

GENERAL EMBRYOLOGY



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WHAT IS EMBRYOLOGY ?

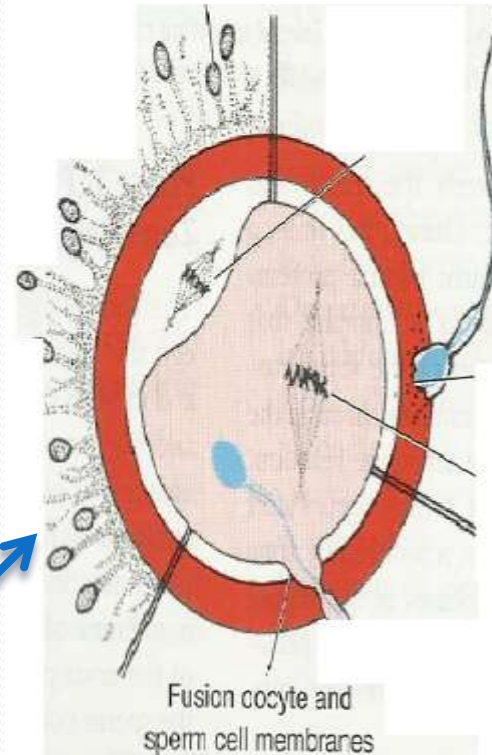
Is the science that deals with the **development** of the embryo from a single **cell** to a baby in **9 months**

Development begins with

FERTALIZATION

What is fertilization?

Fertilization is the process by which *the male gamete* the **sperm** and *the female gamete* the **oocyte** unite to form the **zygote**



Why do we need the union of **two cells** to form **the zygote**?

According to the number of chromosomes in the nucleus of the human cells we
Have **two** types :

1- Somatic cells

2- Reproductive cells (also called sex cells)

A somatic cell (*soma body*) is any cell of the body other than a germ cell.

A germ cell is a gamete
(sperm or oocyte)
or any precursor cell destined to become a gamete

Somatic cells : contain two sets of chromosomes:
first set contains 23 chromosomes coming from the *mother* called *maternal*
The second set contains 23 chromosomes coming from the *father* called *paternal*

Therefore, Somatic cells called

diploid

cells (*dipl-* double; *-oid form*), symbolized $2n$



The two chromosomes that make up each pair are called homologous
chromosomes

(*homo-* same) they

contain similar genes arranged in the same (or almost the same) order

What are

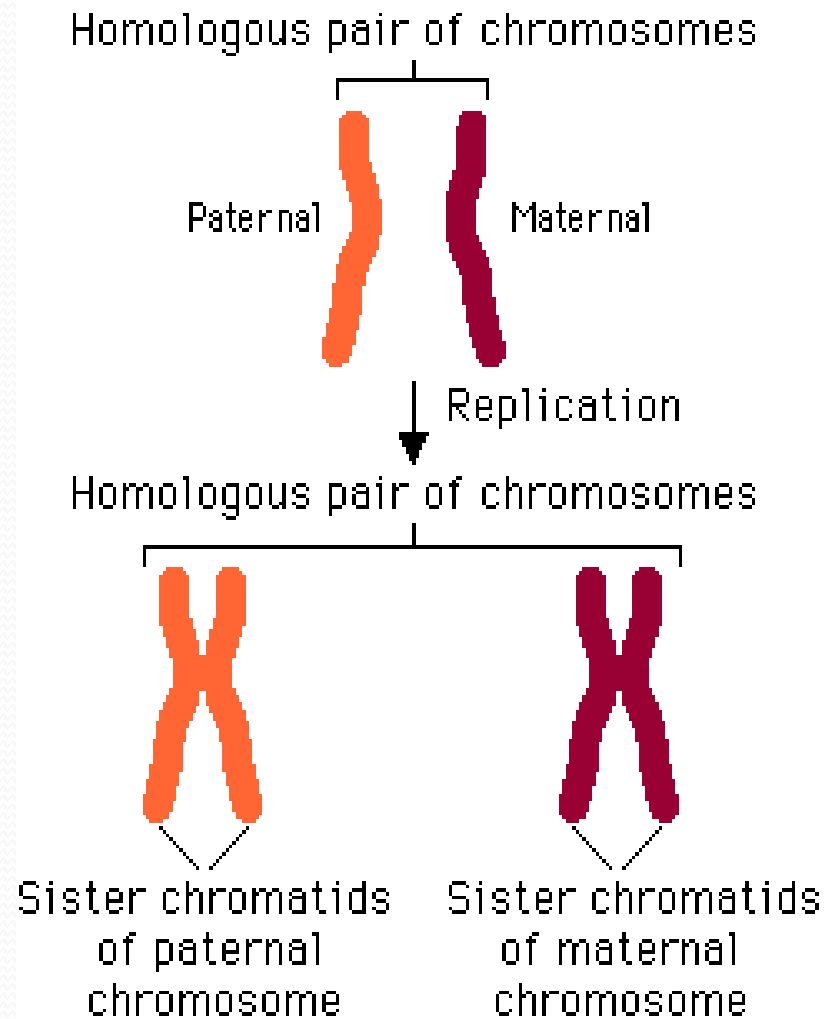
HOMOLOGOUS CHROMOSOMES

When examined under a light microscope generally they look very similar.

The exception to this rule is one pair of chromosomes called the **sex chromosomes, designated X and Y.**

In females the homologous pair of sex chromosomes consists of two large X chromosomes; in males the pair consists of an X and a much smaller Y chromosome

Note : If the sex pair is XX the individual is **genetically female** If the sex pair is XY the individual is genetically **male**

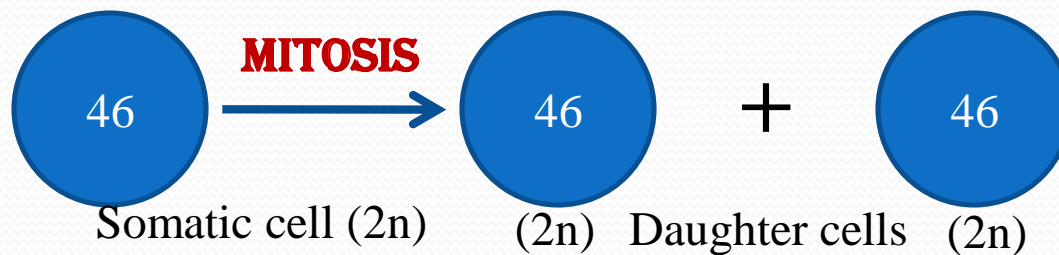


Where can we find somatic cells?

All the cells in the human body are somatic except the *sperm* and the *oocyte*

How they divide?

Somatic cells divide by **mitosis** for growth and to replace cells that die from tear and wear



**MITOSIS CONSERVES
CHROMOSOMES NUMBER**

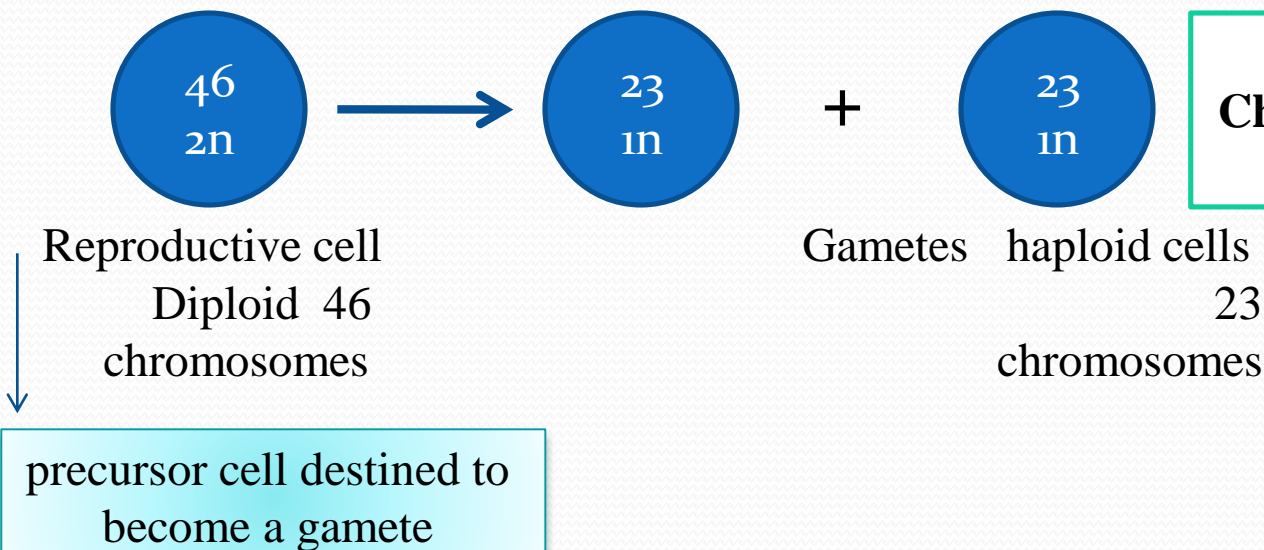
important

2- Reproductive cells (also called sex cells)

Reproductive cells develop in gonads (ovaries in female and testes in male)

They contain only **23** chromosomes that is why they are called **haploid** cells ($1n$)

Reproductive cells divide by **meiosis**



Meiosis does not conserve
Chromosomes number instead
It reduces it by half

important

Thus,

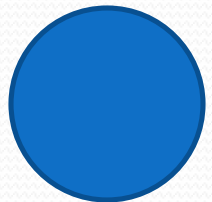
It is impossible for a female to reproduce here self simply because here sex cells (the **oocytes are haploid (23, 1n)**)

It is impossible for a male to reproduce him self simply because his sex cells (the **sperms are haploid (23, 1n)**)

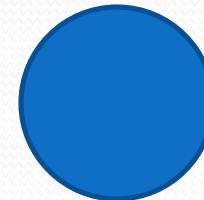
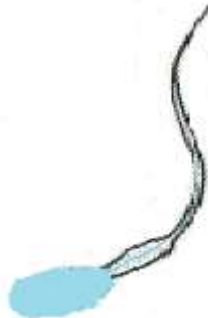
What to do?

Union

Fertilization



+



Oocyte
23 chromosomes
1n, haploid

Sperm
23 chromosomes
1n, haploid

Zygote
46 chromosomes
2n, diploid

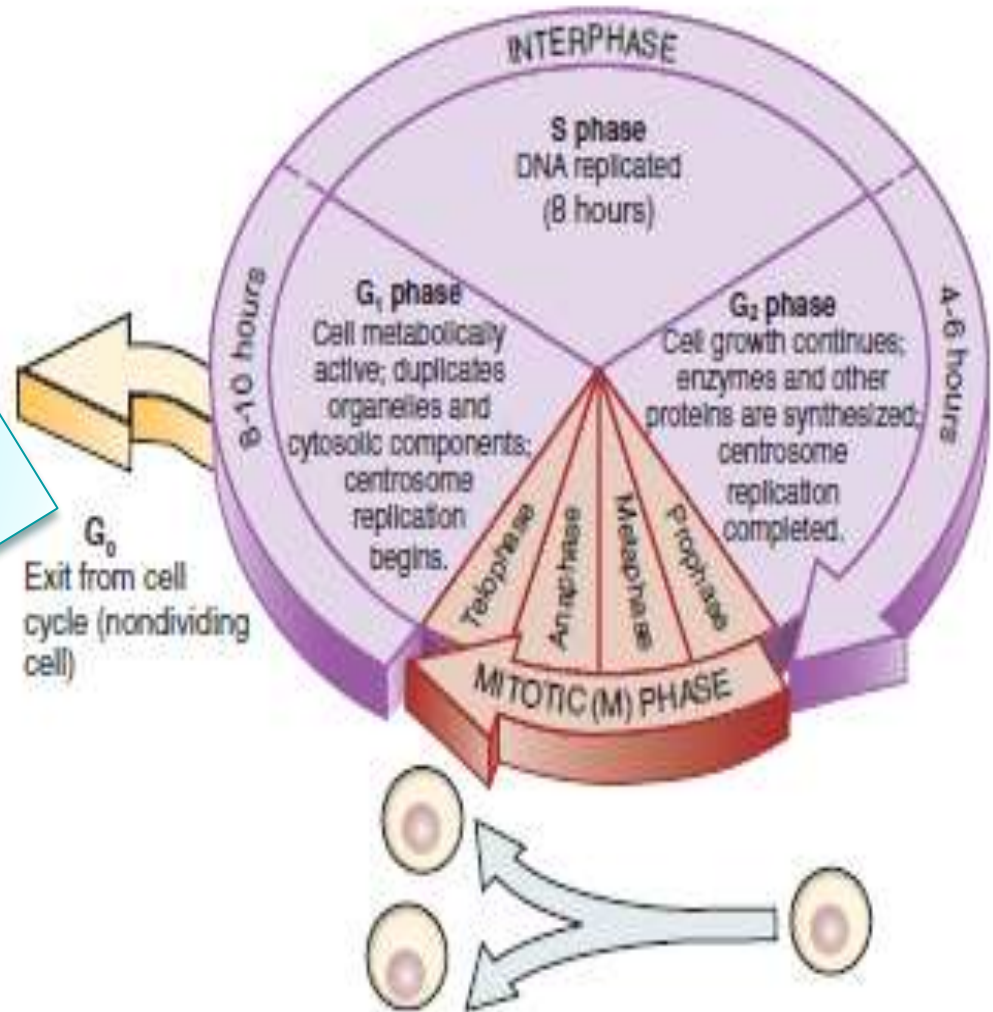
Cell Division

When a cell reproduces, it must replicate (duplicate) all
its
chromosomes to pass its genes to the next
generation of cells

The cell cycle consists of two major periods:

INTERPHASE, when a cell is not dividing,

and the **MITOTIC (M) PHASE**, when a cell is dividing





INTERPHASE

The Interphase

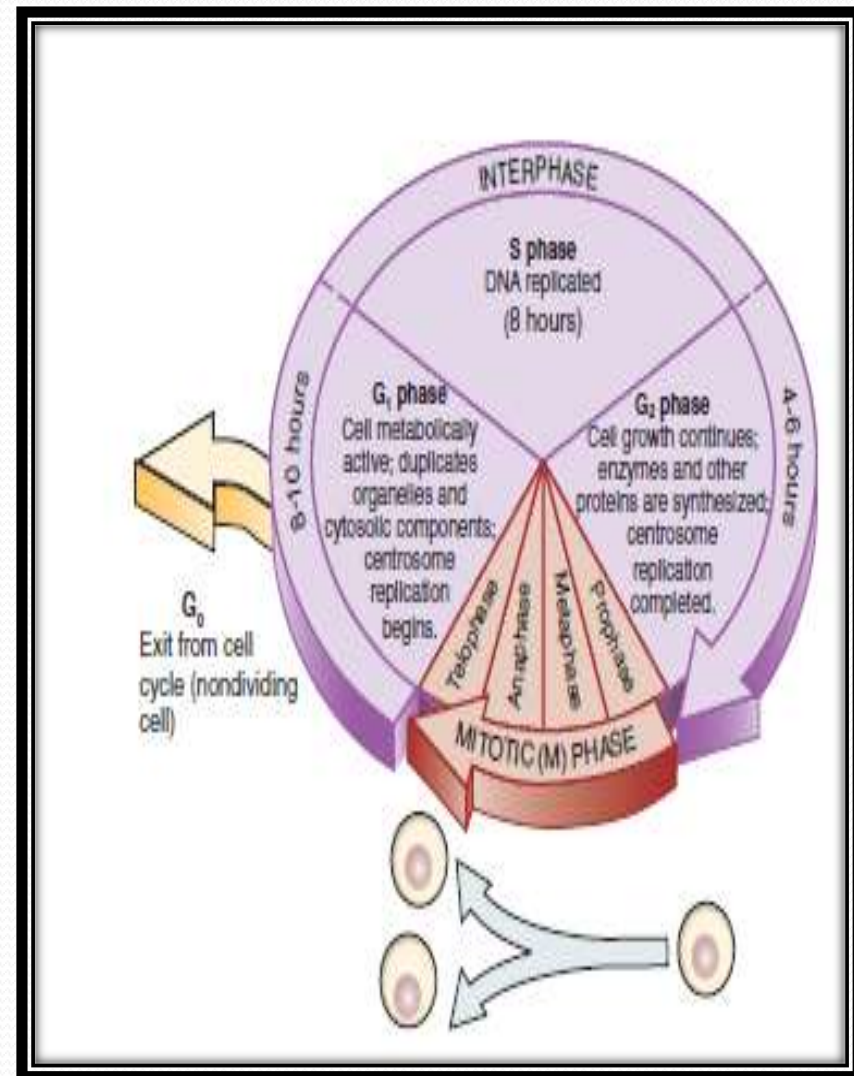
Interphase is a state of high metabolic activity; it is during this time that the cell does most of its growing.

During **interphase**

- 1- **The cell replicates its DNA**
- 2- Produces additional organelles and cytosolic components

Interphase consists of three phases:

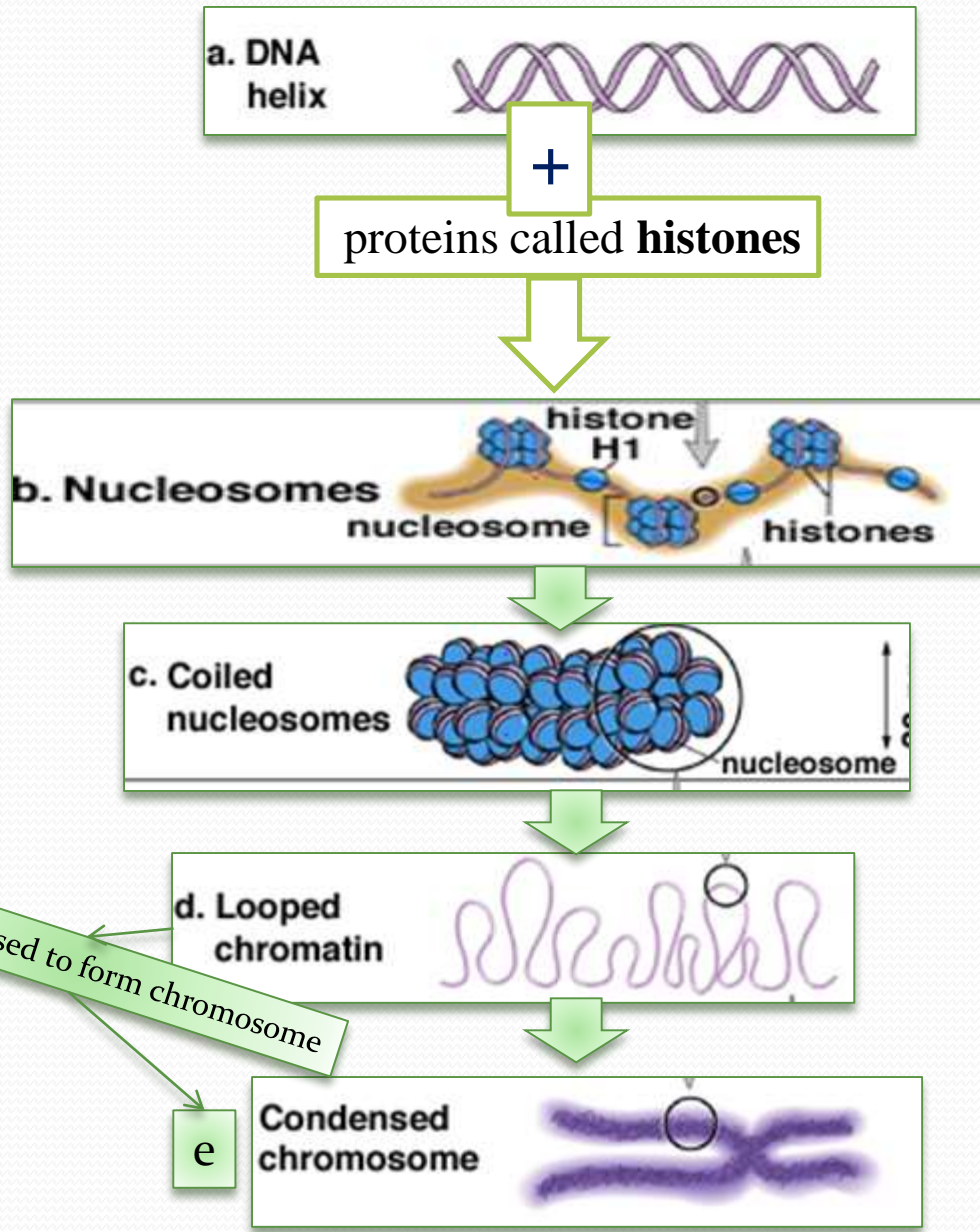
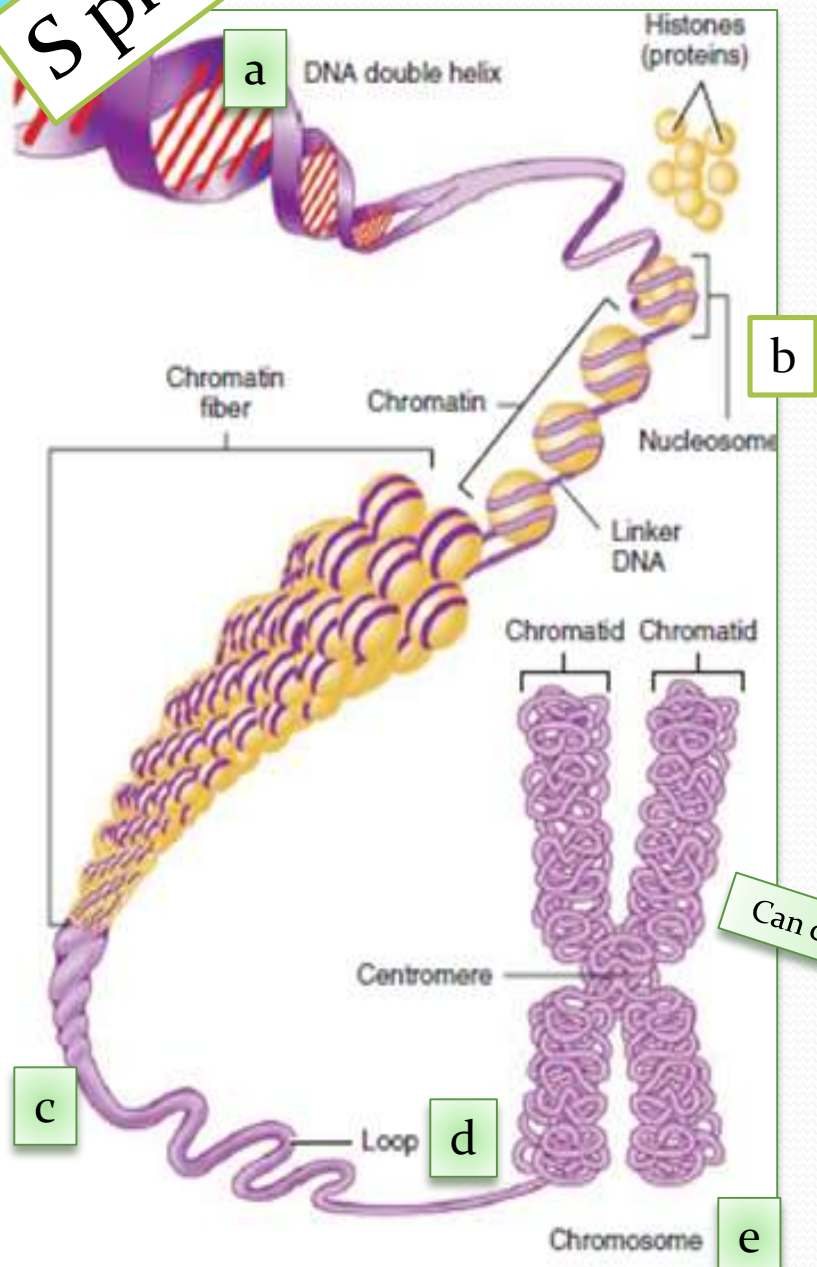
- 1- **G1 phase**
- 2- **S phase**
- 3- **G2 phase**



S phase

Cells before division undergo DNA synthesis during the Interphase

Chromosome's structure

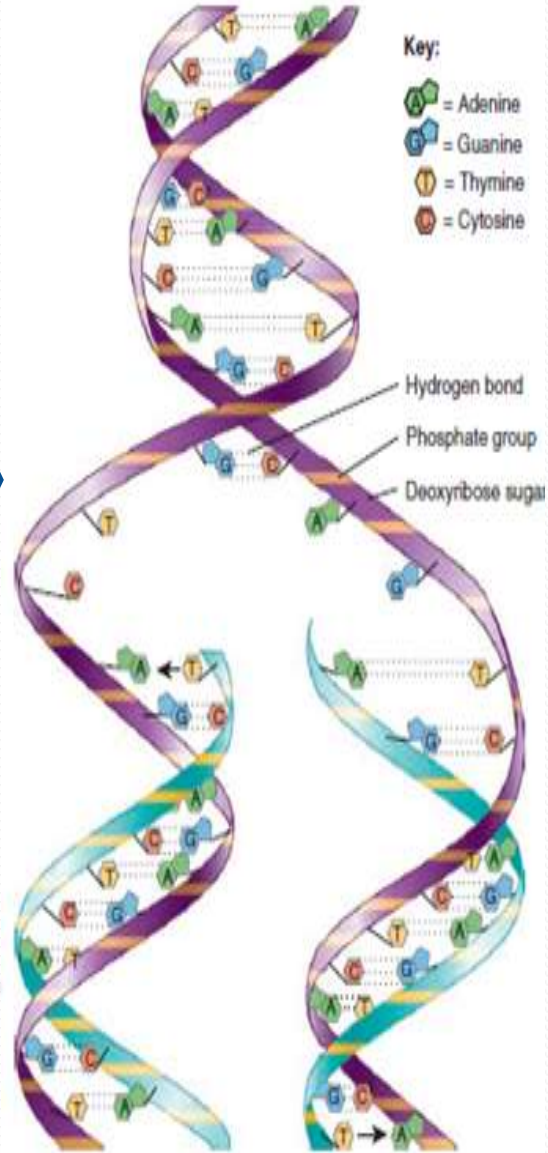


Interphase

DNA

replicates (duplicates)

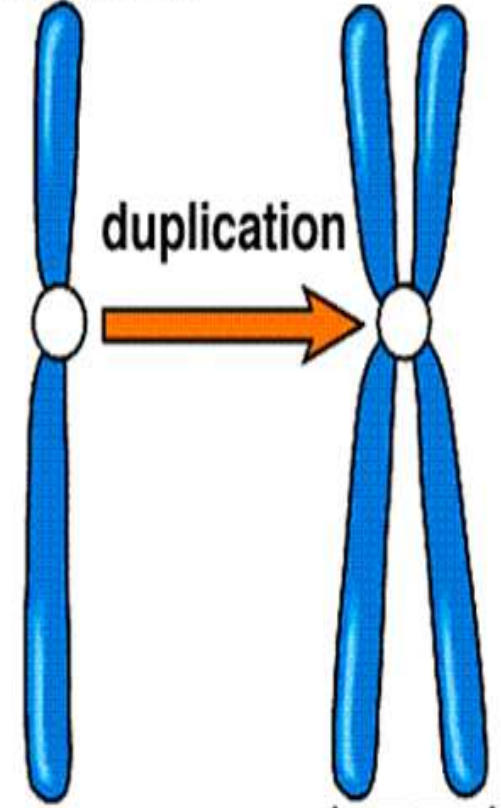
The **S** stands for **synthesis** of DNA.



