

AP Biology: Unit 5 Cheat Sheet

by kmz_2022 via cheatography.com/145729/cs/31651/

Comparing Reproduction				
ASEXUAL vs. SEXUAL				
asexual	similarities	sexual		
- 1 parent	- creates offspring	- 2 parents		
- identical genetics	offspring gorw/develop	- genetic variation		
- quicker & less energy	- use DNA	- slower & more energy		
advantages		advantages		
- well-adapted (stable environment)		- increased diversity		
- large # offspring quickly		- adapt to changed (unstable environment)		
- reliable (fewer steps)		- genetic variation		
- no need for a mate				
- less time & energy				
MEIOSIS vs. MITOSIS				
meiosis	similarities	mitosis		
- 2 divisions = 4 daughter cells	- interphase	- 1 division = 2 daughter cells		

Comparing Reproduction (cont)				
- genetic variation		- forms of reprod- uction	- genetically identical	
- half # chromo- somes (23)		- stage names	- same # chromo- somes (46)	
- sex cells/gametes			- somatic cells	
Meiosis Definit	ions			
- heredity:	transmi	ssion of traits from	1 generation to the next	
- gametes:	haploid reproductive cells (egg/sperm)			
- gene locus:	gene's specific location on the length of a chromosome			
- karyotype:	display of chromosomes matched up with their pairs (based on length)			
homologous chromosomes:	pair of chromosomes that have the same length, stain pattern, and genes controlling the same characteristics			
- sex chromo- some:	chromosome responsible for determining the sex of an individual			
- autosome:	chromo	some NOT directly	involved in determining	
- diploid cell~	2 chron	nosome sets (2n) =	: 46; somatic cells	



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- haploid

cell~

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1 chromosome set (n) = 23; gametes



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Meiosis Background

Meiosis I is called?

Meiosis II is called?

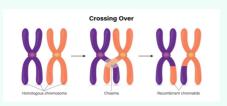
Original source of all genetic diversity?

→ mutations

Term for different version of a gene?

alleles

Variation



- crosing over: genetic rearrangment between sister chromatids by exchanging corresponding segments of DNA
- combining DNA from 2 parents into a single chromosome = variation
- independent assortment: each pair of homologous chromosomes are positioned independently of other pairs
- → each daughter cell represents 1 outcome --- formula: 2ⁿ
- random fertilization~
- \Rightarrow fusion of gametes (2²³ x 2²³) = *variation*

Genetics Definitions

blending genetic material contributed by both parents mixes
 hypothesis~ (like paint)
 particulate parents pass on genes that retain their separate

identities in offspring (like deck of cards)

Genetics Definitions (cont)

- true-b-	plants, that after many generation of self-pollination,
reeding	have produced only the same variety as the parent
strain~	(homozygous)

 hybridiza- crossing of 2 true-breeding varieties tion:

- P genera- the true-breeding parent individuals

tion:

- F1 hybrid offspring arising from a parental cross ('first

generation: filial)

- F2 offspring from the interbreeding of the F1 generation

generation: (second filial)

- homozy- 2 identical alleles

gous:

ZVOOUS:

heterozygous:

physical/physiological traits

2 different alleles

phenotype:

- genotype: genetic makeup/set of alleles

- testcross: breeding an organism of unknown genotype w/

homozygous recessive to determine the unknown

genotype

- cross between 2 organisms that *heterozygous* for the

monohybrid

cross:

- dihybrid cross between 2 organisms that are heterozygous for

cross: both traits



hypothesis~

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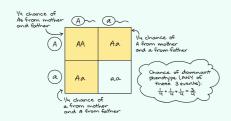
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Mendel's 3 Laws

- Law of Dominance: Some alleles are dominant and cover up the recessive alleles.
- **2.** Law of Segregation: An organism has two alleles for each gene but they can only pass on one.
- Law of Independent Assortment: Genes found on separate chromosomes are inherited independently of each other.

Blood Types	
A	I^AI^A/I^Ai
В	I^BI^B/I^Bi
0	ii
AB	I^AI^B
alleles present=	3
recessive allele=	0
codominant alleles=	A & B

Probability



- multiplication rule of probability: probability of independent events occurring at the same time is the *product of their individual proababilities*
- addition rule of probability: probability of mutually exclusive events occurring is the sum of their individual probabilities

Incomplete Dominance



- incomplete dominance: phenotype between both parents

Codominance

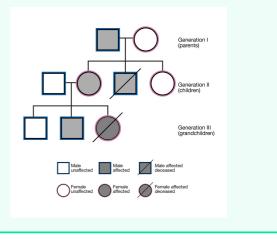


- codominance: 2 alleles that each affect the phenotype in separate ways
- \rightarrow R' \rightarrow protein \rightarrow item A \rightarrow red
- $\downarrow R \rightarrow \text{protein} \rightarrow \text{item B} \rightarrow \text{pink}$

Sex-linked Genes

- male = XY
- female = XX
- X inactivation: most of 1 X chromosome in each cell becomes inactivated
- Barr bodies: inactive X chromosome condenses
- -gene SRY (sex-determining region of Y): Y gene results in a male
- makes transcription factor that binds to other genes causing them to transcribe
- sex-linked genes: genes on sex chromosomes
- father passes Y to sons & X to daughters
- mother passes X to sons/daughters
- sex determination in birds:
- → sex chromosome in *egg*
- *→* male = ZZ
- *→ female =* ZW

Pedigree Chart





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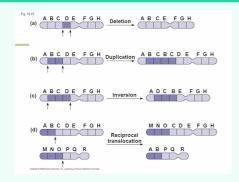
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Chromosomal Mutations



- large scale chromosomal mutation = change in phenotype
- three possible causes of mutations:
- 1. physical/chemical disturbances
- 2. errors during meiosis
- 3. random mutation

Chromosomal Disorders		
- nondisjunction:	pair of homologous chromosomes/sister chromatids fail to separate	
- aneuploidy:	1 or more chromosomes have extra copies/deficient number	
- monosomic (monosomy):	diploid cell that has <i>1 copy</i> of a chromosome (not 2)	
- trisomic (trisomy):	diploid cell that has $3 copies$ of a chromosome (not 2)	
- polyploidy:	organism has more than 2 complete chromosome sets	
- Down syndrome:	extra chromosome 21 (trisomy)	
- Klinefelter syndrome:	extra X chromosome in males (XXY)	
- Turner syndrome:	loss of X chromosome in females (X0)	
- XXY:	extra Y chromosome in males	
- XXX:	extra X chromosome in females	

Genomic Imprints

genomic imprints:

expression of an allele in offspring depends on whether the allele is inherited from mother/father

When does it occur?

during gamete formation

What chemical change does it involve?

methyl group added to cytosine nucleotides (inactivates alleles)

Which organelles contain their own DNA?

mitochondria & chloroplasts

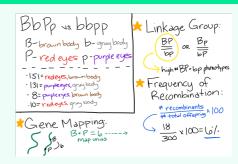
When do these organelles get passes to offspring?

during fertilization, mitochondrial DNA from mother

Why are genetic defects of mitochondrial genes likely to affect the functioning of the nervous/muscular system?

they are the most susceptible to energy deprivation (which ATP comes from mitochondria)

Linkage Group



- linkage group: all the loci that "move together" in inheritance
- frequency of recombination: the frequency that crossing over will occur between two genes
- gene map: chromosome map that shows the relative locations of genes
- **linked genes**: located close enough together on a chromosome that they tend to be inherited together
- 'wild type': phenotype most commonly observed



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