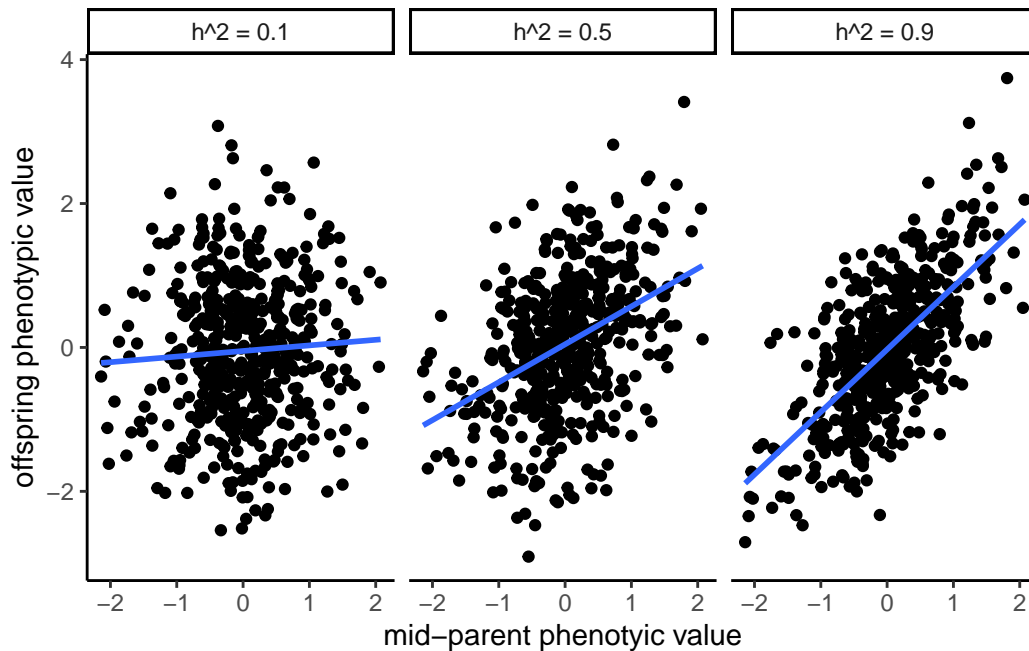
Using the height data, we can calculate the mid-parent–offspring covariance and mid-parent variance and then use that as an estimate for the heritability.

$$\text{cov}(\frac{P_d + P_s}{2}, P_o) = 12.57$$

$$\text{var}(\frac{P_d + P_s}{2}) = 22.04$$

$$\hat{h}^2 = 12.57 / 22.04 = 0.57$$

Parent and offspring phenotypes become more highly correlated as heritability increases.



Wray, N. & Visscher, P. (2008) [Estimating trait heritability](#). Nature Education 1(1):29

Mini review: What assumptions have we made when estimating h^2 ?

Parents' environments are not similar: $\text{cov}(e_d, e_s) = 0$

No assortative mating: $\text{cov}(d, s) = 0$

Parents do not transmit their environments: $\text{cov}(e_o, e_d) = 0$, $\text{cov}(e_o, e_s) = 0$

No gene-environment correlation: $\text{cov}(d, e_d) = 0$, $\text{cov}(s, e_s) = 0$

No genetic nature: $\text{cov}(d, e_o) = 0$, $\text{cov}(s, e_o) = 0$

No inbreeding: $\text{cov}(d, d') = 0$, $\text{cov}(s, s') = 0$

Genetic effects are additive: $Y = a + a' + e$

Genetic influence is the same for both sexes: $\text{var}(d + d') = \text{var}(s + s')$

Generalising to other relatives

Heritability can also be estimated from resemblance between different types of related pairs.

The general equation is:

