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1 Evolutionary Thinking

Introduction (Lecture)

- Essential questions of evolutionary biology:
 - Why do organisms look so different?
 - Why develop elaborate sexual traits?
 - Why do organisms senesce?
- ▶ Evolution is mainly an historical science and thus must relay on other methods of reconstructing the past or making inferences about evolutionary forces.
- ▶ **Proximate**: a question about a mechanistic cause; provides an immediate explanation about how a mechanistic cause functions.
- ▶ **Ultimate**: why, or the reason, a trait or organism is the way it is; an evolutionary explanation.
- Example of proximate vs ultimate in Galapagos finches:
 - Proxmate: developmental growth factor is increased/decreased in some birds.
 - Ultimate: different habits are selected on breaks that maxmize food gathering ability.
- ▷ Evolutionary biology's approach to answering questions:
 - **Empirical data**: observation studies, experiments; the *comparative method*.
 - **Theory**: predictions that use models and mathematical reasoning which can be be tested with empirical data.
- Described Discribing Overview of the components of evolution by natural selection:
 - o Genetic variation exists, via mutations.
 - Mutations are heritable.
 - The is an advantage to survival and/or reproduction from the mutation.
 - Individuals with the advantage in survival/reproduction are selected for.

10 Studying Adaptation

Hypothesis Testing: Oxpeckers Reconsidered

- ▶ **Adaption**: a trait, or a suite of traits, that increases the fitness of its possessor.
- No hypothesis for the adaptive value of a trait should be accepted simply because of its plausibility.
- Dependent of the provided a safe of the provided as a safe of the pr
- Experiments on cattle were done to test whether this observation was true:
 - Results show red-billed oxpeckers have no effect tick loads of cattle.
 - Red-billeld oxpeckers maintained open wounds, even enlarging existing wounds to feed on the cattle's blood.
 - Red-billed oxpeckers removed hosts' earwax; whether this is good of bad is unclear.
 - Even these results must remain in question, as cattle are not the native host for the birds.
- Do Other important points to remember:
 - Differences among populations or species are not always adaptive.
 - Not every trait is adaptive.
 - Not every adaptation is perfect, often the adaption just happened to work well enough or by chance better than other adaptations.

Experimental Design

- ▶ Defining and testing effective control groups is critical.
- ▶ Treatments of controls and experimental measures must be handled as close to exactly alike as possible.
- ▶ Randomization is a key technique for equalizing miscellaneous effects and a tool to avoid bias.
- ▶ Reproduction is essential in order to help remove potential outlier effects.
 - Allows for greater understanding of precision, accuracy, and variation by providing more data for statistical tests.

4 Evolutionary Trees

How to Read an Evolutionary Tree

- ▶ **Phylogeny**: aka evolutionary tree or phylogenetic tree, is a diaggram showing the history of divergence and evolutionary change. Essentially, it's the genealogical relationships of organisms based on descent with modification.
 - **Taxa**: the units you are analyzing, e.g. certain species or DNA sequences.
 - Character: a feature or trait present among the taxa of interest, e.g. teeth of mammals or nucleotides of DNA sequences.
 - · Character state(s): an alternative condition of a character, which are able to evolve one to another, e.g. pointed/flat teeth of mammals.
 - Ancestral character: a trait that was possessed by the common ancestor.
 - Derived character: a trait the was not possessed by the common ancestor and instead evolved in at least one of the descendants.
 - **Synapomorphy**: derived character state shared by two or more taxa and used to define a clade of taxa.
 - · **Autapomorphy**: derived character state in only one taxon.
 - Outgroup: a taxon or taxa that are used to root the phylogeny or determine ancestral character states.
 - Ingroup: the set of taxa that are the focus of the phylogeny.
 - Nodes: points at which the tree splits; represents mutations, speciation events, or character changes.
 - Anagensis: descent with modification, but no speciation.
 - **Cladogenesis**: speciation, origin of clades.
 - Clade: also known as a monophyletic group, an ancestor and all of its descendants.
 - Paraphyletic group: a group of organisms consisting of an ancestor and some of its descendants.
 - Sister: a taxa or clade that are most closely related to each other; they
 share the most recent common ancestor.
- ▶ **Homology**: similarity due to common descent; continuity of a trait, character, or character state through time.
 - Homologous trait: found in a taxa that inherited the trait from a common ancestor.

