

INTRODUCTION TO GENETICS

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Genetics I

MA 205 A

First week

10/5/2023



Outline

- Genetics
- subdivisions of genetics
- Genetic terms



Objectives

- ∘ Define (genetics, gene,.....)
- Compare homozygous and heterozygous



- Genetics: is a branch of biology concerned with the study of heredity and variation.
- It aims to explain how characteristics of living organisms are passed on from one generation to the next.
- Genetic science deals with:
- Transmission of traits from parent to offspring.
- Expression and variation of those traits
- The structure and function of the genetic material



• Four major subdivisions of genetics:

- Classical genetics: describes how traits (physical characteristics) are passed along from one generation to another.
- Molecular genetics: the study of the chemical and physical structures of DNA, RNA and protein.
- **Population genetics**: is a subfield of genetics that deals with genetic differences within and between populations.
- Quantitative genetics: a highly mathematical field that examines the statistical relationships between genes and the traits they encode.



- Gamete: gametes are sex cells (in animal: sperm and ovum; in plants: pollen nucleus and ovum).
- Heredity: The transmission of characteristics from parents to offspring.
- Trait: specific characteristics that varies from one individual to another.
- Gene: genes are short lengths of DNA found on chromosomes. They code for specific proteins.



- Allele: Alleles are different versions of a particular gene.
- **Dominant allele:** An allele whose phenotypic effect is the same in both heterozygous and homozygous conditions.
- Recessive allele: An allele whose phenotypic effect is masked in the heterozygous by another dominant allele

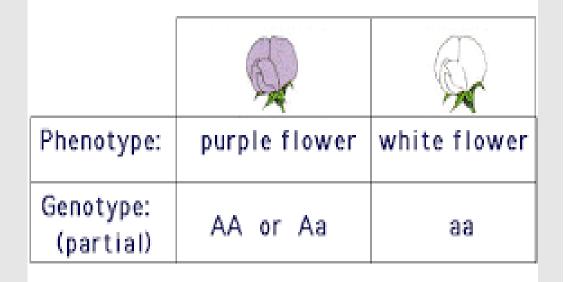


- Homozygous: homozygous individuals are individuals that possess either the same two copies of the dominant allele or two copies of the recessive allele.
- **Heterozygous:** heterozygous individuals are individuals that possess two different alleles of a gene.
- Carrier: heterozygous individual who does not display symptoms of a recessive genetic disorder but can transmit the disorder to his or her offspring.

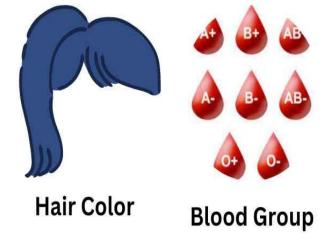


- **Genotype**: refers to the sets of genes carried by an organism that are capable of being passed on to the next generation.
- **Phenotype:** Observable characteristics of an organism, resulting from interactions between the genotype and environment.
- **Karyotype:** The general appearance of the chromosomes of an organism with regard to number, size and shape.

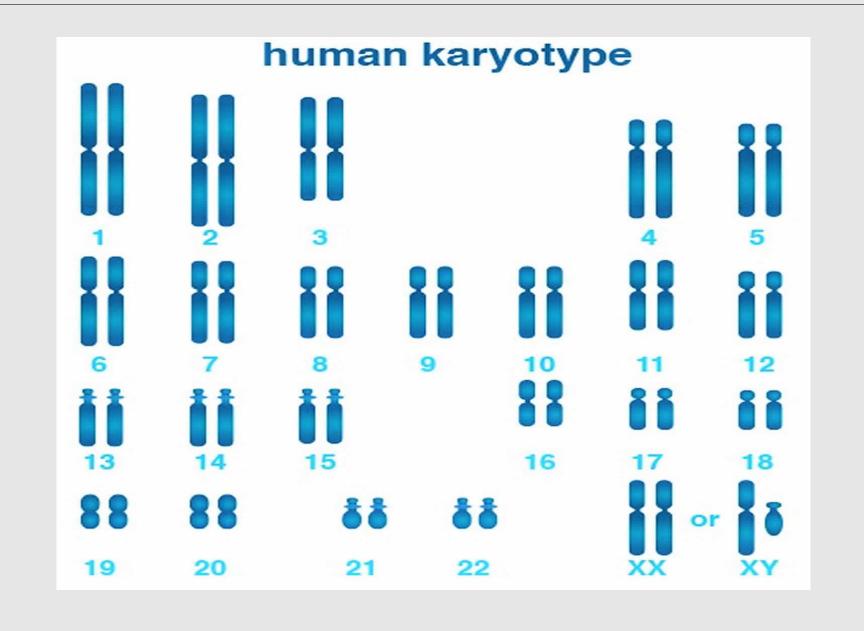
Genotype and Phenotype



Phenotype









- **Diploid**: a cell that contains both sets of homologous chromosome.
- **Haploid**: a cell that contains only a single set of chromosomes.
- Mutation: change in the nucleotide sequence of DNA.
- X-linked: pattern of inheritance, refers to an allele carried on the X chromosome.
- Chromatids: one of the two identical sister parts of a duplicated chromosome

- Genome: The complete set of chromosomes with their associated genes.
- Chromosome: chromosomes are thread like structures of DNA, carrying genetic information in the form of genes. They are located in the nucleus of cells
- **Autosomal chromosome**: in humans, the 22 pairs of chromosomes that are not the sex chromosomes (XX or XY).
- Sex chromosomes: pair of chromosomes involved in sex determination; in males, the XY chromosomes; in females, the XX chromosomes.



References:

- Gangane, SD. (2017) Human genetics, 5th edition. Elsevier. India.
- Pal, G.P. (2011) Medical genetics. Aitbs publishers and distributors, India.
- Peter, J.R. (2016) i Genetics. 3rd edition. Pearson education, India.