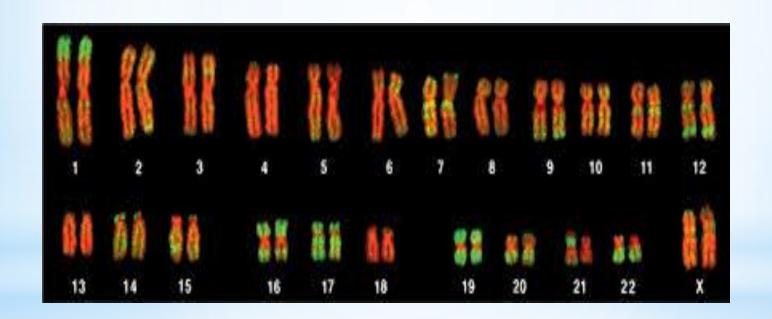
Lec 8 \ The Mutation



Structural Chromosomal mutations

Mutation is defined as a change *
occurring in the genetic material and
occurs at the gene or chromosome level
At the gene level, a mutation is defined
as any change in the nitrogenous bases
of DNA, which in turn leads to harmful
results

Natural Induced

Types of mutations *

There are mutations that lead to several changes in the parts that make up the internal structure of one of the chromosomes, which leads to a defect in its functioning from the normal state, including:

structural Chromosomal mutations

There are mutations that lead to several changes in the parts that make up the internal structure of one of the chromosomes, which leads to a defect in its functioning from the normal state, including:

Deletion: When a piece of a chromosome is deleted, this chromosome loses part of its active genes and becomes smaller in size than its normal counterpart

Inserion: Some mutations lead to the separation of one of the pieces from a specific chromosome, and it happens to be attached to another chromosome, which adds new genetic content to this chromosome.

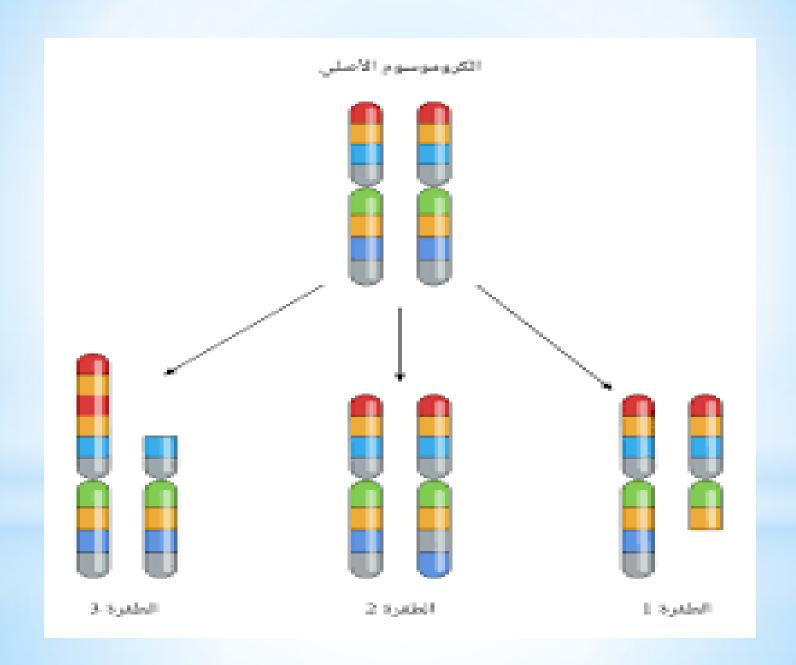
*

1\ structural Chromosomal mutations

Duplication: Duplication of one of the same chromosome segments occurs, which leads to an increase in the expression of these duplicated genes beyond the normal limit

Inversion: Some mutations cause a change in the arrangement of genes, making it inverted compared to the normal form and arrangement

Translocation: The transfer of segments between chromosomes occurs when a segment of one chromosome is lost and attached to another chromosome, leading to the appearance of genes in places other than their normal places.

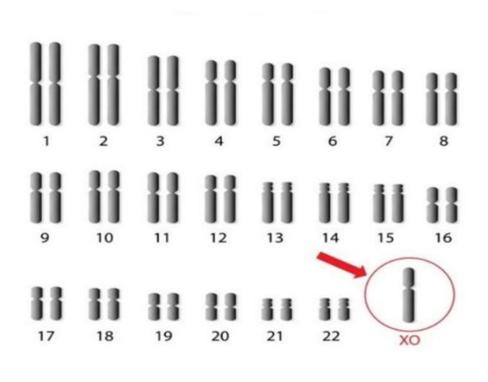


Numerical chromosomal mutation

It means mutations that lead to an increase or decrease in the number of chromosomes, not their structure, and can be divided, depending on the variable number, into:

1- Chromosome decrease occurs when the female loses one of the sex chromosomes number 23 (45X) Turner syndrome

Turner syndrome *





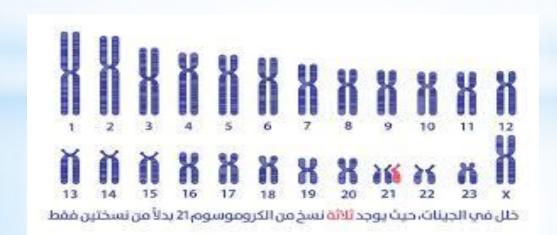






Down syndrome*

2- An increase in one chromosome is called a triple condition, as in Down syndrome (Mongolian), where an increase in body chromosome number 21 (47) occurs Down syndrome





Klinefelter syndrome

3- An increase in the X sex chromosome 23 found in males, as in Klinefelter syndrome (47XXY)