- ▶ **Homoplasy**: or analogous, similarity in the characters or traits in different taxa due to convergent evolution, parallelism, or reversal, but not due to common descent.
 - **Convergent evolution**: similar traits due to selective forces and not shared ancestry.
 - **Parallelism**: convergent evolution in recently diverged taxa.
 - Reversal: derived traits or character states that revert to the ancestral form

Inferring Phylogenetic Trees

- ▶ **Parsimony**: relationships that require the smallest number of character changes are most likely to be correct.
 - Based on shared and derived traits(synapomorphies).
 - Reconstruction using parsimony:
 - 1. Code characters.
 - 2. Make up a taxon×character matrix.
 - 3. Search for synapomorphies, and the shortest tree.
 - Outgroups can help polarize (ancestral vs derived) the characters.
 - Treelength: a measure of evolutionary change using parsimony.
 - Shortest tree length produces most parsimonious tree.
 - Length determined by number of synapomorphies.
 - Homoplasious characters increase tree length.
- ▶ Distance Methods: converts a sequence alignment to genetic distances between pairs of sequences.
 - Branch length is proportional to genetic differences.
- ▶ **Maximum likelihood**: a method of estimating the parameters of a probability distribution by maximizing a *likelihood function*.
 - One of the more dominant means of statistical inference.
 - Likelihood: measure of goodness of fit of a statistical model to a sample of data for given values of the unknown parameters.
 - \circ P(D|H); probability(P), Data(D), Hypothesis(H)
 - **Bayesian**: uses the likelihood function to create a quantity called the *posterior probability* of trees using a model of evolution based on prior probabilities in order to produce the most likely tree.

 Bootstraping: creating a value that indicates how many times out of 100 (normally) that the same branch was observed when repeating the phylogenetic reconstruction on re-sampled (pseudoreplicated) set of dat.

6 Mechanisms of Evolutionary Change

Hardy-Weinberg Equilibrium

- ▶ **Population**: a group of interbreeding individuals and their offsring.
- ▶ **Gene pool**: the set of all genes, or genetic information, in any population.
- ▶ **Genotypic frequency**: number of individuals with a given genotype divided by the total number of individuals in the population.
 - The proportion (i.e., 0 < f < 1) of genotypes in a population.
- ▶ **Allele frequencies**: relative frequency of an allele at a particular locus in a population.
 - **Locus**: a fixed position on a chromosome where a particular gene of genetic marker is.
 - Monoploids: frequency of an allele is the result of the number of copies of the allele divided by sample size.
 - p = i/N
 - p: frequency | i: copies of alleles | N: sample size
 - Diploids: frequency of alleles within three possbile genotypes at a locus with two alleles.
 - $-p = f(AA) + \frac{1}{2}f(AB)$ frequency of A-allele
 - $-q = f(BB) + \frac{1}{2}f(AB)$ frequency of B-allele
 - Allele frequency can always be calculated from genotype frequency, whereas the reverse requires the *Hardy-Weinberg principle* of random mating apply.
- ▶ **Hardy-Weinberg principle**: allele and genotype frequencies in a population will remain constant in the absence of evolutionary influences.
 - Allele frequencies do not change from one generation to the next.
 - Genotypic frequencies after one generation of random mating: $p^2 + 2pq + q^2$
 - Evolutionary influences: genetic drift, mate choice, assortative mating, natural selection, sexual selection, mutation, gene flow, meiotic drive, genetic hitchhiking, population bottleneck, founder effect, and inbreeding.
 - Most of these influences will be discussed later.