Old strand New strand New strand Old strand

Can condensed to form chromosome

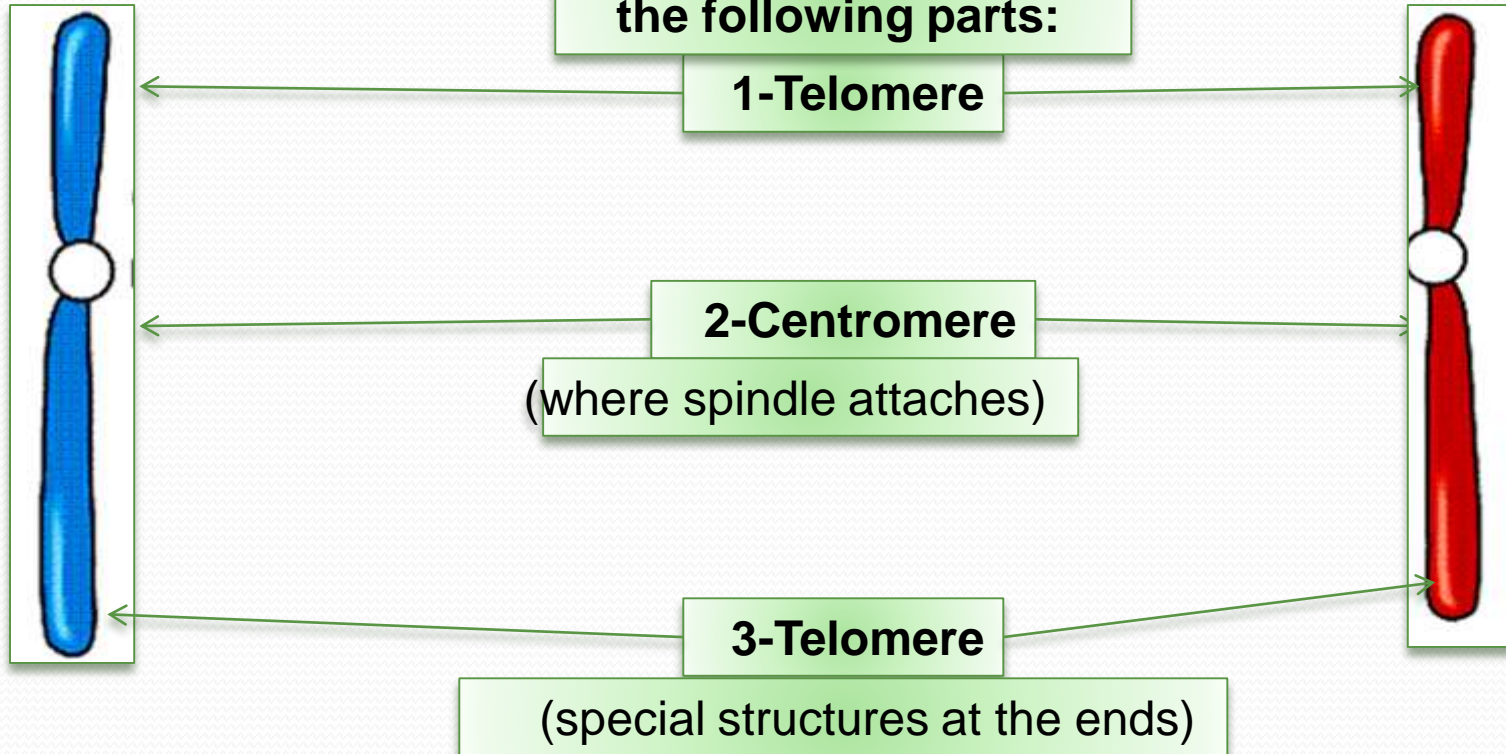
chromosome with one chromatid



sister chromatids

The chromatin of nucleus condense into a **chromosome**

Each chromosome has the following parts:

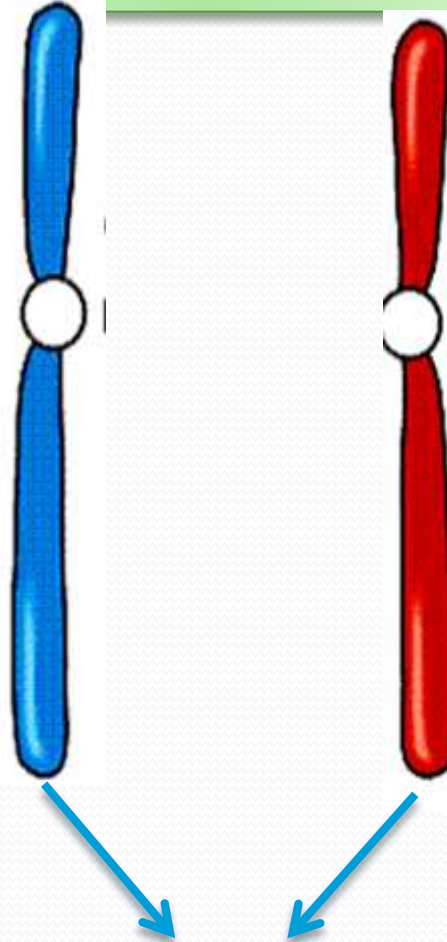


depending on the stage of the cell cycle chromosomes come in 2 forms:

1- The monad form consists of a single chromatid, a single piece of DNA containing a centromere and telomeres at the ends.

2- The dyad form consists of 2 identical chromatids (sister chromatids) attached together at the centromere

The chromatin of nucleus condense into a **chromosome**



One chromosome coming from the father called paternal

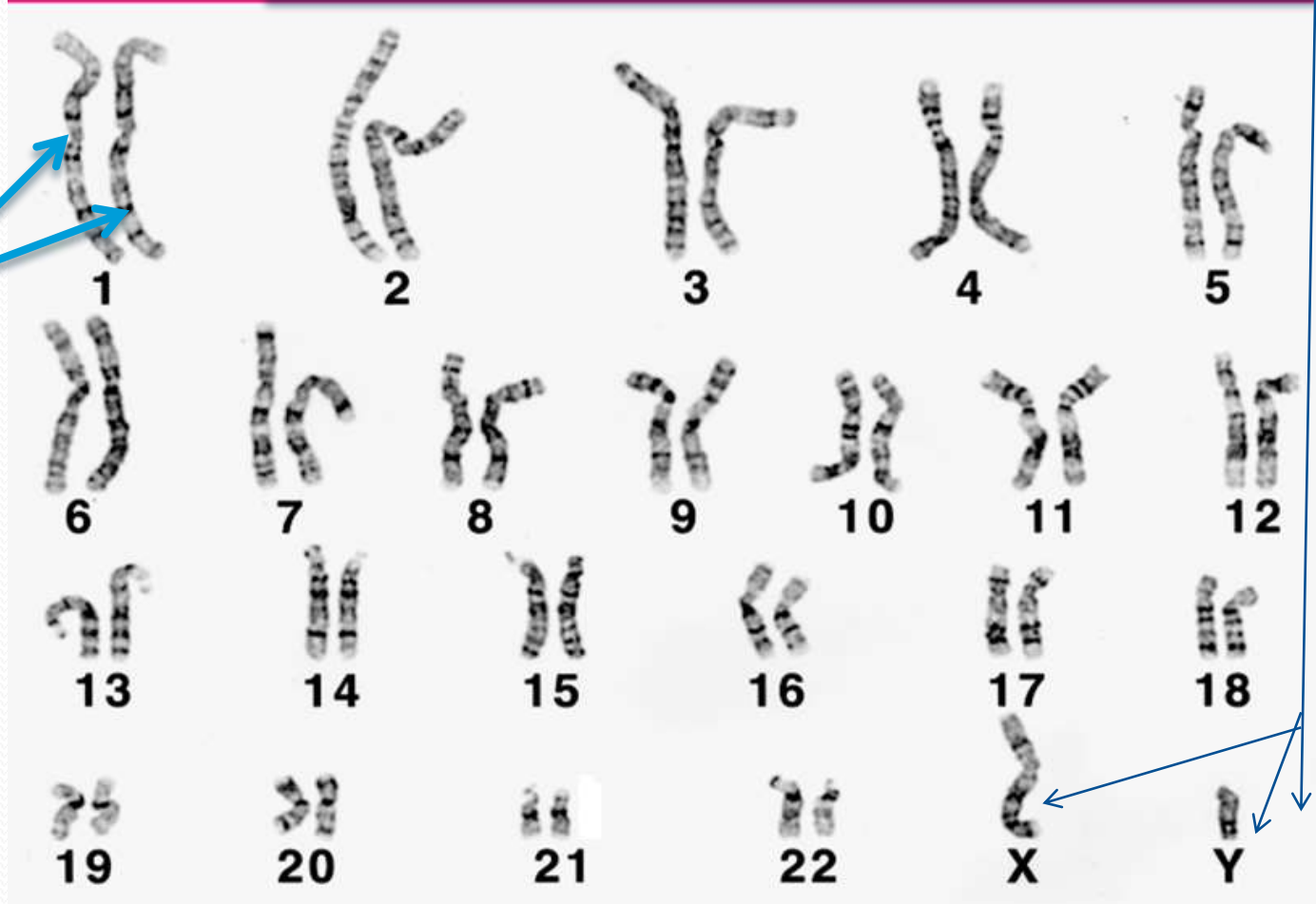
One chromosome coming from the mother called maternal

HOMOLOGOUS CHROMOSOMES

HOMOLOGOUS CHROMOSOMES

Notice that chromosomes number 23 are not homologous, what does this mean?

Notice that in this picture
There are two chromosomes
Numbered 1 and
etc. These
Chromosomes are
called homologous
chromosomes; one
comes from the
mother and the
other comes from
the father
During
fertilization



Picture of the 46 chromosomes (23 pairs of chromosomes)

G phases

Because the G phases are periods when there is no activity related to DNA duplication, they are thought of as gaps or interruptions in DNA duplication.

The G1 phase is the interval between the mitotic phase and the S phase.

During G1,
the cell replicates most of its
organelles and cytosolic components
but
not its DNA.

➤ Replication of centrosomes also begins in the G1 phase.


For a cell with a total cell cycle time of 24 hours, G1 lasts 8 to 10 hours. However, the duration of this phase is quite variable. It is very short in many embryonic cells or cancer cells. Cells that remain in G1 for a very long time, perhaps destined never to divide again, are said to be **in the G0 phase.**

Most nerve cells
are in
the G0 phase. Once a cell enters the S phase, however, it is committed to go through the rest of the cell cycle.

The G2

phase is the interval between the S phase and the mitotic phase. It lasts 4 to 6 hours.

During G2, cell growth continues, enzymes and other proteins are synthesized in preparation for cell division, and replication of centrosomes is completed.



MITOTIC (M) PHASE

Cell Division

Somatic cells

by

Mitosis

Reproductive cells

by

Meiosis

To be discussed later

Cell Cycle

Interphase
(Cell is not dividing)

Mitotic phase
(Cell is dividing)

Consists of four stages:

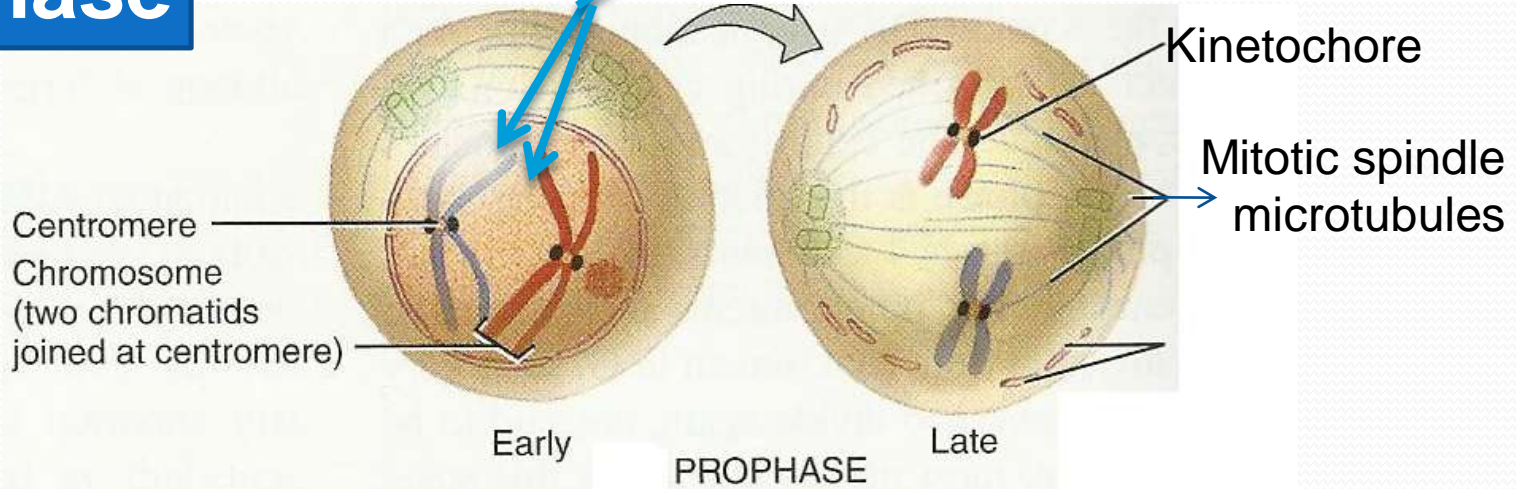
- 1-Prophase
- 2-Metophase
- 3-Anaphase
- 4-Telophase

Consist of three phases:
1- The G1 phase
2- The S phase
3- The G2 phase

Prophase

2n

it has two sets of Chromosomes (two copies)
One chromosome is paternal and the other maternal



Centromere (a constricted region holds the chromatid pair together)

Outside of each centromere is a protein complex called Kinetochores

Later in prophase tubulins in the pericentriolar material of the **centrosomes** start to form the mitotic spindle