$$h^2 = \frac{b}{r}$$

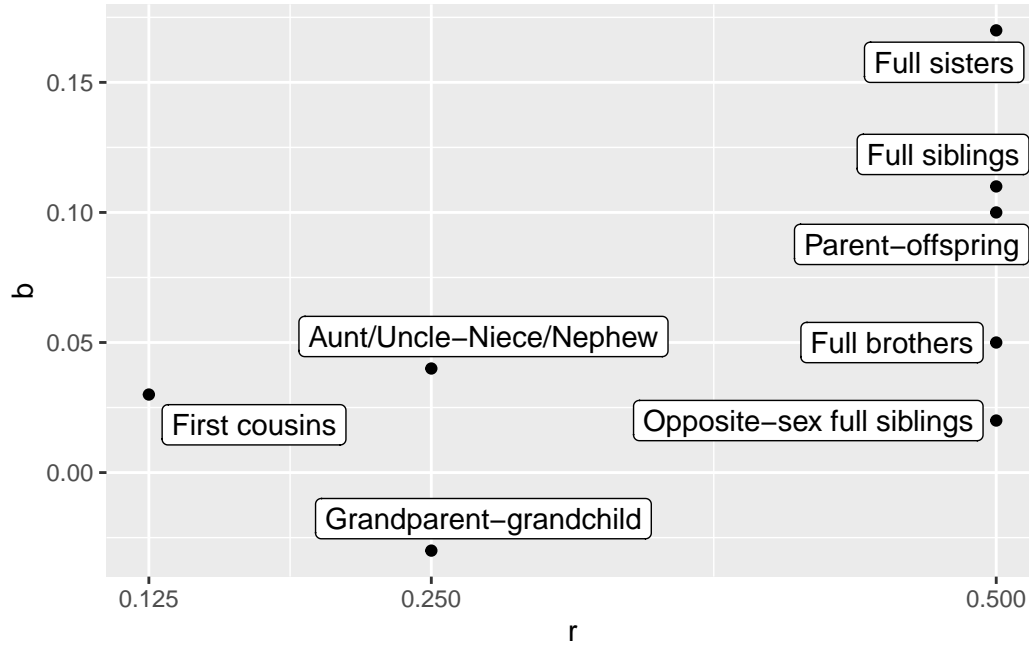
b = regression coefficient

r = relatedness coefficient (“coefficient of additive variance”)

In this notation we used normal r to represent relatedness, to distinguish it from italic r for correlation coefficient.

Example data: depression scores

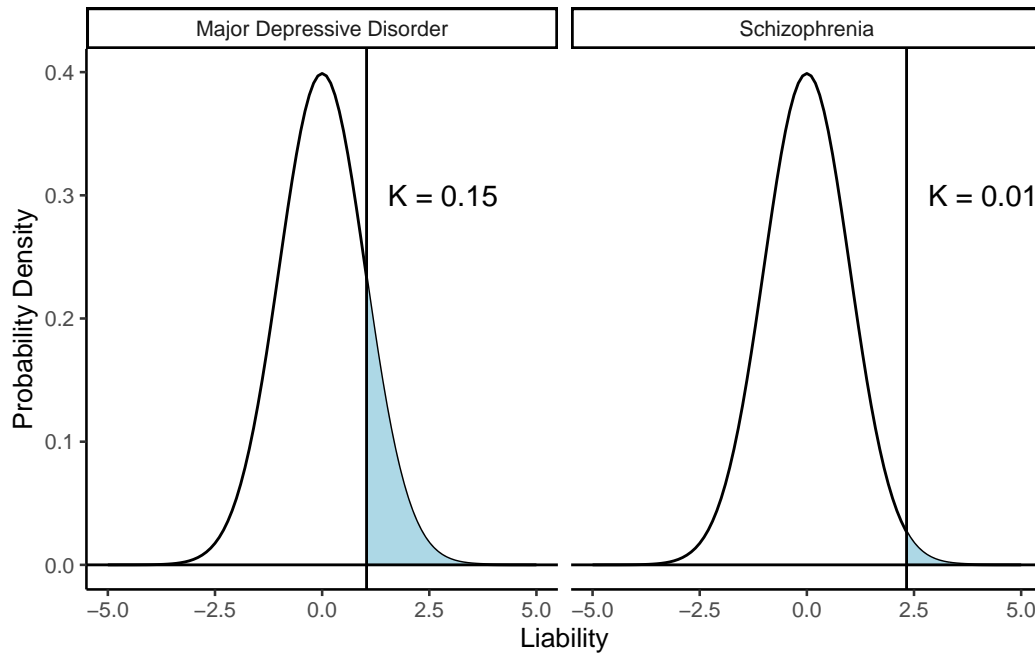
Correlation of depression scores for different pairs of relatives



Fernandez-Pujals AM et al. (2015) Epidemiology and Heritability of Major Depressive Disorder, Stratified by Age of Onset, Sex, and Illness Course in Generation Scotland: Scottish Family Health Study (GS:SFHS). *PLOS ONE* 10(11): e0142197. doi:[10.1371/journal.pone.0142197](https://doi.org/10.1371/journal.pone.0142197)