Selection

- > Fitness: success at which a organism produces fertile offspring.
- ▶ Competition: an interaction between organism in which the fitness of one is lowered by the presence of another.
- > **Selection**: the act on a heritable phenotypic trait due to competition.
 - o Can be members of the same of different species.
 - Not always directional and adaptive, instead selection pressure is applied and removes the less fit variants.
 - Can be classified in different ways, such as effect on a trait, on genetic diversity, by life cycle, by unit of selection, or by the resource in competition.
 - Most effective on large populations.

By Effect on a Trait

- **Stabilizing selection**: the simplies case in which selection acts to hold a trait at a stable optimum.
 - Reduces the individuals in the trails of the trait's distribution, reducing variation.
- **Directional selection**: favours extreme values of a trait.
 - Directional selection on a continuous trait changes the average value of the trait in the population.
 - Can reduce variation in the population, generally not by large amounts though.
- **Disruptive selection (diversifying selection)**: acts during transition periods when current mode is sub-optimal, but alters trait in more than one direction.
 - Univariate: when the trait is both quantitatively favoured in either direction and can lead to speciation.
 - Generally increases the variance on continuous traits.
 - May be more common than generally recognized.
- All three increases the mean fitness of the population.

By Effect on Genetic Diversity

- Purifying selection: aka negative selection; acts to remove genetic variation from the population.
- **de novo** mutation: introduces new variation and opposes negative selection.
- Balancing selection: acts to maintain genetic variation, even in absence of *de novo* mutation by frequency-dependent selection.
 - Frequency-dependent selection: fitness that depends of the phenotypic or genotypic composition of a population.
 - · Positive: fitness increases as frequency of the trait increases.
 - Negative: fitness decreases as the frequency of the trait increases.
 - Overdominance, aka heterozygote advantage: when a combination of alleles confers a selective advantage over individuals with one allele.
 - Underdominance, aka heterozygote disadvantage: when the heterozygote has lower fitness than either homozygote.

By Life Cycle Stage

- Viability selection: aka survival selection: increases probability of survival.
 - Can act to improve probability of survival before and after reproduction.
- Fecundity selection: increases the rate of reproduction given survival.
 - May be split into sub-components including sexual selection, gametic selection, gamete viability, compatability selection, and zygote formation.

Mutation

- ▶ **Mutation**: alteration in the nucleotide sequence of the genome of an organism.
 - May not produce discernible phenotypic changes.
 - The ultimate source of genetic variation.
 - Have several types of changes, from no effect, to small changes, or complete loss of function.

Large-Scale Structural Mutations

- Gene duplications, aka amplifications: repetition of a chromosomal segment or attachment of extra piece of chromosome to another, leading to multiple copies of chromosomal regions.
- o Deletions of large chromosmal regions.
- Fusion genes: mutations that join previously separated genes into one new distinct gene.
- Chromosmal rearrangement: large scale changes in structure of chromosomes, leading to speciation in isolated, inbred populations. Includes:
 - Chromosomal translocations: interchange of genetic parts from nonhomologous chromosomes.
 - Chromosomal inversions: reversing the orientation of a chromosomal segment.
 - Non-homologous chromosomal crossover.
 - Interstitial deletions: inverse of fusion genes; removes a segment of DNA joining distant genes.
- Loss of heterozygosity: loss of one allele, by deletion or genetic recombination, in a organism that previously had two different alleles.

Small-Scale Mutations

- Point mutation: a single nucleotide base change, that can result in a variety of effects.
- o Insertions: add one or more extra nucleotides into the DNA.
 - Usually caused by transposable elements, or errors during replication or repeating elements.
 - Can causereading frame shift, possibly effecting how many codons are read, and thus altering the gene product.
- **Deletions**: remove one or more nucleotides from the DNA.
 - Also can cause a reading frame shift like insertions.
 - Generally irreversible.
- **Substitutions**: exchange of a single nucleotide for another.
 - Often classified as transitions or transversions.
 - Generally a purine (A-G) for a purine, or a pyrimidine (C-T) for a pyrimidine.
 - Can be reversed by another point mutation.

