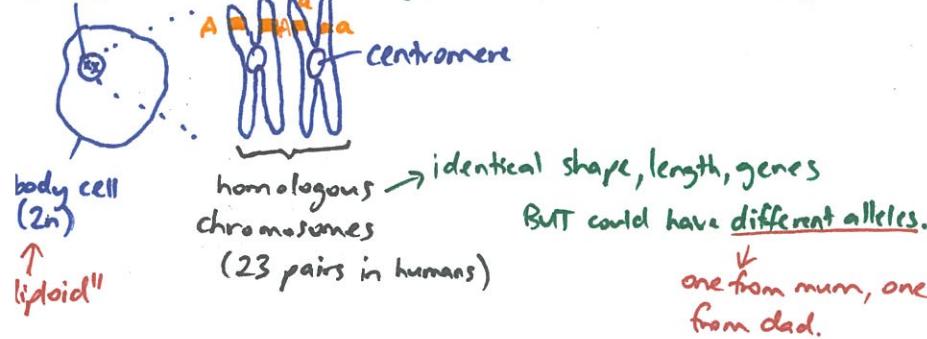
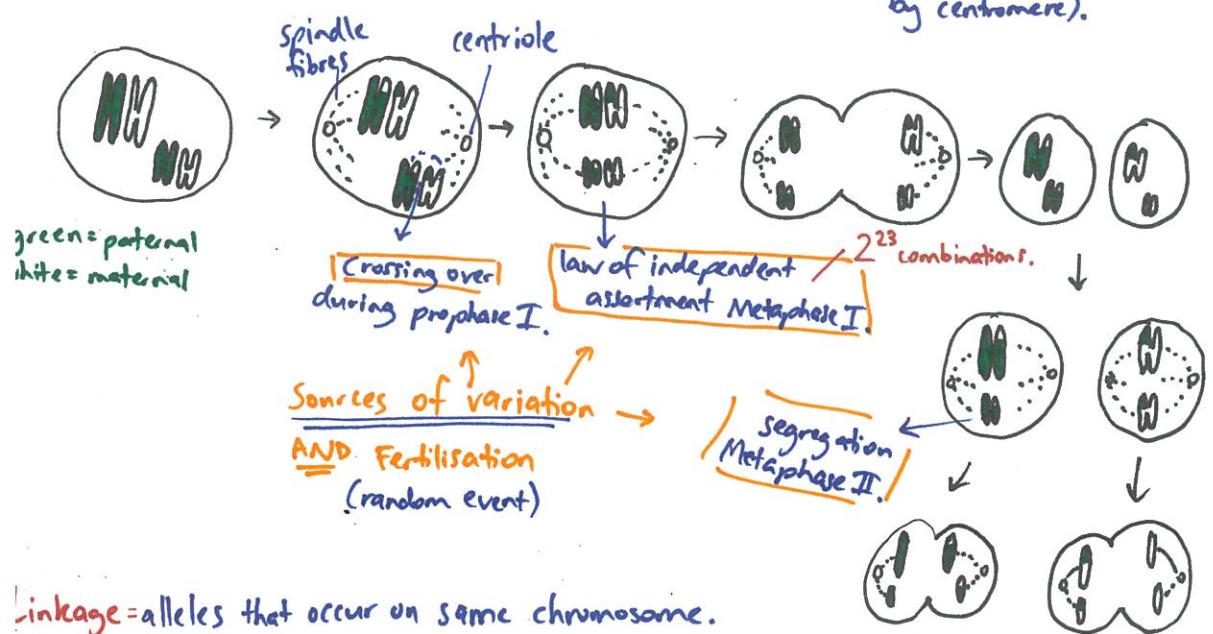


- Meiosis**
- 2 divisions, 1st set = homologous pairs separated, 2nd = chromatids separated.
  - occurs in sex organs (ovaries & testes)
  - produces gametes (sperm + ova)  $\leftarrow$  "haploid"
  - result = 4x daughter cells  $n=23$



\* BEFORE meiosis can begin, each chromosome must be replicated (DNA replication) to form 2 chromatids (held together by centromere).



Linkage = alleles that occur on same chromosome.

↳ inherited together.

↳ can be separated by crossing over.

