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## Guided Notes <br> Unit 6: Classical Genetics

## Chapter 6: Meiosis and Mendel

## I. Concept 6.1: Chromosomes and Meiosis

a. Meiosis: $\qquad$
i. (In animals, meiosis occurs in the sex organs-the testes in males and the ovaries in females.)
b. Somatic Cells and Gametes
i. Somatic cells: $\qquad$
ii. Gametes: $\qquad$ (example: egg and sperm cells)
c. Homologous Chromosomes
i. Cells from males and females of the SAME species have the SAME number and types of chromosomes.
ii. Homologous chromosomes: $\qquad$ ___ (one chromosome in the pair comes from the mother, one comes from the father)

d. Chromosomes
i. Humans have $\qquad$ homologous pairs of chromosomes.

1. (So you have $\qquad$ chromosomes in total.)
ii. Autosomal chromosomes: chromosome pairs 1-22 in humans
iii. Sex chromosomes: $\qquad$ ; the 23rd
pair of chromosomes; occurs in two forms, called $X$ and $Y$
2. Male mammals $=$ $\qquad$
3. Female mammals $=$ $\qquad$
e. Karyotypes
i. Karyotype: a display of $\qquad$ of an
individual (46 in humans)

f. Diploid and Haploid Cells
i. Diploid (2n): $\qquad$
4. Example: body cells
ii. Diploid number: the total number of chromosomes in the cell (46 in humans)
iii. Haploid (n): $\qquad$
5. Example: gametes (sperm \& egg cells)

g. Human Life Cycle
i. $\qquad$ $+$ $\qquad$ $\rightarrow$
(fertilization)
ii. Zygote: has $\qquad$
develops into a sexually mature adult with trillions of cells produced by mitosis

h. The Process of Meiosis
i. Meiosis is different from mitosis.
ii. Meiosis produces $\qquad$ new offspring cells, each with one set of chromosomes-thus half the number of chromosomes as the parent cell.
iii. (Mitosis produces $\qquad$ offspring cells, each with the same number of chromosomes as the parent cell.)
iv. Meiosis produces four haploid cells, rather than two diploid cells. Also, meiosis involves two divisions, whereas mitosis involves only one.

II. Concept 6.2: Process of Meiosis
a. Meiosis consists of two distinct parts - $\qquad$ and $\qquad$ .
i. In meiosis I, homologous chromosomes, each composed of two sister chromatids, are separated from one another.
ii. In meiosis II, sister chromatids are separated much as they are in mitosis. However, the resulting cells are haploid rather than diploid.

b. Interphase



c. Meiosis
i. The process of meiosis is completed, producing $\qquad$ cells as a final result.
ii. Each gamete made from meiosis can have a large variety of possibilities for genetic variation.
6. (Remember, human gametes have 23 chromosomes, so the possibilities are numerous! This is why you and your siblings are not identical.)

d. Review: Comparison of Mitosis and Meiosis
iii. Mitosis
7. Provides for $\qquad$
8. Produces daughter cells that are genetically identical to the parent cell
9. Involves $\qquad$ division of the genetic material in the nucleus (usually accompanied by cytokinesis, producing two diploid cells)
iv. Meiosis
10. Produces $\qquad$ which allows for sexual reproduction
11. Yields haploid daughter cells with only one set of homologous chromosomes
12. Involves $\qquad$ nuclear divisions, yielding four haploid cells
13. Duplicated homologous chromosomes form tetrads
14. The chromosome number in each of the two daughter cells is haploid
v. Both Mitosis and Meiosis
15. Chromosomes duplicate only once, in the preceding Interphase
16. Meiosis II is basically identical to mitosis. (The sister chromatids separate, and each cell divides in two.)
17. Both make it possible for cells to inherit genetic information in the form of chromosome copies

III. Concept 6.3: Mendel and Heredity
a. Vocabulary
i. Trait: $\qquad$
18. (Example: a plant's trait of having red flowers, or yellow flowers, etc.)
ii. Genetics: $\qquad$
b. Gregor Mendel: an Austrian monk who found the $\qquad$
i. (He showed that traits are inherited as discrete units!)
c. Prior to Mendel
i. Blending hypothesis: hypothesis in 1800 s to explain how offspring inherit traits from both parents
19. (Example: red and yellow flowered plants could produce an orange flowered plant)
ii. BUT this was $\qquad$ !
d. Mendel's Work
i. Mendel bred pea plants and recorded $\qquad$ in
the offspring for 7 years.
ii. True-breeding Plants
20. Mendel fertilized true-breeding pea plants - meaning when they $\qquad$ , the offspring are identical to the parent pea plant.
a. (Example: purple-flowered pea plant makes purple-flowered pea plant offspring)
b. (Your book refers to "true-breeding plants" as purebred.)