Impact on Protein Sequence

- Effect of mutation depends heavily on where it occurs, particularly in a coding or non-coding region.
- Regulator sequences, e.g. promoters, enhancers, silencers, can alter gene expression but are less likely to alter protein sequence.
- Frameshift mutation: caused by insertion or deletion of nucleotides that is not divisible by three, resulting in a different translation from the original.
- **Synonymous substitution**: a condon replacement with another that codes for same amino acid.
 - Silent substitution: no phenotypic difference after a synonymous substitution.
- Nonsynonymous substitution: a codon replacement that codes for a different amino acid.
 - Missense mutation: codon replacement that renders the resulting protein nonfunctional.
 - Nonsense mutation: codon replacement that results in a premature stop codon that produces a truncated and often nunfunctional protein.

Migration

- ▶ **Gene flow**: movement of alleles, or genetic variation, between populations.
 - If the rate of gene flow is high enough, then two populations are considered to have equivalent allele frequencies and thus a single population.
 - o Constrains speciation by combining gene pools of the groups.
 - May result in the addition of novel genetic variants in the gene pool.
- ▷ Gene flow is expected to be lower in species that:
 - have low mobility or dispersal.
 - o occur in fragmanted habits.
 - have long distances 2between populations.
 - have small population sizes.
- ▶ **Allopatric speciation**: when gene flow is blocked by physical barriers that inhibit gene flow.
- ▶ **Sympatric speciation**: result of gene flow that is blocked due to non-physical barriers that inhibit gene flow.

Genetic Drift

- ▶ **Genetic drift**: the change in the allele frequencies in a population due to random sampling.
 - Not influenced by environmental factors.
- ▶ May cause certain gene variants to become fixed or lost by chance.
- ▶ Generally drives populations towards genetic uniformity over time, decreasing heterozygosity.
- ▷ Only mutation or gene flow can introduce new alleles, which acts against genetic drift.
- ▶ **Founder effect**: result of sampling error which has an increased likelyhood on populations with low numbers.
 - By chance certain alleles can be dominant when they otherwise wouldn't be in a new founding population.
 - o Often acts to drastically increase rate of genetic drift.
- ▶ **Genetic bottleneck**: a sharp reduction in the size of population due to environmental events.
 - o Can essentially cause a founder effect, though it's not a new population.

Coalescent Theory

- **Coalescent theory**: how gene variants sampled from a population may have originated from a common ancestor.
 - Assumes no recombination, no natural selection, no gene flow in the simplilest case.
- Aims to look backward in time by merging allels into a single ancestral copy according to a random process in coalescence events.
- Many theoretical genealogies are made in order to compare to observed data in order to test assumptions about demographic history of a population.
 - Used to make inference about population genetic parameters, such as migration, population size, and recombination.
- **Coalescent time**: number of preceding generations where the coalescence took place, not calender time.
 - Estimation of the time can be made multiplied by $2N_e$ with the average time between generations.
 - Time to coalescence for a pair of allels at a locus is dependent on population size.

- Formula:
$$P_c(t) = \left(1 - \frac{1}{2N_e}\right)^{t-1} \left(\frac{1}{2N_e}\right)$$

 Can also be used to model the amount of variation in DNA sequences expected from genetic drift and mutation.

Molecular Evolution

- ▶ **Molecular evolution**: the process of change in the sequence composision of cellular molecules across generations.
- ▶ **Polymorphism**: occurrence of two of more clearly different morphs, or alternative phenotypes, in the population of a species.
 - **Substitution**: when allels become fixed or lost in a population and polymorphism is ended.
 - Substitution rate (k): $k = sN\mu$
 - -s = probability of fixation.
 - $N\mu$ = mutation rate of population.

Recombination

- **Recombination**: the process that results in genetic exchange between chromosomes or chromosomal regions.
 - Can also cause mutations due to misalignment after recombination.
 - Gene repair: a type of recombination that is the product of DNA repair that corrects damage using a homologous template.
 - Often responsbile for homogenizing sequences of duplicate genes over long periods of time, which reduces nucleotide divergence.
- **Genetic hitchhiking**: change in allele frequency not because of natural selection, but due to proximity to a gene undering selective sweep.
 - Selective sweep: a beneficial mutation that increases frequency and generally becomes fixed.