Recurrence risk to relatives

$$\lambda_R = \frac{P(\text{affected}|\text{relative affected})}{P(\text{affected in population})} = \frac{K_R}{K}$$

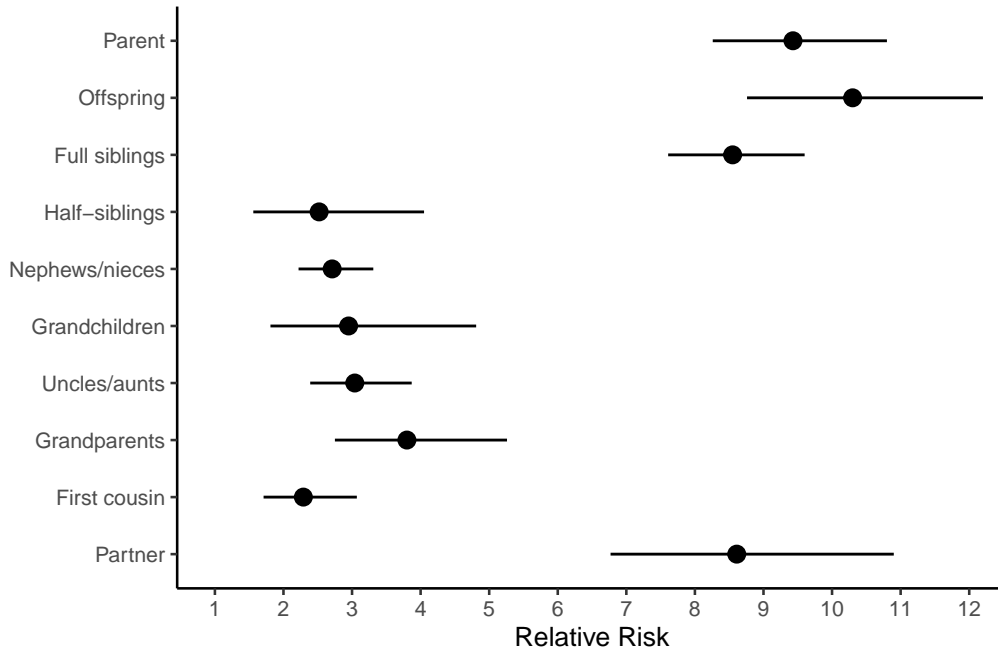


Represents how much more likely to are to be affected by a disorder given that a relative is affected, compared to someone from the general population.

Example:

- $K_{\text{sib}} = P(\text{affected}|\text{sibling affected}) = 0.09$
- $K = P(\text{affected in population}) = 0.02$
- $\frac{K_{\text{sib}}}{K} = \frac{0.09}{0.02} = 4.5$

Recurrence risk for schizophrenia



Recurrence risk to relatives for schizophrenia in Sweden, which has a baseline risk of $K = 0.47\%$.

Lichtenstein, P. et al. Recurrence risks for schizophrenia in a Swedish National Cohort. *Psychol. Med.* 36, 1417–1425 (2006). doi:[10.1017/s0033291706008385](https://doi.org/10.1017/s0033291706008385)

Recurrence risk and heritability

- Score unaffected = 0, affected = 1
- If population prevalence is K , then phenotypic variance is $K(1 - K)$ (Bernoulli distribution)

- Y = score of individual (proband)
- Y_R = score of relative of proband
- Expectation: $E[Y] = E[Y_R] = K$
- $K_R = E[Y_R | Y = 1]$
- Probability that both Y and $Y_R = 1$: $E[YY_R] = K \times K_R$

$$\text{cov}(Y, Y_R) = E[YY_R] - E[Y]E[Y_R] = K \times K_R - K^2$$

- James, J. W. Frequency in relatives for an all-or-none trait. *Ann. Hum. Genet.* 35, 47–49 (1971). doi:[10.1111/j.1469-1809.1956.tb01377.x](https://doi.org/10.1111/j.1469-1809.1956.tb01377.x)
- Risch N. Linkage strategies for genetically complex traits. I. Multilocus models. *Am J Hum Genet.* 1990 Feb;46(2):222-8. PMID: 2301392

$$= K \times K_R - K^2 = K(K_R - K) = K^2 \left(\frac{K_R}{K} - 1 \right) = K^2(\lambda_R - 1)$$