21. (Mendel experimented with these seven pea plant characters.)
iii. Cross-Fertilization
22. Cross-fertilization: $\qquad$
$\qquad$


Cross seen above: purple flower x white flower
iv. Hybrids

1. Hybrid: $\qquad$
2. Monohybrid cross: a pairing in which the parent plants differ in only one (mono) character
v. Mendel's Experiment
3. Mendel crossed the $\qquad$ generation to produce the $\qquad$ generation.
4. Notice the results:

vi. Mendel's Conclusion
5. F1 generation: all plants had purple flowers
6. F2 generation: some plants had purple flowers and some had white

7. Mendel concluded that the trait for the white flower had $\qquad$
vii. Cross Results
8. Mendel performed many crosses to help him observe inheritance patterns.

FIGURE 6.10 MENDEL'S MONOHYBRID CROSS RESULTS

| $\mathrm{F}_{2}$ TRAITS | DOMINANT | RECESSIVE | RATIO |
| :--- | :--- | :--- | :--- |
| Pea shape | 5474 round | 1850 wrinkled | $2.96: 1$ |
| Pea color | 6022 yellow | 2001 green | $3.01: 1$ |
| Flower color | 705 purple | 224 white | $3.15: 1$ |
| Pod shape | 882 smooth | 299 constricted | $2.95: 1$ |
| Pod color | 428 green | 152 yellow | $2.82: 1$ |
| Flower position | 651 axial | 207 terminal | $3.14: 1$ |
| Plant height | 787 tall | 277 short | $2.84: 1$ |

viii. Mendel's Three Conclusions

1. Traits are $\qquad$ .
2. Organisms inherit $\qquad$ copies of each gene, $\qquad$
$\qquad$ .
3. The two copies $\qquad$ .
a. (Numbers 2 and 3 make up Mendel's Law of Segregation.)

## IV. Concept 6.4: Traits, Genes, and Alleles

a. Genes
i. Gene: $\qquad$
$\qquad$
ii. Each gene has a locus, a $\qquad$
$\qquad$ on a pair of homologous
chromosomes.

b. Alleles
i. Alleles: $\qquad$

1. (Example: allele for blue eyes (b) vs. allele for brown eyes (B))
2. Each parent donates one allele for every gene.
ii. Homozygous: two alleles that are the $\qquad$ at a specific locus
3. (Example: BB or bb )
iii. Heterozygous: two alleles that are
$\qquad$ at a specific locus
4. (Example: Bb)

Homozygous alleles are identical to each other.


Heterozygous alleles are different from each other.
c. Genotype and Phenotype
i. Genotype: genetic makeup or combination of alleles $\qquad$ _)

1. Example: PP
ii. Phenotype: an observable trait (
2. Example: purple flowers
d. Allele Dominance
i. Dominant allele: an allele in a heterozygous individual that
$\qquad$
3. (Example: allele for brown eyes (B))
ii. Recessive allele: an allele in a heterozygous individual that
$\qquad$
$\qquad$
4. (Example: allele for blue eyes (b))


## V. Concept 6.5: Traits and Probability

a. Punnett Square
i. Punnett square: a grid system for predicting all possible genotypes resulting from a cross - it gives the ratio of possible $\qquad$

b. Monohybrid Cross
i. Monohybrid cross: examine the inheritance of $\qquad$
ii. Examples:

c. Testcross
i. Testcross: a cross between an organism with $\qquad$ genotype and an organism with $\qquad$ phenotype

d. Dihybrid Cross
i. Dihybrid cross: examines the inheritance of $\qquad$

ii. Independent Assortment

1. Mendel's dihybrid crosses led to his second law, the $\qquad$
$\qquad$ . This law states
that allele pairs separate $\qquad$ of each other during meiosis.