Neutral Theory

- **Neutral theory of molecular evolution**: most evolutionary changes occur at the molecular level.
- Most variation is due to random genetic drift of mutant alleles that are selectively neutral.
- Compatible with phenotypic evolution, as phenotypes are driven by molecular changes.
- o Most mutations are neutral with respect to fitness.
- A minority of mutation are advantageous.
- Substitution rate predicted to be neutral, equal to per-individual mutation rate, independent of population size.
- \circ K_A/K_s test used to determin direction selection based on evolutionary history.
 - K_A : number of nonsynonymous substitutions (replacement).
 - K_S : number of synonymous substitutions (silent).
 - $K_A > K_s$ signals for diversifying selection.
 - $K_A < K_s$ signals for purifying selection.

Molecular Clocks

- **Molecular clocks**: the average rate at which species' genomes accumulates neutral mutations over time.
 - A linear rate is often easy to establish.
 - Used to measure evolutionary divergence.

Nonrandom Mating

- ▶ **Inbreeding**: production of offspring from closely genetically related individuals.
 - Results in homozygosity, which can increase chances of offspring being affected by deleterious or recessive traits.
 - Inbreeding depression: the reduced fitness in a given population due to inbreeding.
 - Usually caused by population bottlenecks or the founder effect.
 - Can also result in purging of deleterious allels through purifying selection.
 - Can allow for the expression of advantageous phenotypes, which if outweighs the disadvantages, then could potentially lead to speciation.
 - **Coefficient of inbreeding**: the probability that two alleles at any locus in an individual are identical by descent.
 - Nonrandom mating does not alter allele frequencies and not a mechanism of evolution.
 - Can alter the frequencies of genotypes, changing the distribution of phenotypes in a population, which can alter patters of natural selection.
- > **Assortative mating**: mating based on phenotypic factors.
 - Can play a role in sympatric speciation.
 - A form of sexual selection.
 - Can be either positive of negative, selecting for similar or different phenotypes respectively.

8 Evolution at Multiple Loci

Linkage Equilibrium and Diesquilibrium

- ▶ **Linkage equilibrium**: when the genotype of a chromosome at one locus is independent of its genotype at another locus.
- ▶ **Linkage disequilibrium**: the non-random association of alleles at different loci in a given population.
 - Occurs when frequency of the association between loci's different alleles is higher or lower than expected.
- ▶ **Haplotype**: a group of alleles in an organism that are inherited together from a single parent.
 - Used to mean the collection of specific alleles that represent a phenotype and likely to be conserved.
 - Also can be used to mean a set of linked single-nucleotide polymorphism alleles that are associated statistically.
- ▶ Factors that influences disequilibrium: selection, rate or genetic recombination, mutation rate, genetic drift, system of mating, population structure.
 - Undertanding linkage desequilibrium in a genome can be a powerful signal of the population genetic processes that structure it.
 - Selection, genetic drift, assortative mating, and population admixture act to create disequilibrium.
 - Recombination and outbreeding act to reduce disequilibrium.
- ▶ Level of linkage disequilibrium can between A and B can be quantified by the coefficient of linkage disequilibrium, D_{AB} .
 - o Formula: $D_{AB} = P_{AB} P_A \overline{P_B}$
 - \circ P_{AB} : the frequency with which both occur together on same gamete, or the frequency of the AB haplotype.
 - \circ P_AP_B : product of the probabilities give the probability they occur together.
 - When there is a difference, the magnitude of the coefficient rises, indicating linkage disequilibrium.
 - Strong recent selection can be indicated by linkage disequilibrium of allels located next to neutral allels.