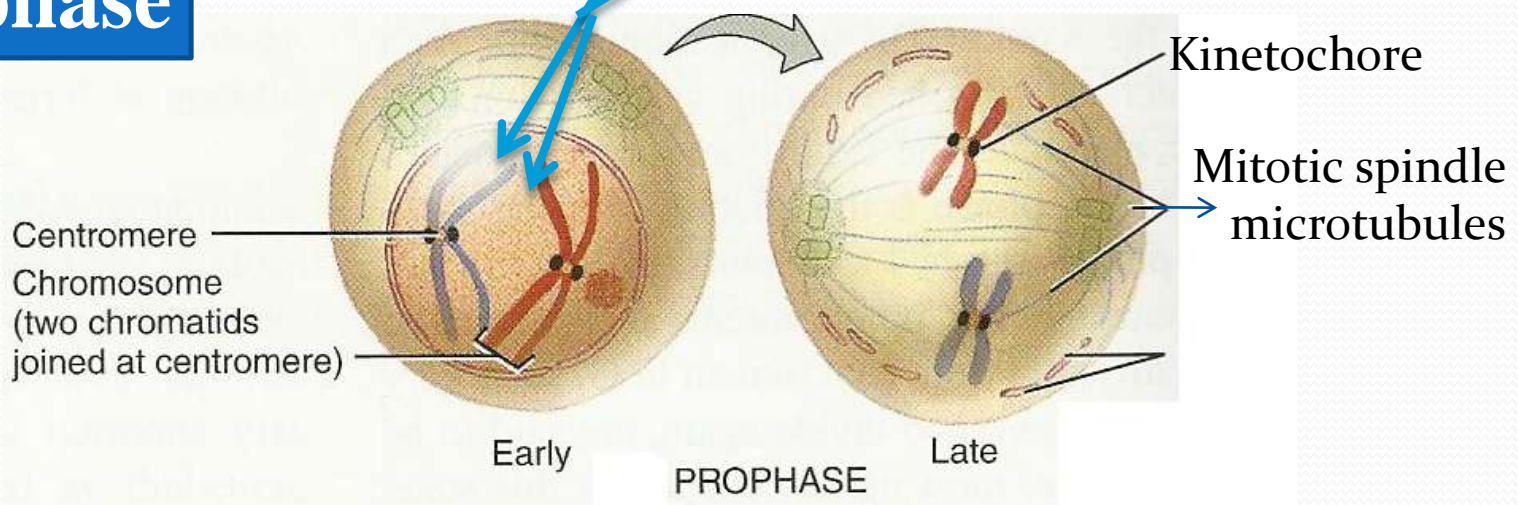
The mitotic spindle **attaches** to the Kinetochores

The mitotic spindle (microtubules) lengthen they push the centrosomes to the poles

it has two sets of
Chromosomes (two copies)
One chromosome is
paternal and the other maternal

2n

Prophase



Centromere (a constricted region holds the chromatid pair together)

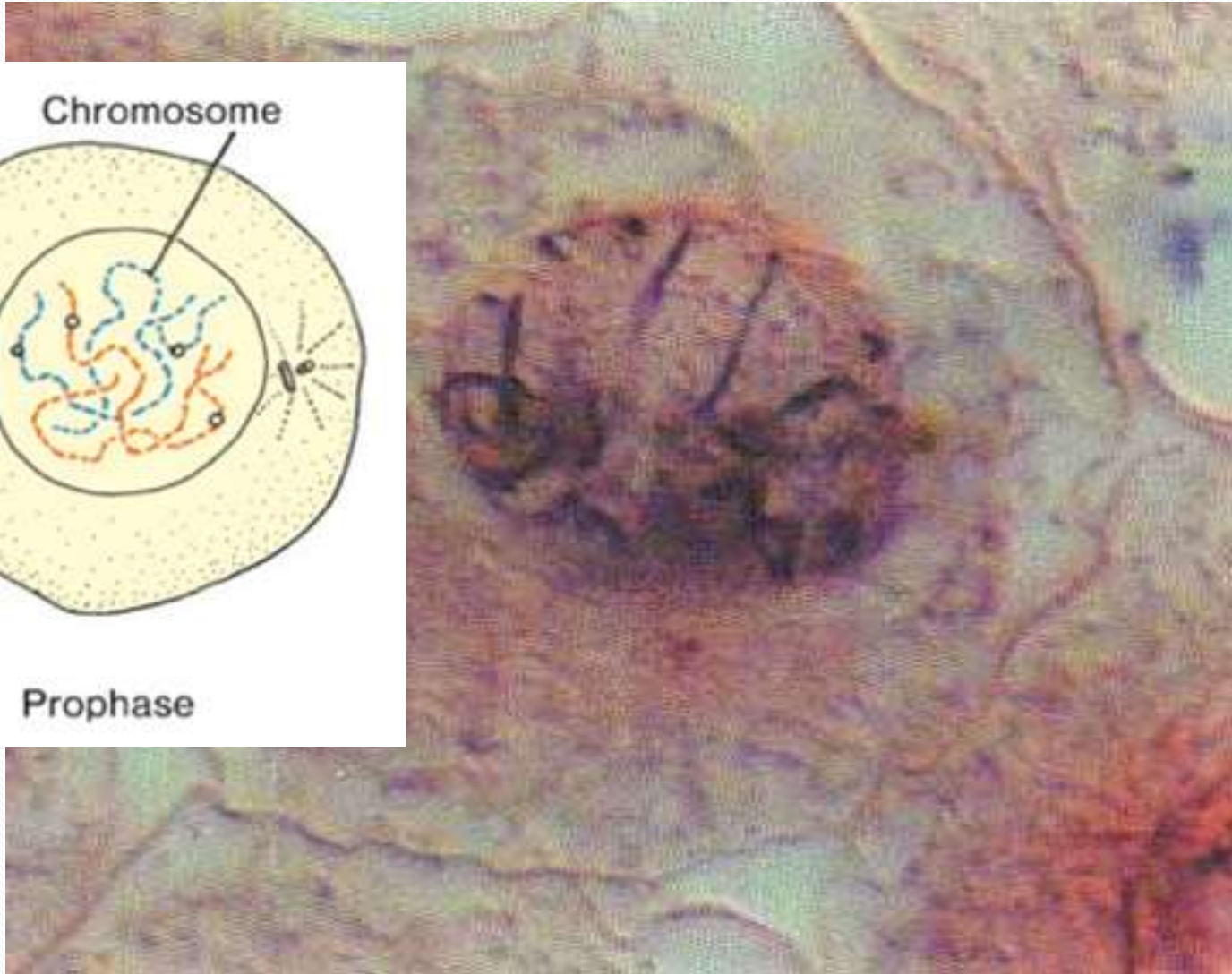
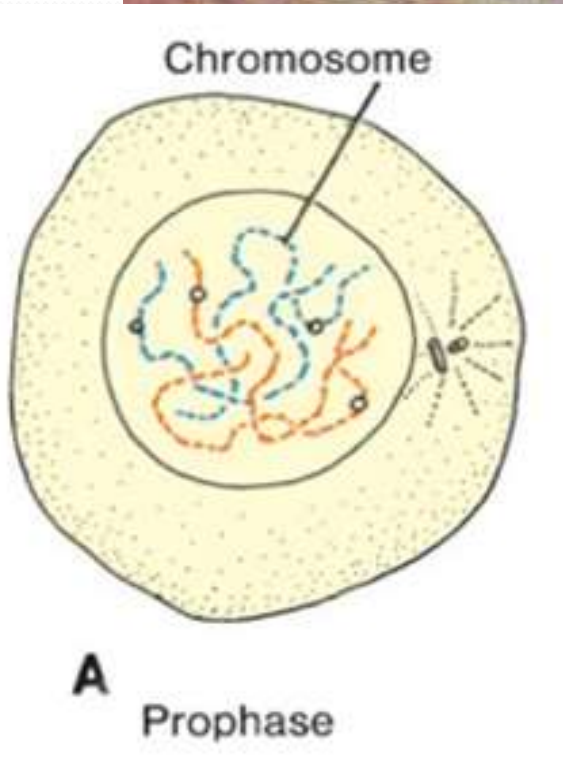
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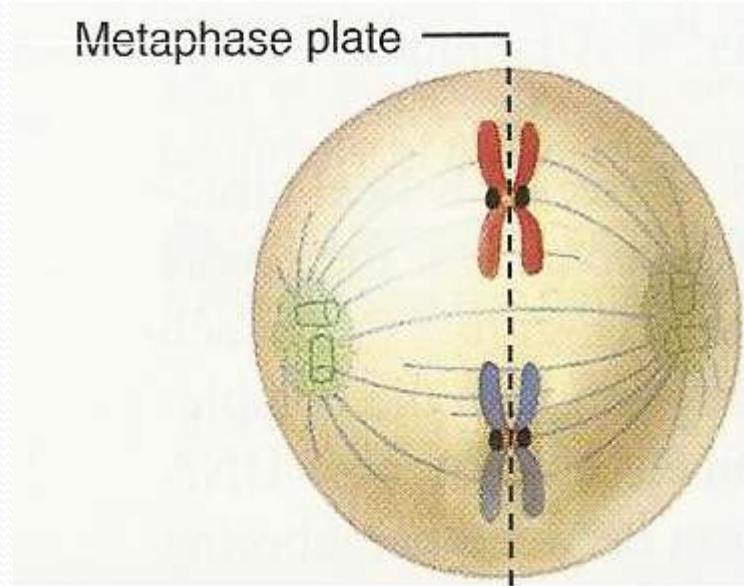
the mitotic spindle **attaches** to the Kinetochores

As the mitotic spindle (microtubules) lengthen they push the centrosomes to the poles

Mitosis Prophase



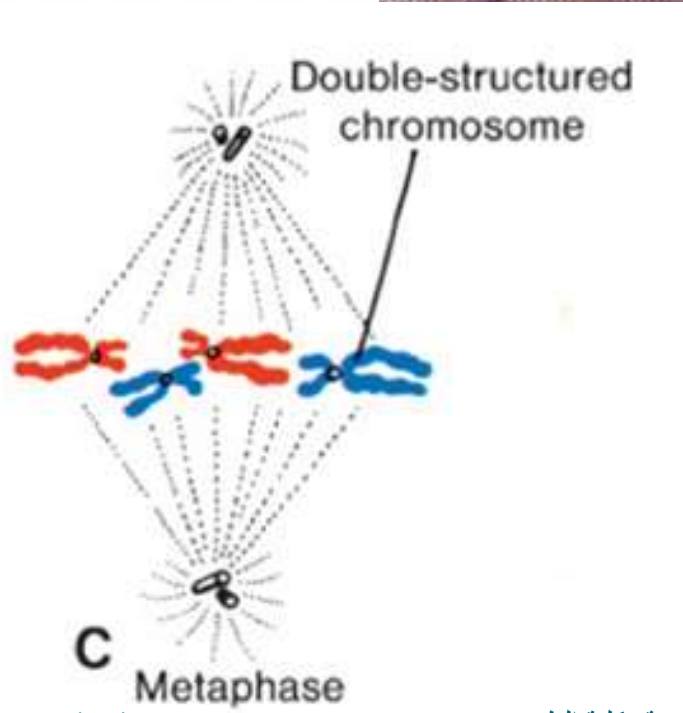
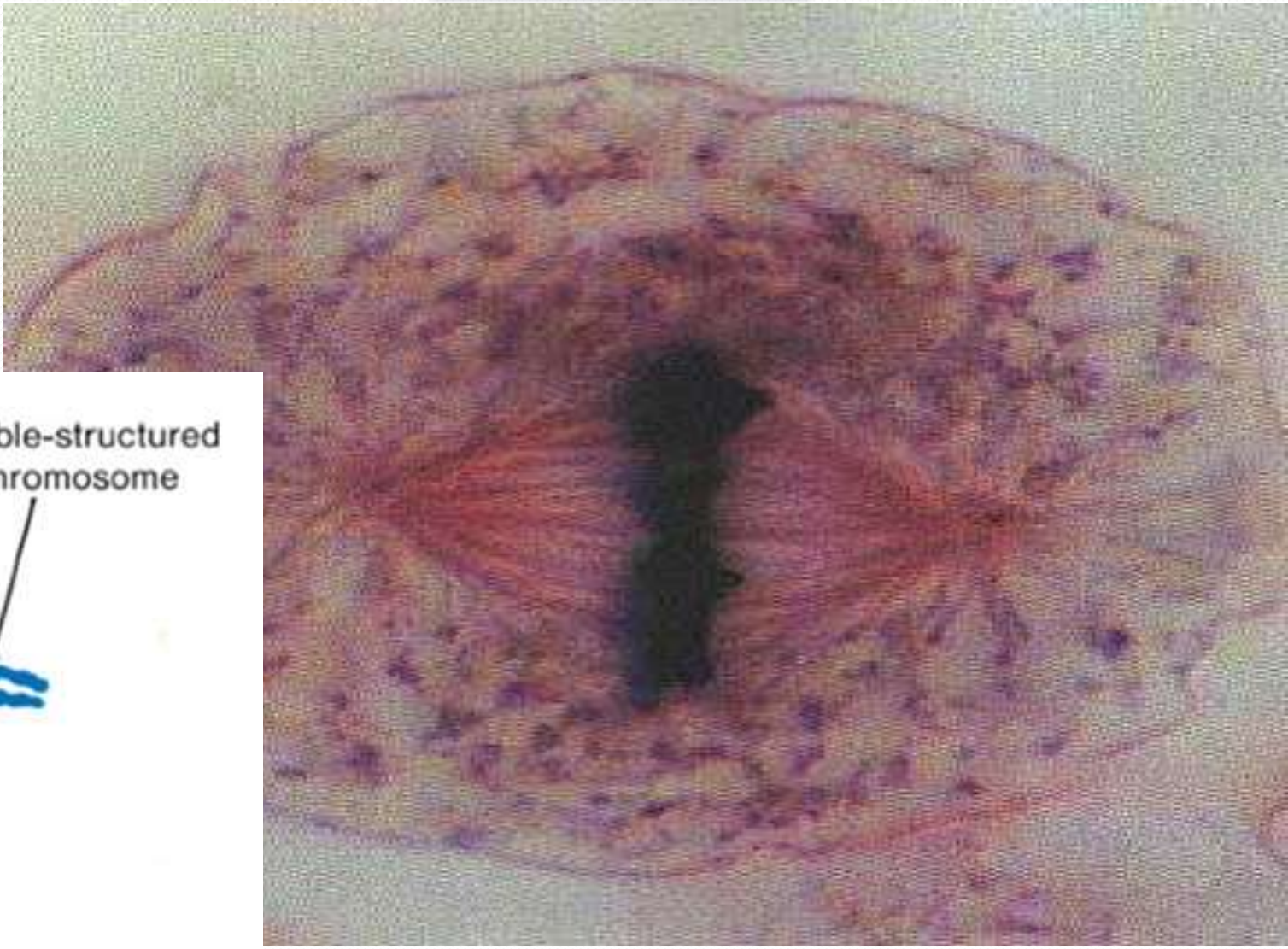
METAPHASE