(Example (above): $R$ can end up with either $Y$ or $y$, and $r$ can end up with either $Y$ or $y$ )
e. Probability
i. Probability:
2. Genotypic ratio: the ratio of the
a. (1 PP : $2 \mathrm{Pp}: 1 \mathrm{pp}$ )
3. Phenotypic ratio: the ratio of the
a. (3 purple : 1 white)


## IV. Concept 6.6: Meiosis and Genetic Variation

a. Genetic Variation
i. Sexual reproduction creates unique combination of genes.
ii. How?
$\qquad$ in
meiosis
a. (Think back to Mendel's law...)

2.
a. AKA - you don't know which gamete will be used to make the zygote!
iii. Unique phenotypes may give a reproductive advantage to some organisms.

## Chapter 7: Extending Mendelian Genetics

I. Concept 7.1: Chromosomes and Phenotype
a. Autosomal Genes
i. Autosomal gene: $\qquad$
$\qquad$ (chromosomes 1-22 in humans)
ii. $\qquad$ copies of each autosomal gene affect phenotype.
iii. Autosomal Disorders

1. Mendel's rules of inheritance apply to autosomal genetic disorders.
2. Carrier: $\qquad$
3. Disorders caused by dominant alleles are uncommon.

| heterozygous <br> parent (Cc), <br> carrier <br> C | heterozygous parent (Cc), carrier C c |  |
| :---: | :---: | :---: |
|  | CC homozygous dominant | Cc <br> heterozygous, carrier |
|  | Cc <br> heterozygous, carrier | CC <br> homozygous recessive, affected |
|  | $\begin{aligned} & =\text { Normal allel } \\ & =\text { Cystic fibros } \end{aligned}$ | minant) <br> lele (recessive) |

b. Sex-Linked Genes
i. Sex-linked genes: $\qquad$ (chromosome 23
in humans)

1. Y chromosome genes in mammals are responsible for male characteristics.
2. $X$ chromosome genes in mammals affect many traits.
ii. Male and Female Genotypes
3. Male mammals: $\qquad$ genotype
a. All of a male's sex-linked genes are expressed.

b. Males have no second copies of sex-linked genes.
4. Female mammals: $\qquad$ genotype
a. Expression of sex-linked genes is similar to autosomal genes in females.
b. X chromosome inactivation randomly "turns off" one X chromosome.
iii. Sex-Linked Cross
5. Example: fruit flies
6. Allele for red eyes $=$ dominant
7. Allele for white eyes $=$ recessive
8. It is extremely rare to find a female with white eyes.
9. Why might this be?

P


White-eyed male

$X^{r} Y$
a. This inheritance pattern is located only on the $X$ chromosome. There is no corresponding eye color locus on the Y .
b. Females (XX) carry $\qquad$ copies for eye color.
c. Males (XY) carry only $\qquad$ for eye color.
i. (SO - a female will have white eyes only if she has the white-eye allele on both her $X$ chromosomes, but males will only need one allele on their one X chromosome.)

iv. Sex-Linked Disorders

1. Sex-linked disorders are disorders that are inherited as $\qquad$
$\qquad$ . (the same way as the white-eye trait in fruit flies)
2. Examples: red-green color blindness and hemophilia (a disease in which blood fails to clot normally)
3. These are more common in $\qquad$ .
a. (If a human male inherits the sex-linked recessive allele from his mother, the allele will be expressed - whereas females must inherit two alleles to exhibit the trait!)
II. Concept 7.2: Complex Patterns of Inheritance
a. Many characters of organisms have more complicated inheritance patterns than those studied by Mendel.
b. Incomplete Dominance (sometimes referred to as Intermediate Inheritance)
i. For some characters of organisms, $\qquad$ .
ii. Incomplete dominance: pattern of inheritance where $\qquad$

4. Example: red and white parents produce F 1 hybrid offspring that are pink - neither the red nor white allele is dominant
iii. This inheritance pattern $\qquad$
5. (This is because the parent phenotypes can reappear in the F2 generation.)