Heritability estimate

$$h^2 = \frac{\text{cov}_R}{rV_P} = \frac{K^2(\lambda_R - 1)}{rK(1 - K)} = \frac{K(\lambda_R - 1)}{r(1 - K)}$$

Recurrence risk of psychiatric disorders

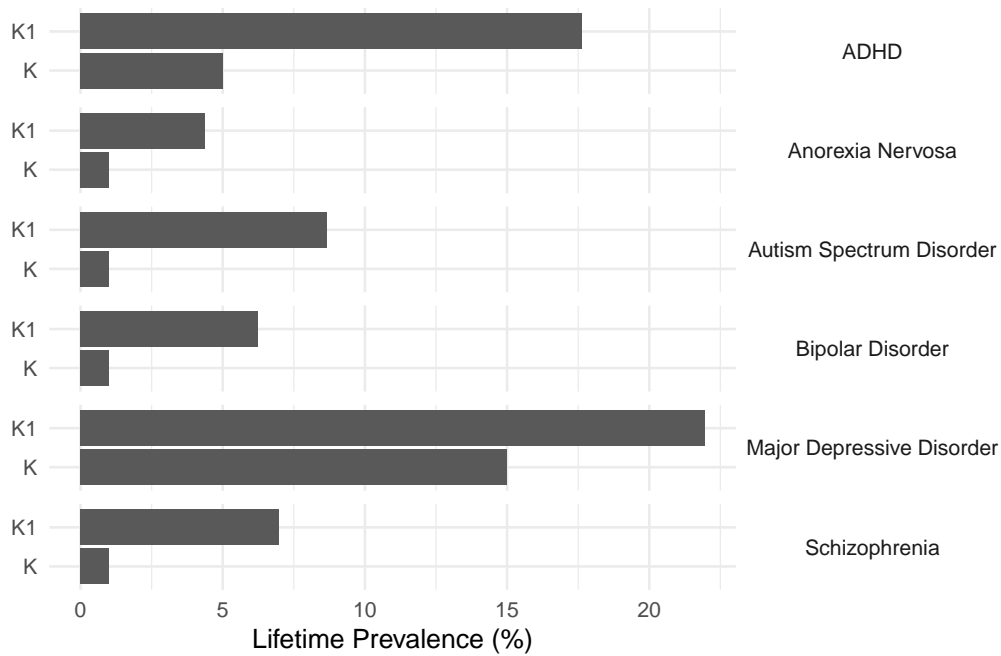
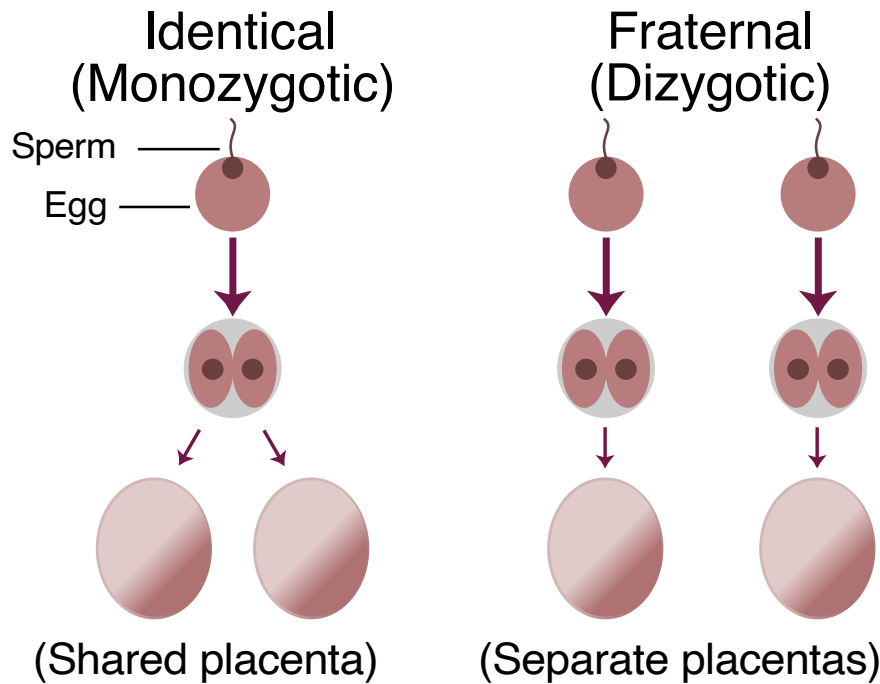


Figure 2: Prevalence in population (K) and with one parent affected (K[1])

Baselmans, B. M. L., Yengo, L., Rhee, W. van & Wray, N. R. Risk in relatives, heritability, SNP-based heritability and genetic correlations in psychiatric disorders: a review. *Biol Psychiatry* (2020) doi:10.1016/j.biopsych.2020.05.034.