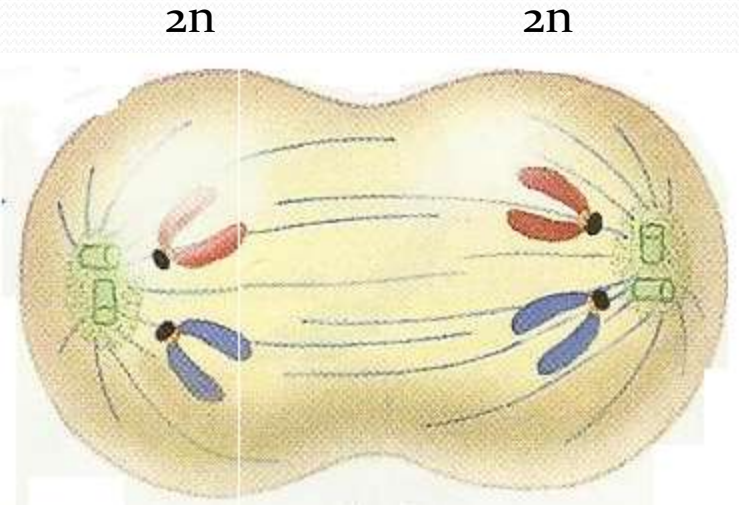
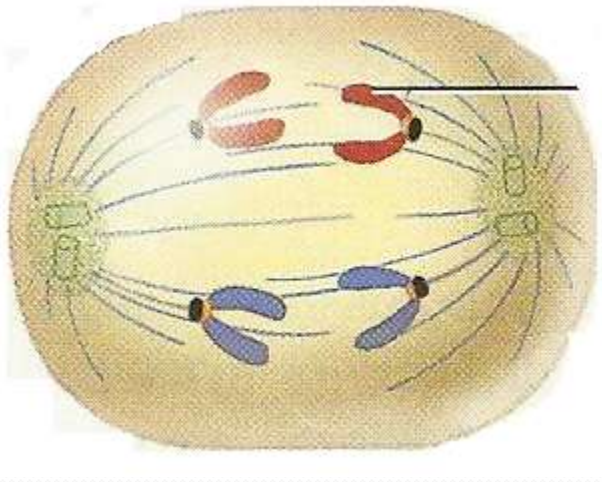
The Kinetochore microtubules align the centromeres at the exact center of the mitotic spindle

This midpoint region called **metaphase plate**

Mitosis Metaphase



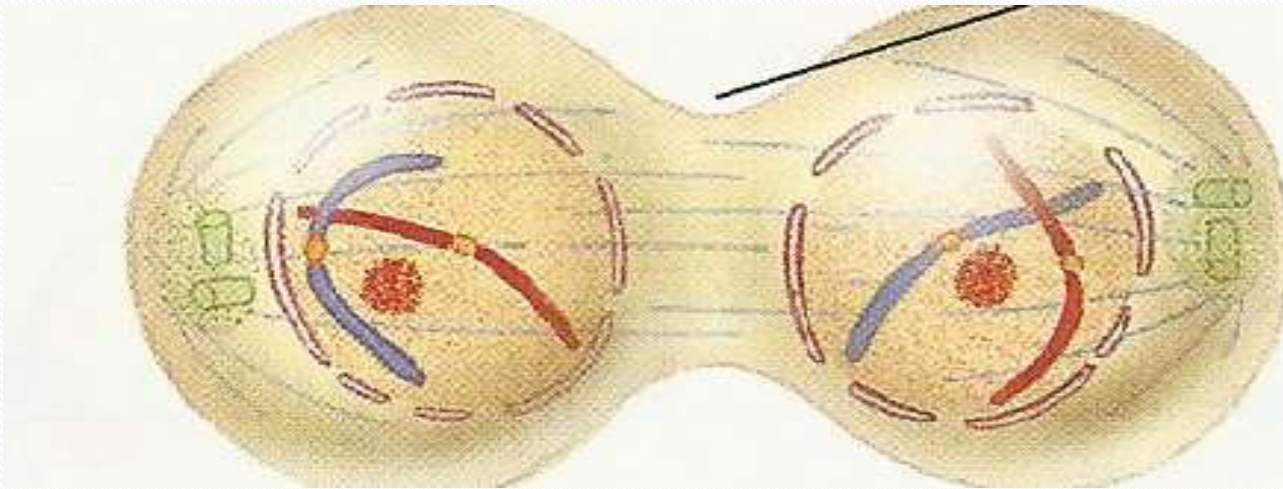
ANAPHASE



The **centromeres split** leading to the **separation** of the two members of the chromatid pair
once separated the chromatids are termed chromosomes



TELOPHASE

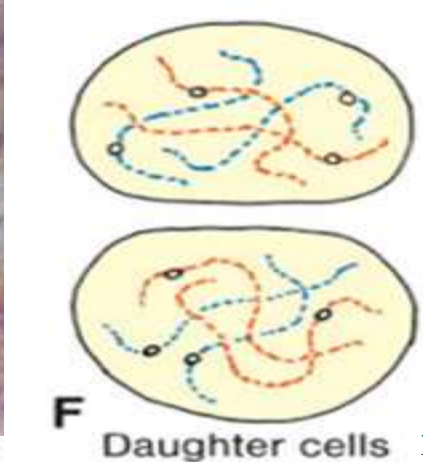
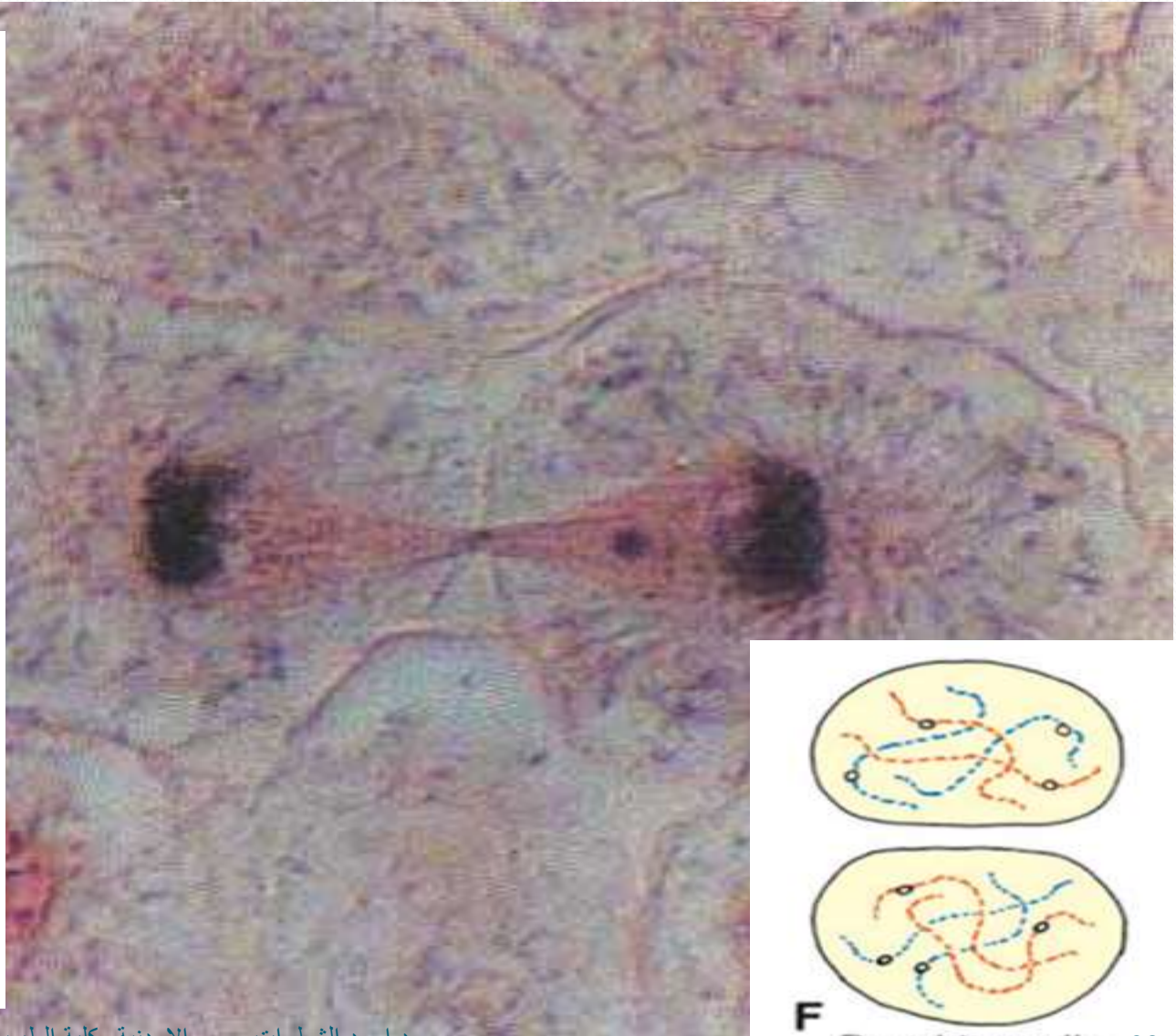
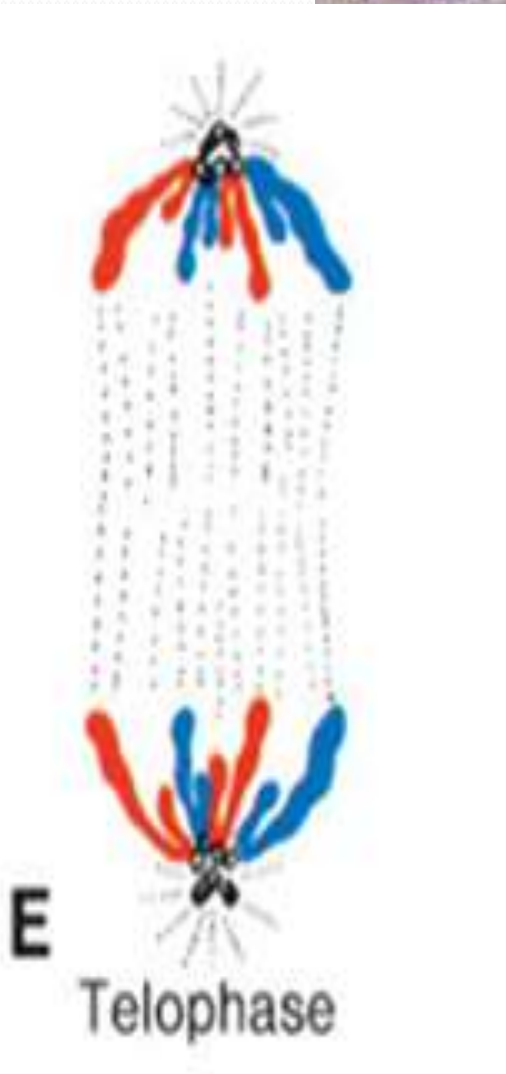


The identical sets of chromosomes now at apposite poles of the cell

A nuclear envelope forms around each chromatin mass

The mitotic spindle disappears

Mitosis Telophase

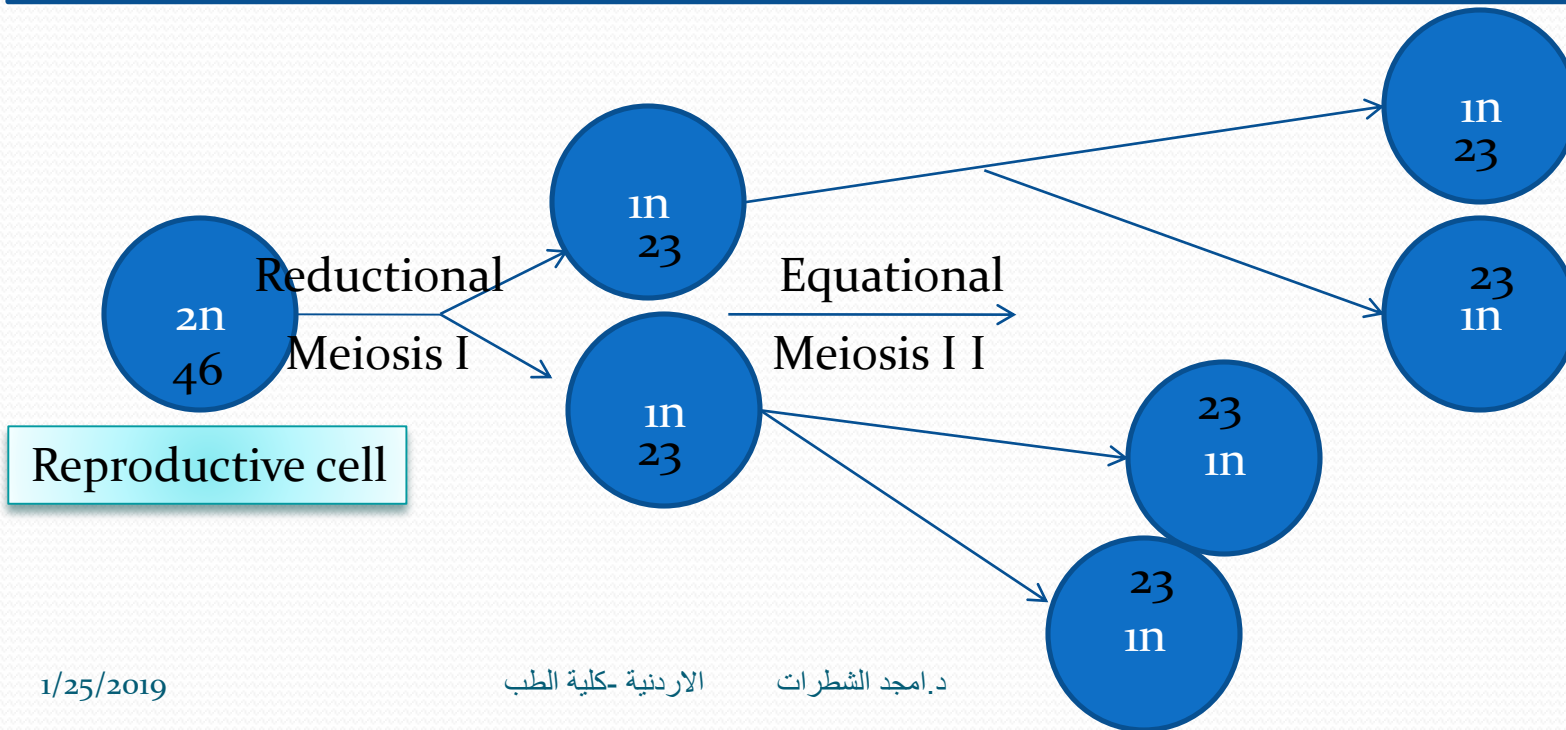


Meiosis

Meiosis occurs in two successive stages :