Recombination's Effect on Linkage Diesquilibrium

- Linkage diesquilibrium (D) will converge to zero depending of the magnitude of the recombination rate (c) between two loci the absence of natural selection, inbreeding, and genetic drift.
- The smaller the distance between the two loci, the smaller the rate of convergence of D to zero.
- Genetic recombination tends to randomize genotypes, thus it tends to reduce frequency of overrespresented chromosome haplotypes and increases underrepresented haplotypes.
- o In short, recombination reduces linkage disequilibrium.

Adaptive Significance of Sex

- ▶ **Parthenogenesis**: a natural form of asexual reproduction in which growth and development of embryos occur without fertilization.
- ▶ **Hermaphrodite**: an organism that has complete or partial reproductive organs and can produce gametes normally associated with both sexes.
 - o Allows the either partner to act as male of female in sexual reproduction.
 - **Sequential**: bon as one sex, but later changes into the opposite.
 - o **Simultaneous**: fully functioning male and female genitalia.

Maynard Smith Assumptions

- 1. A female's reproductive mode does not affect how many offsping she makes.
- 2. A female's reproductive mode does not affect the probability that her offspring will survive.
- Designed to question advantage of either mode of reproduction (asexual vs sexual).
- Predicts asexual females should be dominant reproduction form.
 - Due to the two-fold cost of sex: requires two individuals and some of the offspring are male.
 - Asexual reproduction of females would have all of the individuals reproduce and pass on genes, leading to quick dominance.
- The prediction is not seen, so it raises question of what the benifits of sexual reproductive are.

Violations of Maynard Smith Assumptions

- 1. Male parental care can influence how many offspring survives or can be made, though this violatin is rare.
- 2. Presence of self-compatible hermaphrodites and males in same organism means males must:
 - collectively fertilize a proportion of eggs that exceed their own frequency in the population (α) ;
 - and/or produce offsring that have a higher relative fitness (w).
 - Condition for male persistance: $\alpha w > 2$
- o Raises the question how male fertilization increases relative fitness.

Consequences of Sex

- Sexual reproduction: meiosis with crossing over and mating between unrealted individuals.
 - Meiosis results in genetic recombination.
 - Mating between unrealted individuals (outcrossing) results in allelic segregation.
- Segregation tends to restore the population to the Hardy-Weinberg equilibrium.
- Recombination tends to restore the population to linkage equilibrium.

Maintenance of Sex

- More attention is given to the maintenance of sex via its effects on linkage disequilibrium.
- Sex purges deleterious mutations.
 - Genetic load: the burden imposed by the accumulating mutations in a population.
 - Muller's ratchet: mutations should accumulate in asexual lineages.
 - · Lack of ability to purge mutations means that drift has higher chance of removing smaller proportion of mutation free individuals.
 - · Thus, number deleterious mutations increases in individuals remaining, overtime lowering fitness of lineage.
 - A higher mutation rate selects for more frequent outcrossing; thus more males (more sexual reproduction).

- More frequent outcrossing results in more recombination, which opposes genetic drifts tendency of creating linkage disequilibrium.
 - · Recombination recreates lost genotypes caused by random chance due to drift.
- Een if two individuals mate carrying deleterious mutations, then
 there is still a chance that they can produce mutation free offsping.
- Genes responsible for sex are maintained because they help create zero-deleterious-mutation genotypes.
- Sex accelerates adaptive evolution.
 - Natural selection also helps creates linkage disequilibrium, creating chances at losing advantageous genotype.
 - · Sex again introduces recombination allowing for recovery of lost advantageous genotypes.
 - · Recombination can be can also disadvantageous once optimal genotypes arise, raising question how it remains useful.
 - Red Queen hypothesis: sex remains beneficial indefintely in populations subject to ever-changing selection.
 - · Results in coevolutionary arms races.
 - · Genotypes in lower-than-equilibrium frequencies due to previous disadvantage may soon become advantageous.
 - Thus, sex is continually selected for as recombination is continually favored in changing conditions.
 - · Creates an accelrated rate of adaptive evolution in genes that take advantage of changing fitness of genotypes relative eachother.
 - Genes for sex are maintained in populations due to higher frequencies of genetic hitchhiking in genotypes they create.