Estimating environmental effects

Contrast pairs of relatives that have comparable environmental similarity but different genetic similarity.



- Monozygotic (MZ) twins $r = 1.0$
- Dizygotic (DZ) twins $r = 0.5$

[Zygote development figure](#) CC-BY-SA Triply.

Additive genetic and shared environment effects

Add a shared (C or “common”) environment to the basic genetic model, to capture similarity between relatives attributable to environmental factors. E represents the unique, non-shared environment.

$$P = A + C + E$$

$$h^2 = \frac{\text{var}(A)}{\text{var}(P)}, c^2 = \frac{\text{var}(C)}{\text{var}(P)}, e^2 = \frac{\text{var}(E)}{\text{var}(P)}$$

$$h^2 + c^2 + e^2 = 1$$

Twin correlations

$$\text{MZ twins: } r_{\text{MZ}} = h^2 + c^2$$

$$\text{DZ twins: } r_{\text{DZ}} = \frac{1}{2}h^2 + c^2$$

Solve for genetic similarity (h^2)

Calculate difference between MZ and DZ correlations

$$r_{\text{MZ}} - r_{\text{DZ}} = (h^2 + c^2) - (\frac{1}{2}h^2 + c^2)$$

$$r_{\text{MZ}} - r_{\text{DZ}} = h^2 - \frac{1}{2}h^2 + c^2 - c^2$$

$$r_{\text{MZ}} - r_{\text{DZ}} = \frac{1}{2}h^2$$

$$h^2 = 2(r_{\text{MZ}} - r_{\text{DZ}})$$

Substitute h^2 into MZ equation and solve for shared environment similarity (c^2)

$$r_{\text{MZ}} = \underbrace{h^2}_{2(r_{\text{MZ}} - r_{\text{DZ}})} + c^2$$

$$r_{\text{MZ}} = 2(r_{\text{MZ}} - r_{\text{DZ}}) + c^2$$

$$r_{\text{MZ}} - 2(r_{\text{MZ}} - r_{\text{DZ}}) = c^2$$

$$c^2 = r_{\text{MZ}} - 2r_{\text{MZ}} + 2r_{\text{DZ}}$$

$$c^2 = 2r_{\text{DZ}} - r_{\text{MZ}}$$

Therefore from MZ and DZ twin correlations we can estimate:

$$h^2 = 2(r_{\text{MZ}} - r_{\text{DZ}})c^2 = 2r_{\text{DZ}} - r_{\text{MZ}}e^2 = 1 - h^2 - c^2$$