Meiosis I (also known as reductional meiosis) which deals with the number of chromosomes it halves the number of chromosomes

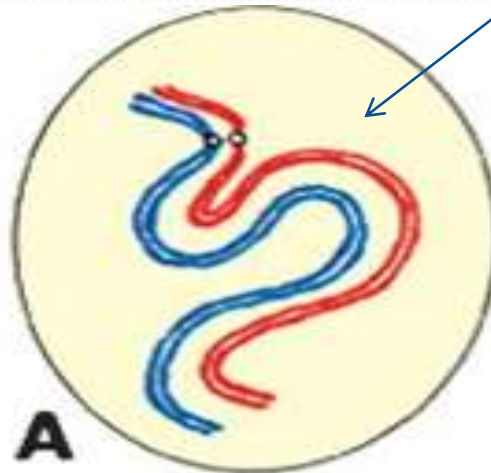
Meiosis II (also known as equational meiosis) which deals with the conditions of chromosomes



Meiosis I is generally divided into four stages:

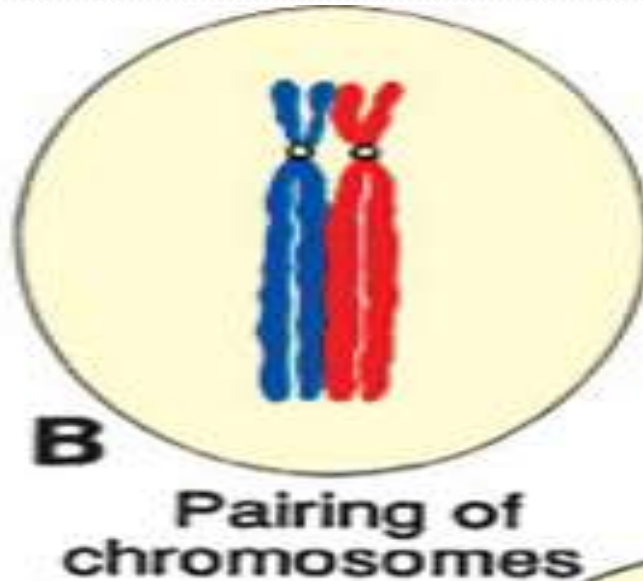
1-Prophase
2-metaphase
3-
Anaphase

1-Prophase is running into stages
*A- **LEPTOTEN** stage, (lepto means long)*
In this stage chromosomes are elongated and
extended and become gradually visible



B- ZYGOTEN stage, (*zygo* means joined)

In this stage identical chromosomes pair up together (synapsis)



C- PACHYTENE stage, (*pachy* means short)

In this stage chromosomes become shorter and more condensed

D- DIPLOTENE stage,

Chromosomes come together and cross each other by certain segments of their bodies forming what we called **CHIASMATA: X- shaped structure**

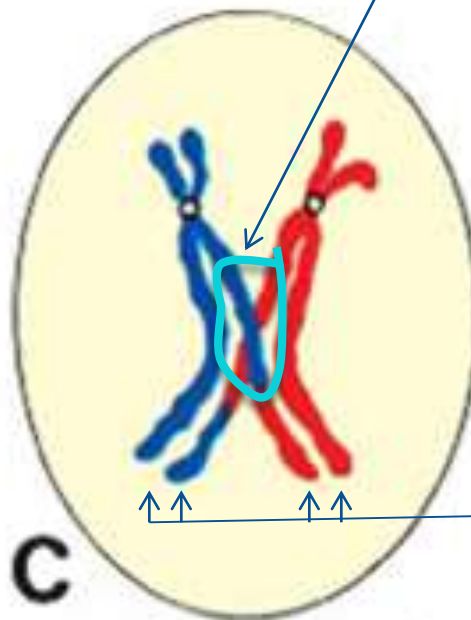
Formed by the junction of two chromatids of the for chromatids (**tetrad**)

In Prophase I

Crossing over of non-sister chromatids

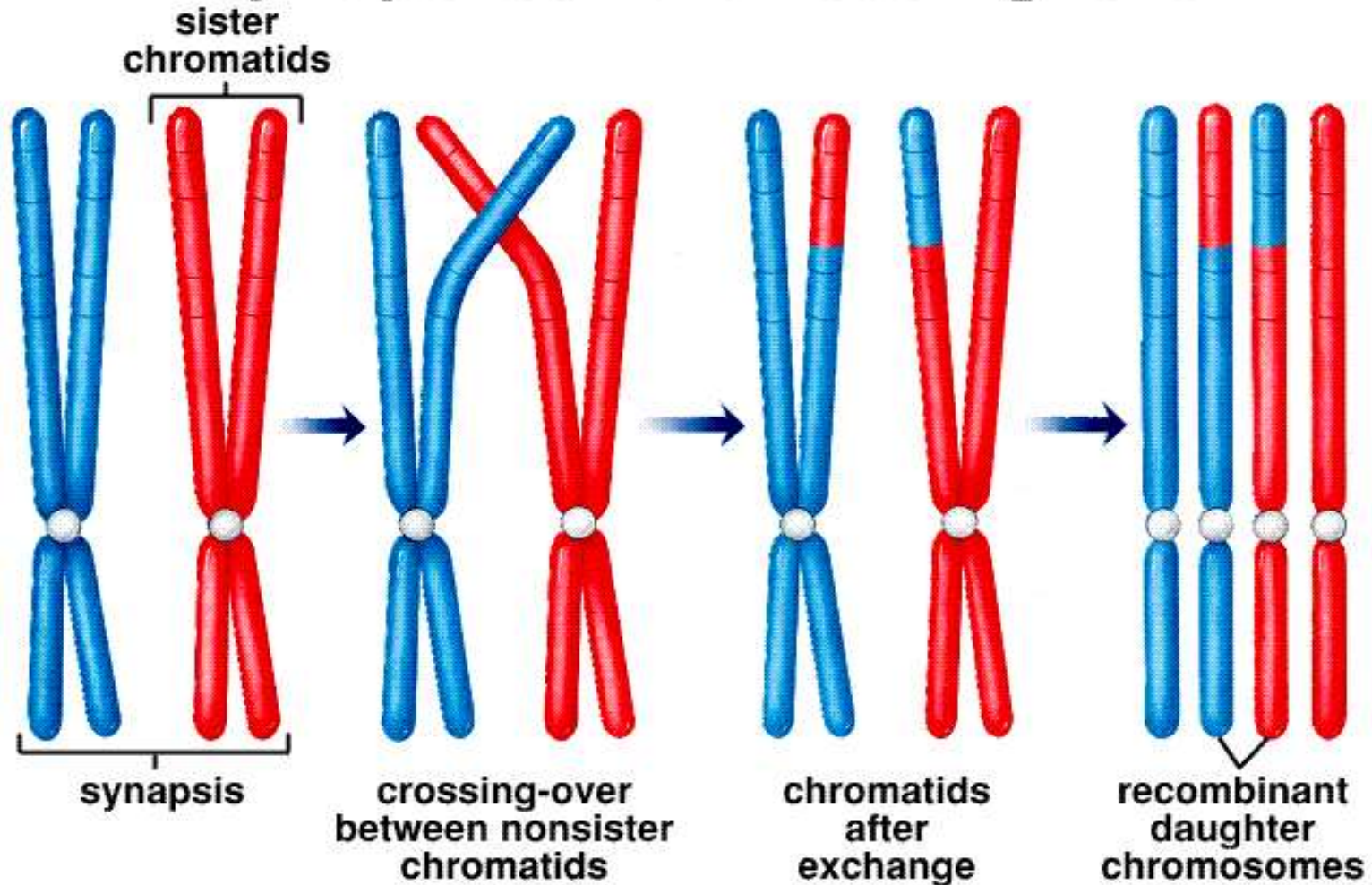
During prophase I, non-sister chromatids can undergo synapsis, in which the chromatids line up side-by-side & exchange genetic information between them

This allows new combination of genetic material which will become part of a new offspring