4x genetically unique gametes {

↳ produces different combinations of alleles.

linked alleles  $\rightarrow$  R<sup>1</sup>R<sup>2</sup>, r<sup>1</sup>r<sup>2</sup>

multiple alleles  $\rightarrow$  Monohybrid - complete dominance  
Inheritance:  $\rightarrow$  Dihybrid - incomplete dominance  
- co-dominance

Monohybrid = one characteristic controlled by one gene

'capital letter' = dominant allele i.e. R

'lower case' = recessive allele i.e. r

heterozygous = Rr or RrTt 2 genes, 4 alleles

homozygous = rr or RR or RRTT or rrtt

Complete dominance: only 2 possible phenotypes.

only one dominant allele needed to be present to have full dominant effect and mask recessive allele (if present)

Multiple alleles: > 2 alleles for a gene. You will only ever find 2 present in genotype. e.g. IAIB in blood grouping. (3x possible alleles).

2x heterozygotes (Rr x Rr)

$\begin{array}{c} R\ r \\ R\ R\ R\ r \\ R\ r\ r\ r \end{array}$

phenotype = 3:1 ratio

expected... this is a chance event!

2x diff. uppercase letters used.

Codominance = for heterozygous individuals, both alleles are equally dominant, both alleles expressed in phenotype, independent of each other.

e.g. cows R = red W = white RW = roan (mixture of some hair red and some white)

## 2.5 Genetic Variation (9/157)

gene = basic unit of heredity, a length of DNA that codes for a protein/trait.

allele = alternative form of a gene.

genotype = genetic make up of an individual, the actual alleles an individual has.

phenotype = expression of the genotype, what is seen

genotype + environment  $\rightarrow$  phenotype

gene pool = all alleles present in a population.

allele frequency = no. of times an allele occurs in a population.

mutation = change in base sequence of DNA. They are spontaneous, random, rare.

↳ could be increased by mutagens (UV, chemicals, environmental factors).

gene mutations = insertion, deletion, substitution.

↳ same sense, mis-sense, non-sense.

↳ source of genetic variation.

↳ if in gametes, mutation is heritable, new mutated allele enters gene pool.... now subject to Natural Selection.

"selected against" = dies/no reproduction.  
"selected for" = survives & reproduces.

Genetic diversity = range of alleles present in gene pool. ↑ no. different alleles = ↑ genetic diversity.

Test cross: breeding w/ homozygous recessive to determine genotype of an individual... if no offspring have recessive trait  $\rightarrow$  original parent must be homozygous.

Incomplete dominance: for heterozygous individuals, neither allele dominates the other, both contribute to produce an intermediate/blended phenotype.

- 3x possible phenotypes e.g. RR = red snapdragon flowers

rr = white "

Rr = pink "

" "

" "

" "

" "

" "

" "

" "

## Population Genetics, Key definitions

Gene Pool: total no. of alleles present in population.

mutation  
Factors affecting frequencies  
natural selection

Evolution: change in allele frequencies over time.  
Process of how new species develop.  
migration  
genetic drift

Mutations: source of new alleles in gene pool. Essential for Evolution

Genetic drift: change in allele frequencies due to CHANCE (not selection)  
↳ prominent in small populations. e.g. recessive allele lost due to BOTTLENECK

Founder Effect: small group colonises geographically isolated area.  
Not a good representation of allele frequencies in original population.  
e.g. seed colonising NZ from Australia.

Bottleneck Effect: pop. rapidly reduced in numbers due to catastrophic event or sudden selection pressure. If pop. recovers  $\rightarrow$  reduced diversity.

Affected by Independent Assort?	increases genetic variation?	Crossing over?	no. of homologous pairs.	no. of possible gametes
Linked:	$\times$	$\checkmark$	1	2
DE DE Dd Dd EE FF Ee Ff	$\times$ $\times$ if Xing over.	$\checkmark$ depends how close	DE de	DE de

Not Linked:	$\checkmark$	$\checkmark$	$\checkmark$	4
DE Addd EE FfFf	$\checkmark$	$\checkmark$	$\checkmark$	DE de

Linkage: ignoring Crossing over... first

Yy Ff Xy yf ff =

1:1:1:1 expected phenotype ratio.

But, if linked, ratio changes due to recombinants.

↓ linkage  $\rightarrow$  linkage ratio

Dihybrid Inheritance: 2 characteristics controlled by 2 genes. Could be linked or not linked. e.g. w/ no linkage

TY YY x Ty yy = all F1 Tt Yy  
(Ty x ty)

Tt Yy x Tt x Yy =

possible gametes = TY, Ty, tY, ty

TY Ty tY ty

TY + + + +  
Ty + + + +  
tY + + + +  
ty + + + +

q: 3:2:1 phenotype ratio.