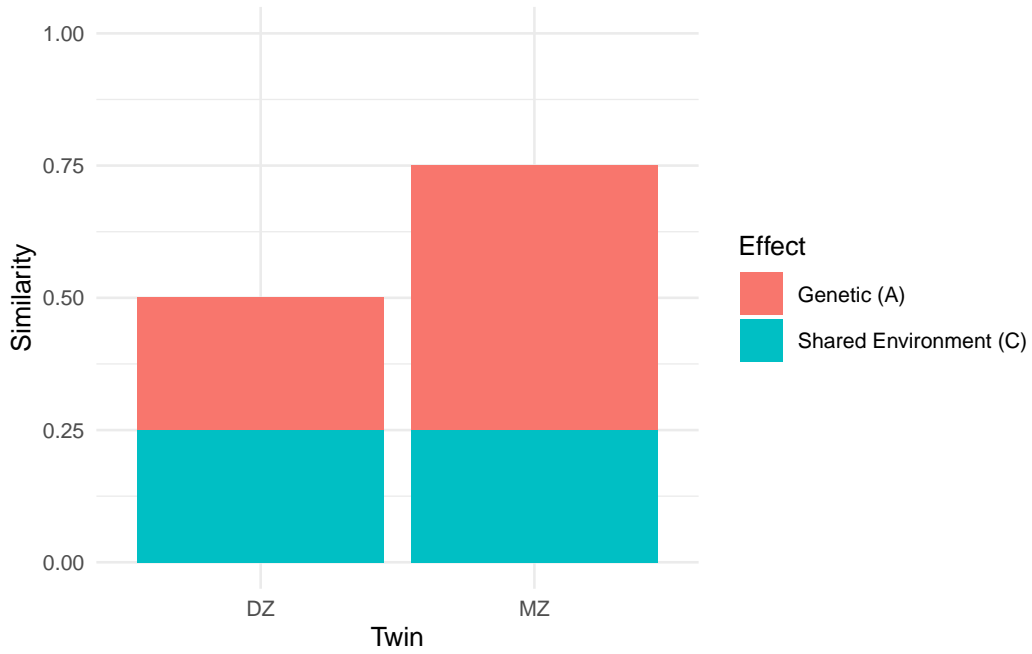
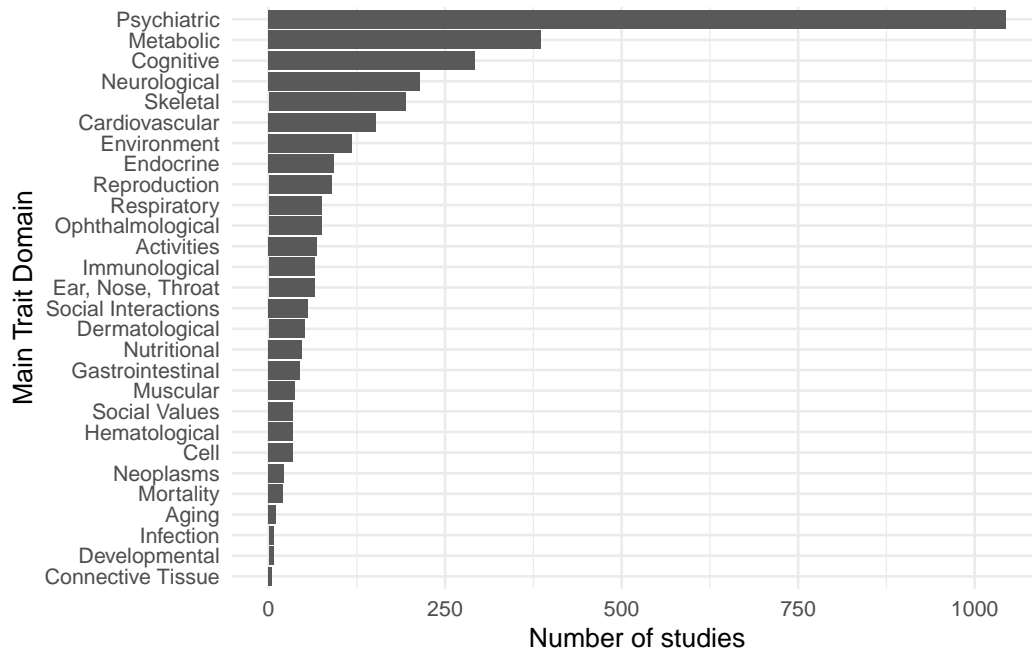


Figure 3: Visualisation with $r[\text{MZ}] = 0.75$ and $r[\text{DZ}] = 0.5$.

What do we know about psychiatric genetics from twins studies



Polderman TJC et al. [Meta-Analysis of the Heritability of Human Traits based on Fifty Years of Twin Studies](#) *Nature Genetics* doi:10.1038/ng.3285