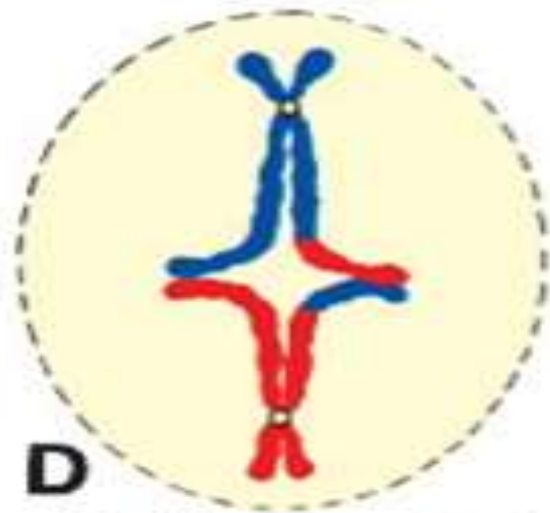


C
Chiasma formation

Synapsis and Crossing-over

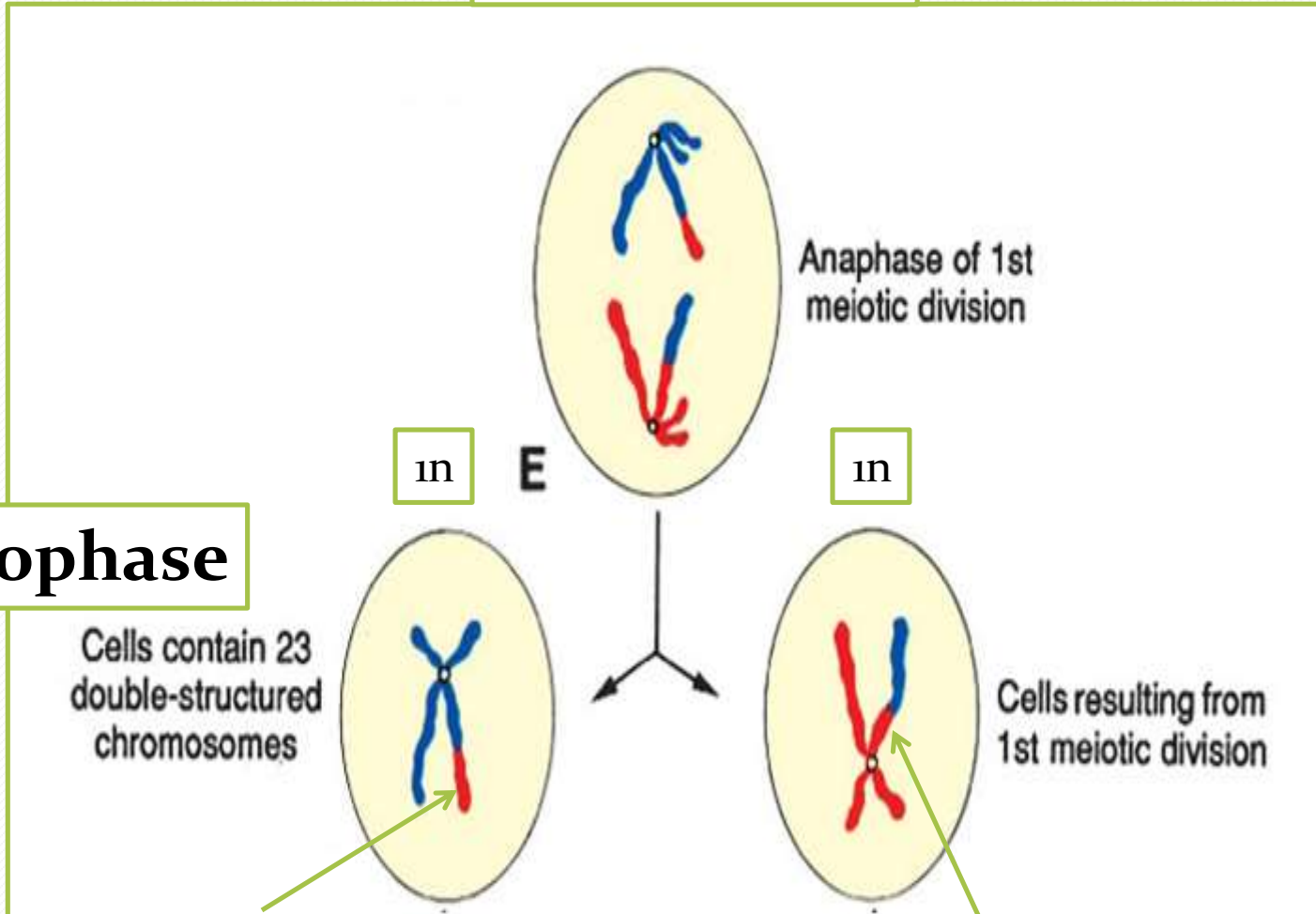


2-Metaphase



D
Pulling apart of
double-structured
chromosomes

3-Anaphase



Anaphase of 1st meiotic division

n

E

n

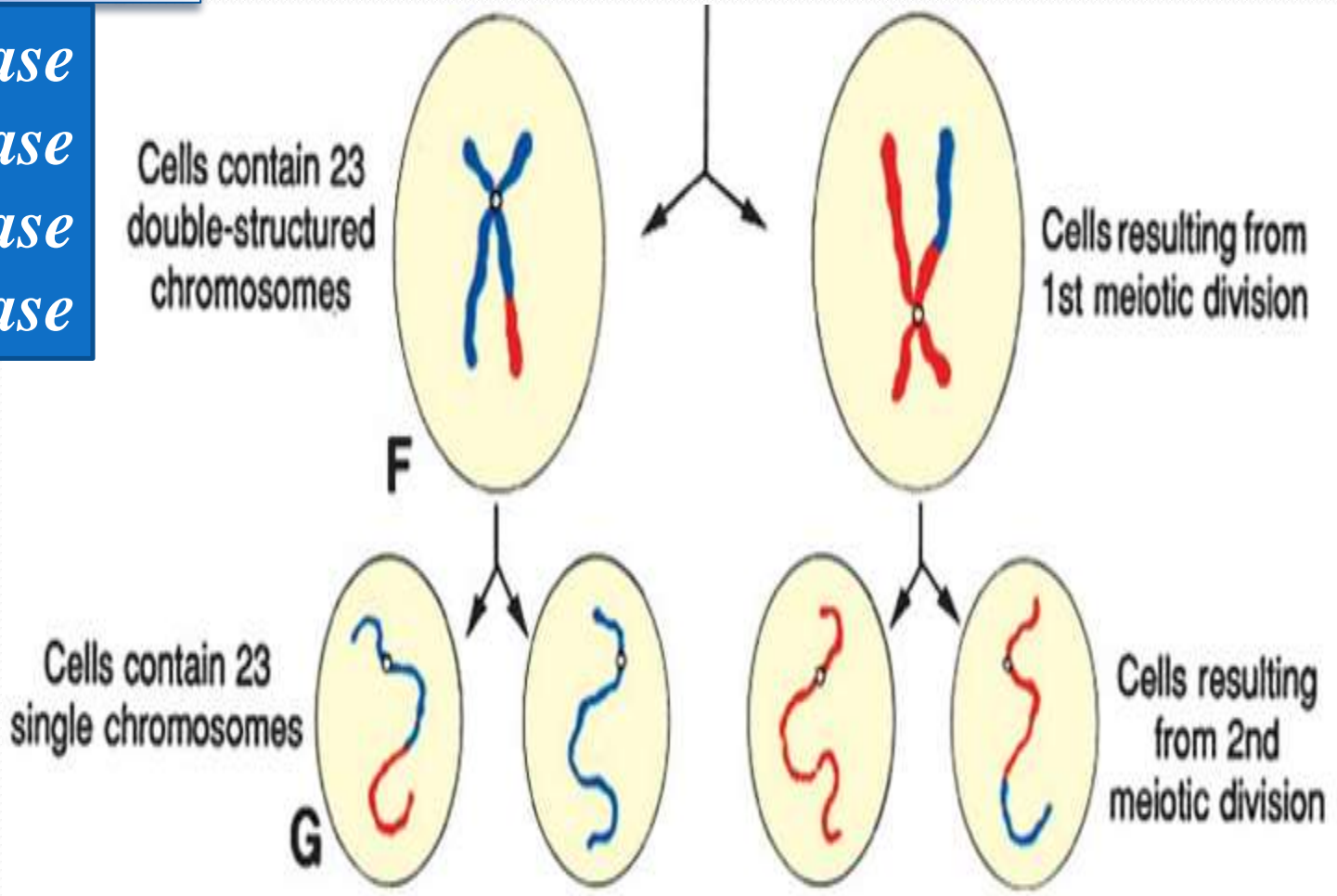
4-Telophase

Cells contain 23 double-structured chromosomes

Cells resulting from 1st meiotic division

Meiosis II runs into 4 stages:

- 1-Prophase*
- 2-Metophase*
- 3-Anaphase*
- 4-Telophase*



Meiosis I
Pairing of chromosomes
Homologous chromosomes separate
Daughter cells are haploid

Mitosis
No pairing
Sister chromatids separate,
Daughter cells are diploid

Chromosomal abnormalities

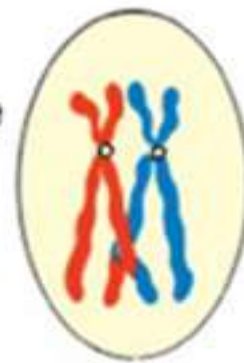
may be

numerical or structural

Abnormalities in chromosome number may originate during meiotic or mitotic divisions.

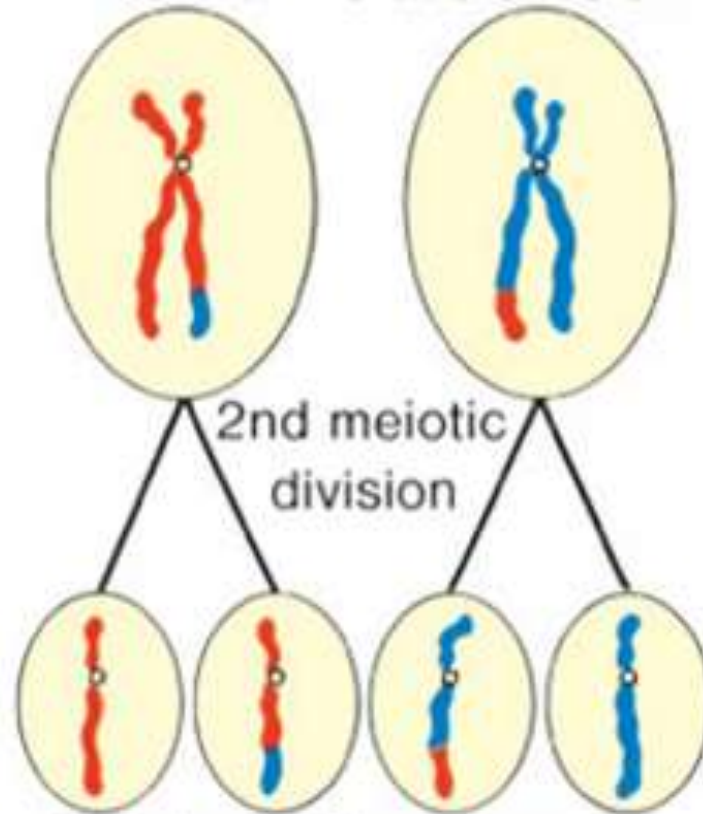
In meiosis, two members of a pair of homologous chromosomes normally separate during the first meiotic division so that each daughter cell receives one member of each pair

Primary oocyte or spermatocyte after DNA duplication
46 double-structured chromosomes



Normal

Normal meiotic division



23 single chromosomes

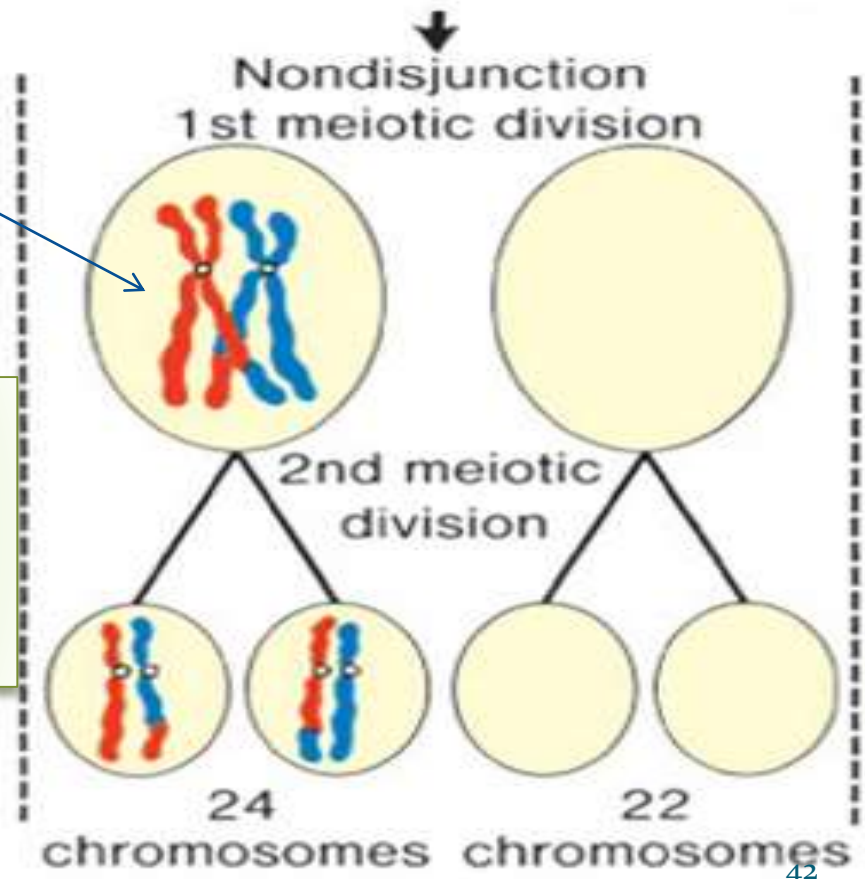
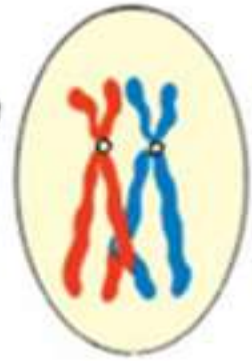
Sometimes, however, separation does not occur
(nondisjunction)

Both
members of a pair move
into one cell .

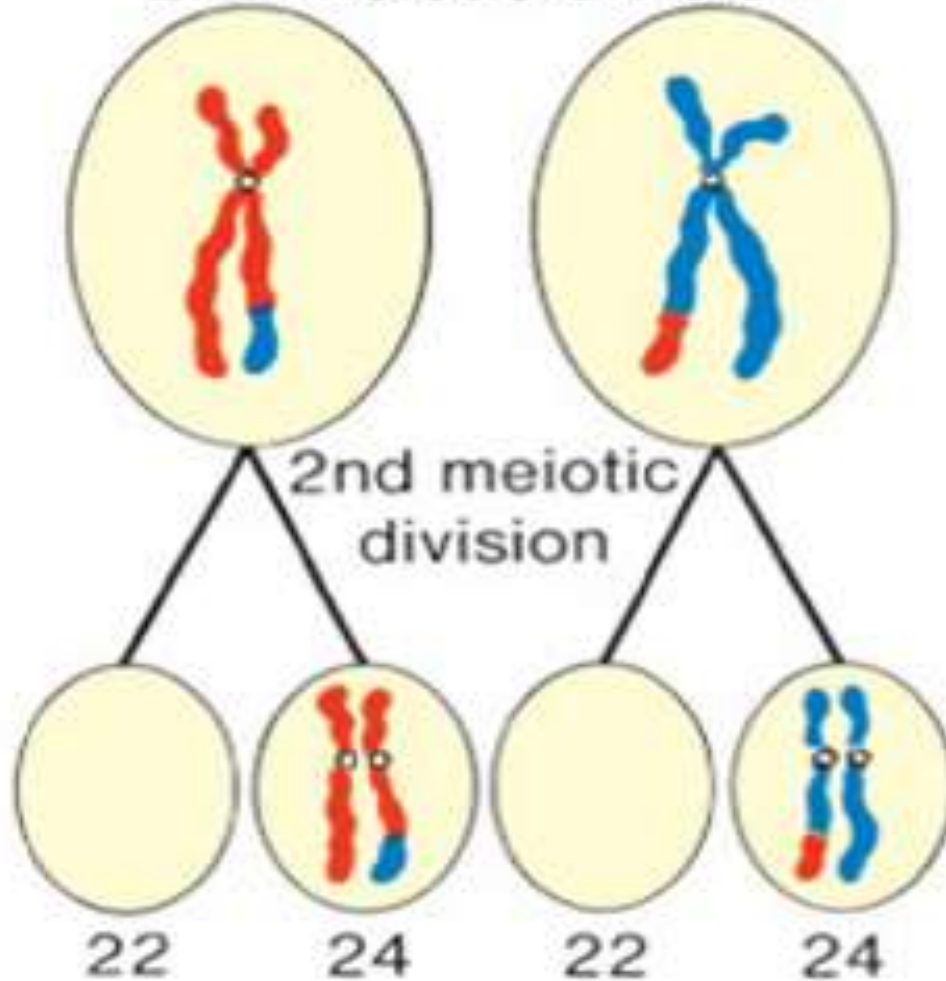


As a result of
nondisjunction of the chromosomes, one
cell receives 24 chromosomes,
and the other receives 22 instead of the
normal 23.

Primary oocyte or spermatocyte
after DNA duplication
46 double-structured
chromosomes



Nondisjunction 2nd meiotic division



Translocations

Sometimes chromosomes break, and pieces of one chromosome attach to another.

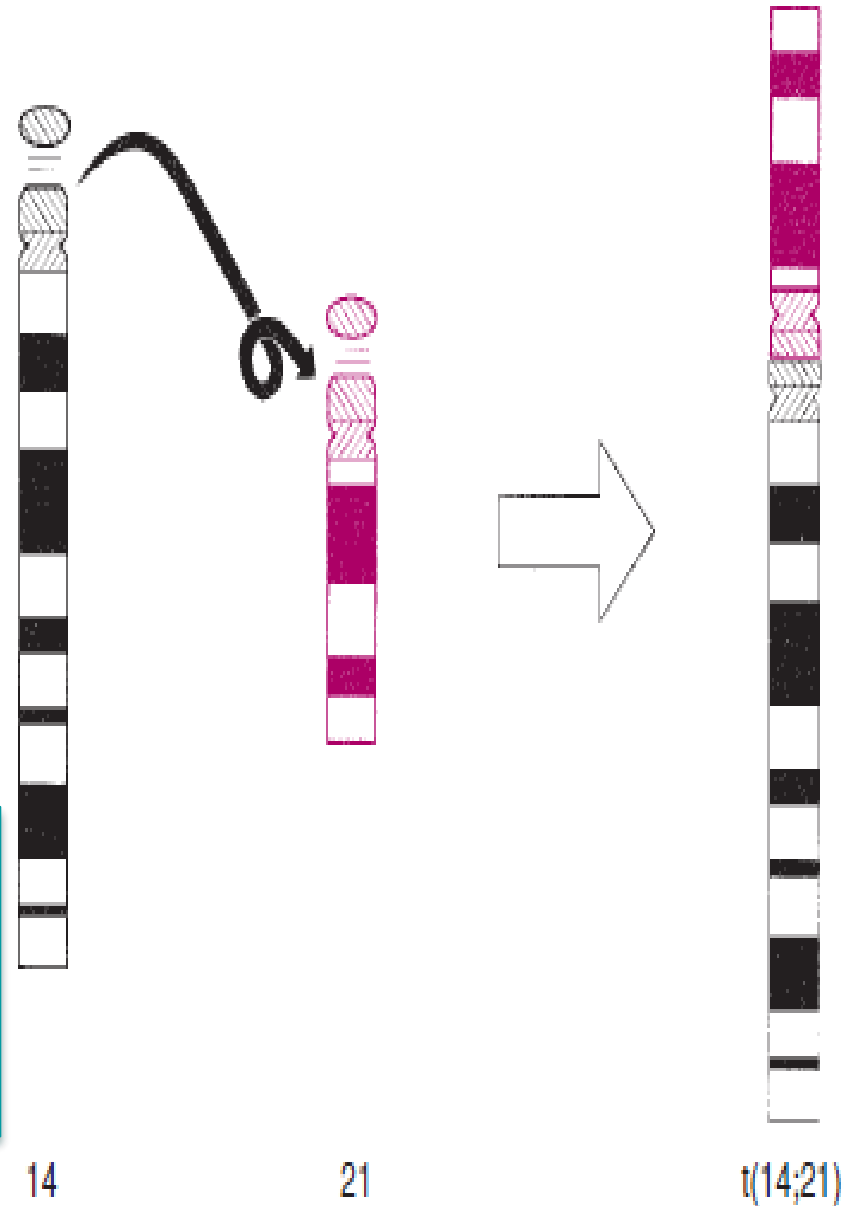
may be

1- **Balanced**, in which case breakage and reunion occur between two chromosomes but no critical genetic material is lost and individuals are normal

2- **Unbalanced**, in which case part of one chromosome is lost and an altered phenotype is produced.