9 Quantitative Genetics

The Nature of Quantitative Traits

- ▶ **Quantitative genetics**: deals with phenotypes that vary continuously due to multilocus traits and environmental factors.
 - Allows for the prediction of how a population will respond to selection, even when we do not know the genetic basis of the trait.
- ▶ Qualitative traits: descrete traits that can be assigned individual categories by observation of simple genetic test.
- ▶ Quantitative traits: continuous traits determined by the combined influence of the genotype at multiple loci and the environment; the focus of quantitative genetics.
 - Study of continuous distribution requires many other statistical methods such as the effect size, mean, and variance, to link phenotypes to genotypes.

Heritable Variation

- ▶ **Heritability**: the degree of variation in a phenotypic trait in a population that is due to genetic variation betwen individuals in that population.
- ▶ **Phenotypic variance (V** $_P$ **)**: the genetic variance (V $_G$) combined with the environmental variance (V $_E$).
 - $\circ V_P = V_G + V_E + V_{GE}$
 - \circ V_{GE} represents variance associated with intereaction of genetic and environmental factors.

Genetic Variance

- Additive genetic variance (V_A) : how much the phenotypic trait is influenced by traits that show an additive effect on the quantitative traits.
 - Measures the magnitude to which individual phenotypic differences can be predicted due to additive effects of allelic substitutions.
 - The greater the additive genetic variation for the trait, the greater is response to selection can be.
- \circ **Dominance genetic variance (V**_D): associated with the dominant gene actions which cover the influence of the recessive alleles at the particular locus.
- **Epistasis (V_I)**: occurs due to statistical interaaction among loci, i.e., gene-by-gene modification.

$$\circ V_G = V_A + V_D + V_I$$

▶ **Broad-sense heritability (H²)**: all genetic contributions to a populations phenotypic variance, including additive, dominant, and epistatic, and maternal/paternal effects.

$$\circ H^2 = \frac{V_G}{V_P}$$

- ▶ Narrow-sense heritability (h²): proportion of the total phenotypic variance that is due to the additive effects of genes.
 - $\circ h^2 = \frac{V_A}{V_P}$
 - o Allows prediction of how a population will respond to selection.
 - The greater the additive genetic variation for the trait, the greater its response to selection can be.

Estimating Heritability

- Heritability is often estimated by measuring correlations between parents and offspring.
- Midparnet: the average of the trait value between parents.
 - The mother's value often has a scaling value applied to help account for developmental sex differences.
- Midoffspring: the average value of a trait among offsping.
- o Correlations are determind through a line of best fit.
 - Heritability of the trait is measued between 0 and 1 determind by the slope of the best fit line plotted against midparant and midoffsping values.
 - Using a method of least-squares linear regression, which minimizes the sum of the squared vertical distances between points, the heritability slope represents narrow-sense heritability.
- If a trait is heritable, monozygotic twins will resemble each other more than dizygotic twins.

- **Selection differential (S)**: the difference between the mean of the selected individuals (\bar{x}) and the mean of the entire population (μ)
 - $-S=\bar{x}-\mu$
 - Can be used to create a statical selection gradient, which is used to represent the strength of selection.
- **Breeder's equation**: evolutionary response (R) can be predicted using narrow-sense heritability and the selection differential.
 - $R = h^2 S$
 - The greater the heritability and selection differential, the greater the response to selection.
- When characters are genetically correlated, selection for one can drag the other along through genetic hitchhiking.
- Selection on the alleles at any single locus affecting a quantitative is often very weak; substantial genetic variation may persist at equilibrium between mutation and selection.
- Low heritability often represents populations effected by strong selection, as selection removes variation.
- Higher heritability represents high variance within the population.

11 Sexual Selection

Sexual Dimorphism and Sex

⊳ Sexual Dimorphism:

Sexual Dimorphism in Humans

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Sexual Selection by Sex

▷ -

Selection on Males: Competition

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Selection on Males: Female Choice

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Selection on Females

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Sexual Selection in Plants