c. Codominance (sometimes referred to as Multiple Alleles)
i. Codominant alleles will $\qquad$ .
ii. Codominant alleles are $\qquad$ -.
6. The ABO blood types result from codominant alleles.
7. Three alleles for blood type in the human population: $\qquad$ , $\qquad$ ,
8. (But note that any one person has only two alleles for blood type.)
9. Result in 6 genotypes \& 4 phenotypes

a. Alleles $I^{A}$ and $I^{B}$ are $\qquad$ .
b. Allele $i$ is $\qquad$ .
iii. The individual's phenotype is not incomplete, but rather shows the separate traits of both alleles.
d. Polygenic Inheritance
i. Polygenic inheritance: $\qquad$

e. Environmental Affect
i. An individual's $\qquad$ as well as on genes.
10. (Example: a tree's genotype does not change, yet the tree's leaves vary in size, shape, and greenness from year to year (depending on exposure to wind and sunlight))
ii. For humans, many phenotypes vary due to environmental factors.
iii. On the other hand - some phenotype with little or no influence from the environment.
11. Example: human blood type
iv. In summary, the product of a genotype is generally not a single, rigidly defined phenotype, but a range of possibilities influenced by the environment.
III. Concept 7.4: Human Genetics and Pedigrees
a. How to Study Inheritance Patterns
i. To study the inheritance of a human trait:
12. $\qquad$
13. Organize the information in a $\qquad$
14. $\qquad$ by applying

Mendel's concepts of dominant and recessive alleles and his principle of segregation
b. Pedigrees
i. Pedigree: $\qquad$

1. Squares $=$ $\qquad$
2. Circles $=$ $\qquad$
3. Colored shapes = individuals that $\qquad$
ii. Parents are connected by horizontal lines, with their children beneath them in the order of birth.

4. Example: Earlobes are either free or attached. This pedigree tracks the occurrence of attached earlobes in three generations of a family.
iii. How do you figure out genotypes of each?
5. Determine the pattern in which the trait occurs
6. (Example: Notice that the first-born daughter in the third generation has attached lobes, although both of her parents have free earlobes. - so the attached-earlobe trait must be recessive. If the trait for attached earlobes were dominant, then at least one of her parents would have attached earlobes.)

7. The genotypes of most family members can be determined.
c. Types of Disorders
i. Disorders inherited as recessive traits
ii. Disorders inherited as dominant traits
iii. Sex-linked disorders (both dominant and recessive)
d. Disorders Inherited as Recessive Traits
i. Most human genetic disorders are recessive.
ii. Carrier: $\qquad$
iii. Example: a particular form of deafness is inherited as a recessive trait.

e. Disorders Inherited as Dominant Traits
i. A $\qquad$ number of human disorders are inherited as dominant traits.
ii. Example: achondroplasia (a form of dwarfism - "little people")
8. (About 1 out of 25,000 people has achondroplasia - all individuals with this disorder are heterozygous. More than 99.99\% of the population is homozygous for the normal, recessive allele so it clear that dominant alleles are not necessarily more plentiful than recessive alleles in a population.)
f. Sex-linked Disorders

i. Females $\qquad$ sex-linked genetic disorders.
ii. Males (XY) $\qquad$ their sex linked genes.
iii. Most sex-linked alleles are located on the X chromosome. A male only receives such sex-linked alleles from his mother. (The homologous Y chromosome is always inherited from the father.)
iv. Therefore, if the phenotype is more common in males, the gene is likely sex-linked.
v. Pedigrees for Sex-Linked Disorders
9. Example: red-green colorblindness
10. The half-filled circles represent
$\qquad$ -.
11. (This pedigree shows the appearance of colorblindness in four generations of a family.)
12. It is rare—but not impossible-for females to exhibit sex-linked (Xlinked) traits.


## Chapter 9: Frontiers of Biotechnology

## I. Concept 9.6: Genetic Screening and Gene Therapy

a. Genetic Screening
vi. Genetic screening: $\qquad$
i. Determines risk of having or passing on a genetic disorder
ii. Used to detect specific genes or proteins
iii. Can detect some genes related to an increased risk of cancer
iv. Can detect some genes known to cause genetic disorders

b. Gene Therapy
vii. Gene therapy: $\qquad$
i. Gene therapy replaces defective or missing genes, or adds new genes, to $\qquad$
ii. Gene Therapy Techniques

1. Genetically engineered viruses used to "infect" a patient's cells
2. Insert gene to stimulate immune system to attack cancer cells
3. Insert "suicide" genes into cancer cells that activate a drug
iii. Gene Therapy Challenges
4. Inserting gene into correct cells
5. Controlling gene expression
6. Determining effect on other genes