An example,

unbalanced translocations between the long arms of chromosomes 14 and 21 during meiosis I or II produce gametes with an extra copy of chromosome 21, **one** of the causes of Down syndrome



at fertilization,
a gamete having 23 chromosomes fuses with
a gamete having 24 or
22 chromosomes, the result is an individual
with either 47 chromosomes

Trisomy

or 45 chromosomes

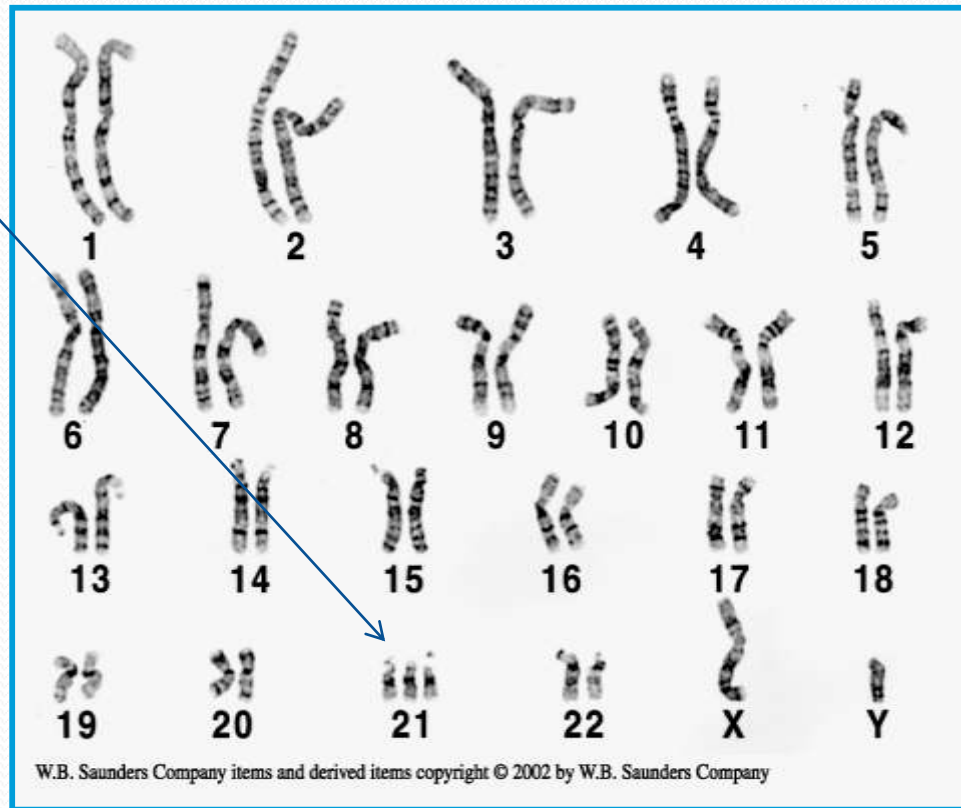
Monosomy

Down syndrome

Trisomy 21

Is usually caused by an extra copy of chromosome 21

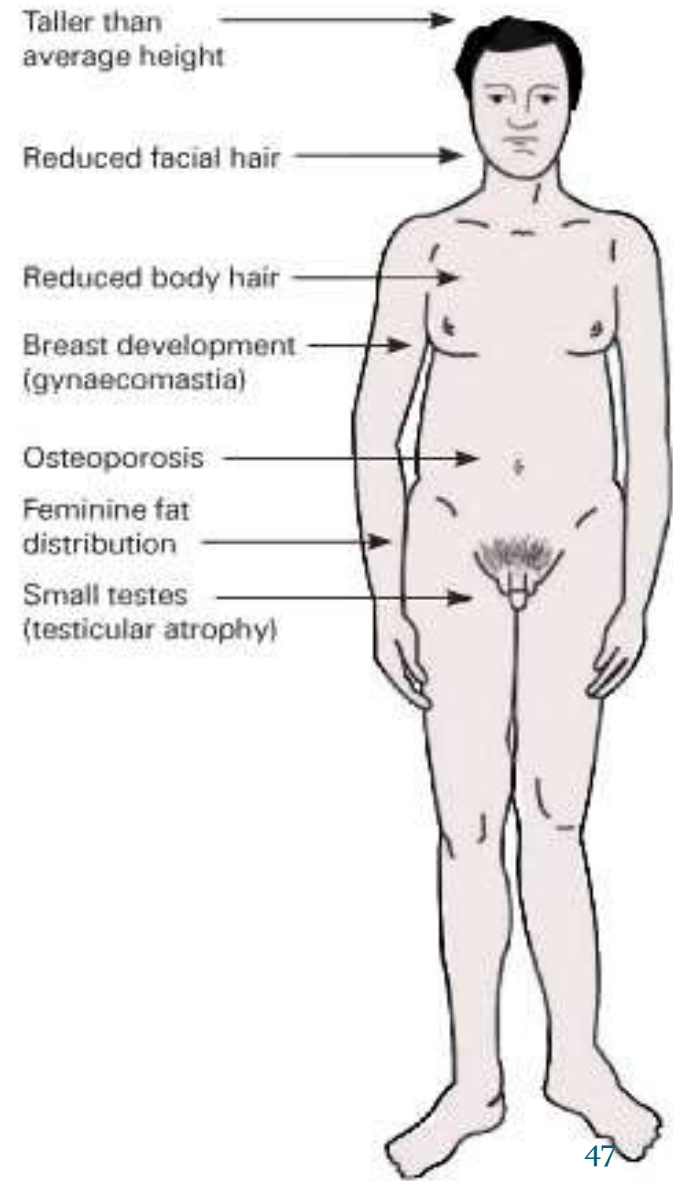
Frequency	Maternal age at birth
1/1250	15-19
1/1400	20-24
1/1100	25-29
1/900	30
1/900	31
1/750	31
1/625	32
1/500	33
1/386	34
1/300	35
1/225	36
1/175	37
1/140	38
1/100	39
1/80	40
1/65	41



90%: Meiotic nondisjunction during meiosis II of oogenesis
 10%: Meiotic nondisjunction during meiosis I of spermatogenesis

Klinefelter's Syndrome

XXY – Phenotypically male with an **extra X** chromosome



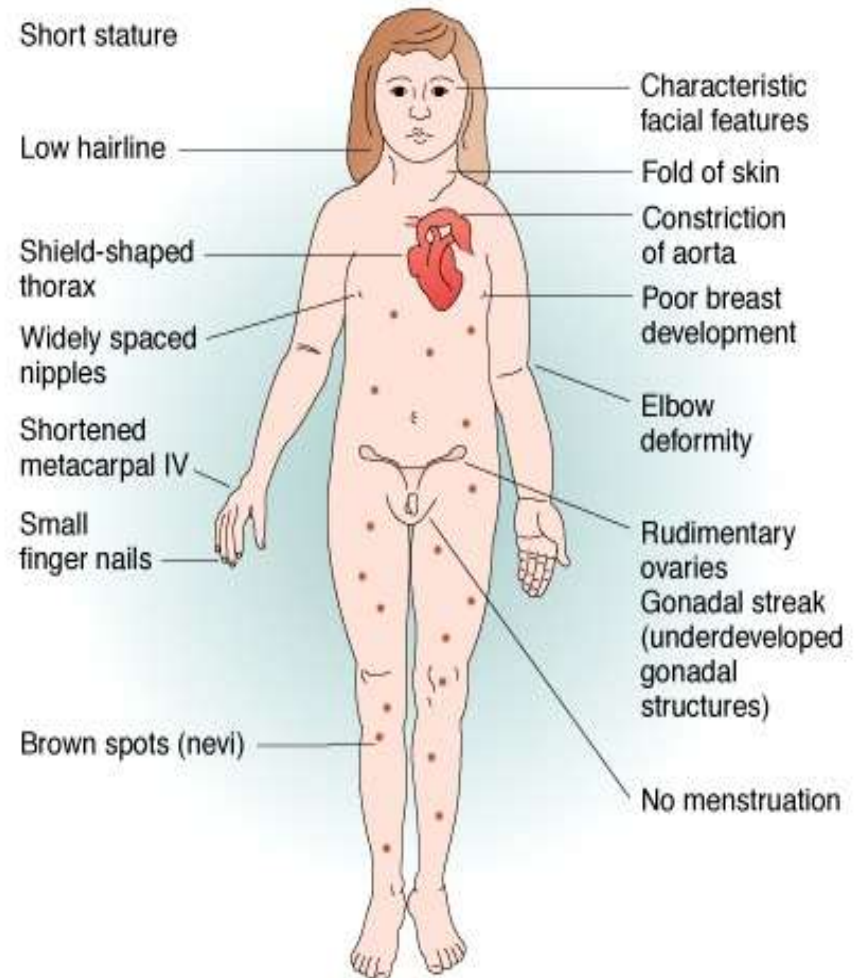
Turner's Syndrome

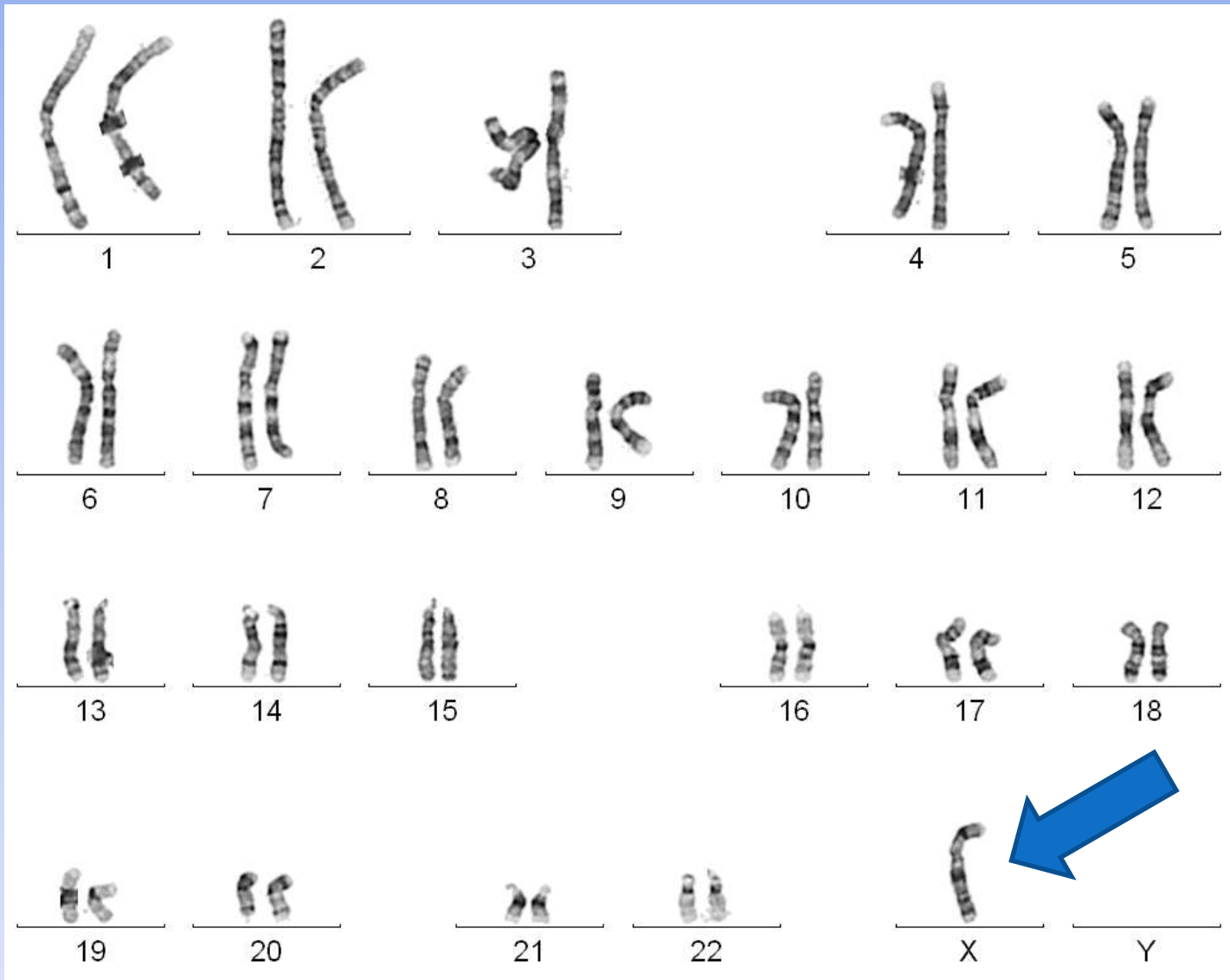
XO – Phenotypically female missing an X chromosome

is the only monosomy compatible with life.

Even then, 98% of all fetuses with the syndrome are spontaneously aborted.

The few that survive are unmistakably female in appearance and are characterized by the absence of ovaries (**gonadal dysgenesis**)





Thank you