Meta-analysis of twin heritability

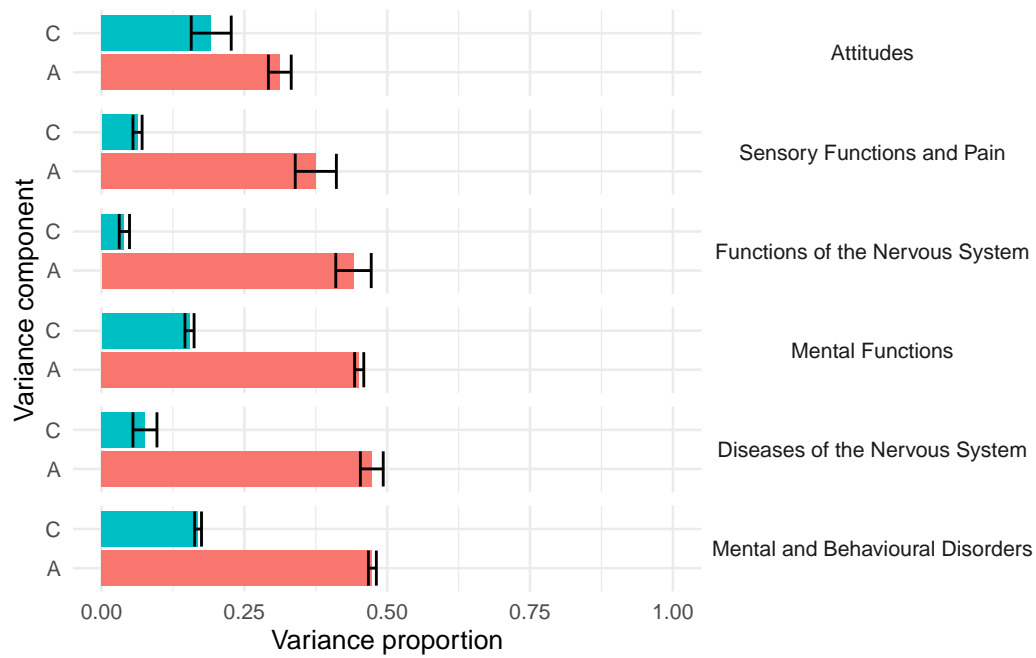


Figure 4: Meta-analysis of phenotype domains

Mental and behavioural disorders are on average moderately heritable with a smaller but substantial portion attributable to the shared environment.

Data from match.ctglab.nl.

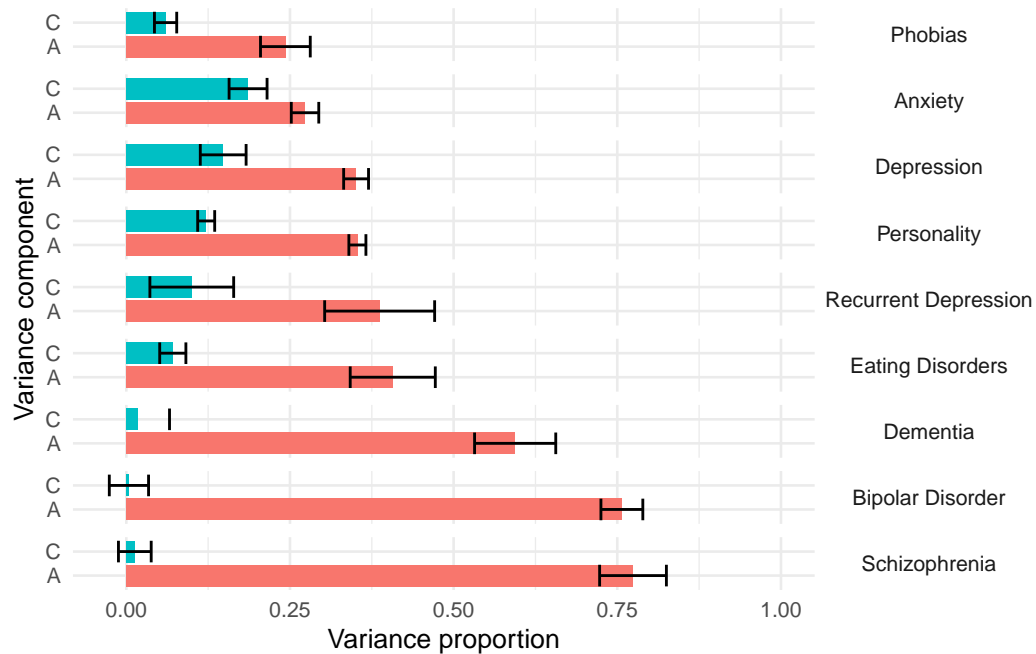


Figure 5: Meta-analysis of psychiatric, neurological, and psychological phenotypes

Genetics of depression and schizophrenia

Major depressive disorder (MDD)

- 1+ in 10 affected
- 7-11 years of life are lost
- Global cause of ill-health

MDD diagnostic criteria

1 of the 2 **cardinal** symptoms with 5 symptoms in total:

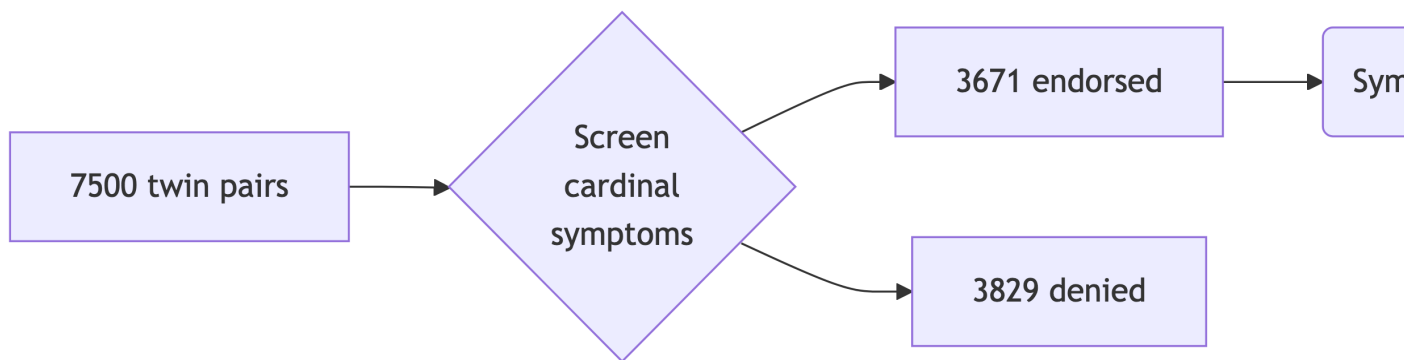
1. **Low mood**
2. **Anhedonia**
3. Increase or decrease in weight or appetite
4. Insomnia or hypersomnia
5. Psychomotor agitation or slowing
6. Fatigue

7. Feelings of worthlessness/guilt
8. Concentration problems
9. Suicidal ideation

Heterogeneous disorder: 227 possible symptom profiles.

Fried, E. I. & Nesse, R. M. Depression is not a consistent syndrome: An investigation of unique symptom patterns in the STAR*D study. *J Affect Disorders* 172, 96–102 (2015).

“Evidence for Multiple Genetic Factors Underlying DSM-IV Criteria for Major Depression”



Kendler, K. S., Aggen, S. H. & Neale, M. C. Evidence for Multiple Genetic Factors Underlying DSM-IV Criteria for Major Depression. *JAMA Psychiat* 70, 599–607 (2013).

Multivariate twin model

- analysed MZ and DZ twin cross-correlations among depression symptoms
- tested 1, 2, or 3 factor models of genetic, shared environment, and unique environment

Best fit model

Genetic factors

1. Psychomotor/cognitive symptoms (psychomotor, guilt, concentration, suicidality)
2. Mood symptoms (low mood, anhedonia)
3. Neurovegetative symptoms (weight, sleep, fatigue)

Individual-specific (unique) environment factors

1. General depression (low mood, anhedonia, weight, sleep, psychomotor, fatigue, concentration)
2. Mood symptoms (low mood, anhedonia, concentration)
3. Cognitive symptoms (guilt, suicidality)

External validators of genetic factor scores

Table 1: Data from Table 4 Kendler, Aggen, & Neale. $p < 0.001$

Validator variable	Cognitive/Motor	Core Mood	Neurovegetative
Neuroticism	0.54	0.27	0.30
Anxiety disorder	0.77	0.49	0.47
Age at onset	-3.00	-1.15	0.37
No. of episodes	0.55	0.16	0.23
Melancholia subtype	1.05	-0.43	1.34
Unreactive mood	0.82	0.10	0.79

Cognitive/Motor factor associated with non-specific liability to internalising disorders

Neurovegetative factor most strongly related to Melancholic subtype more specific to major depression

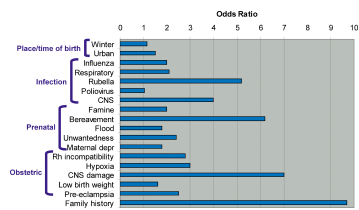
Schizophrenia diagnostic criteria

At least one **core** symptom and two or more symptoms present for a significant period of time

1. **Delusions**
2. **Hallucinations**
3. **Disorganised speech**
4. Grossly disorganised or catatonic behaviour
5. Negative symptoms

Negative symptoms encompass deficiencies in emotional life, motivations, thinking, and relationships

Schizophrenia isk factors



Compare environmental risk factors to family history.

Figure CC-BY Sullivan PF (2005) The Genetics of Schizophrenia. PLoS Med 2(7): e212.
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