

A. OVERVIEW OF MEIOSIS

- **Cell division** that produces **four haploid cells** that are **genetically different**.
- Produces **gametes** (sex cells)
- Has **two** divisions.
- Has **two** main **jobs**:

1. Produces haploid (sex) cells that contain half the normal number of chromosomes.

This ensures that the correct **diploid number** is **restored** at **fertilisation**.

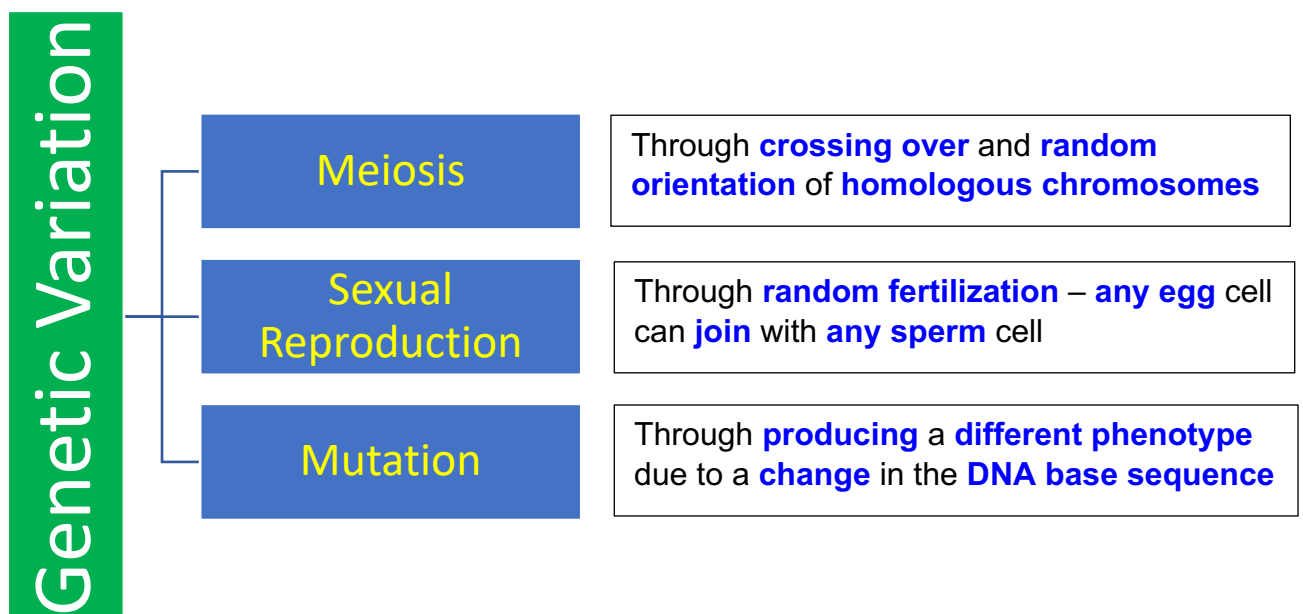
Otherwise, **each generation** would have **twice as many chromosomes** as the **previous one**.

2. Increases genetic variation

Produces **genetically different** cells.

Through **crossing over** and **random orientation** of **homologous chromosomes**

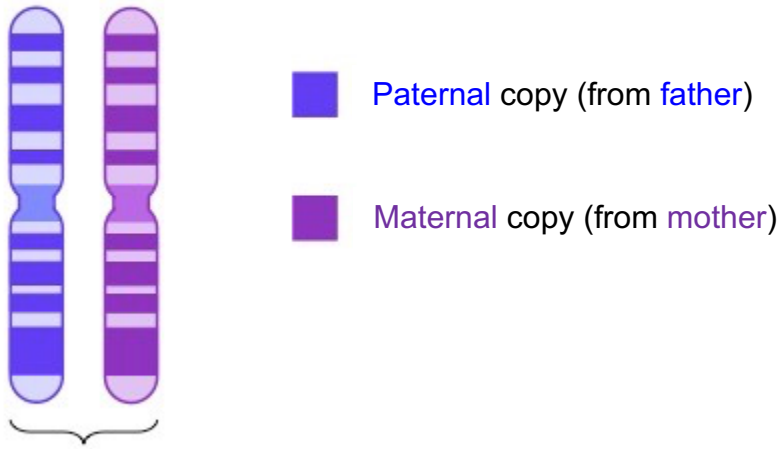
Remember though, that **meiosis** is **one** of **three** methods that **increase genetic variation**:



B. A RECAP ON HOMOLOGOUS CHROMOSOMES

What they are

Here is one pair of homologous chromosomes:



Homologous pair

- They are the **same shape** and **size**
- They carry the **same genes** in the **same sequence**
- They carry **different alleles**
- In each pair, **one** chromosome is from **mum** and the **other** is from **dad**

However, in text books, chromosomes look like the letter '**X**', with a **circle** at the centre.

Why is this?

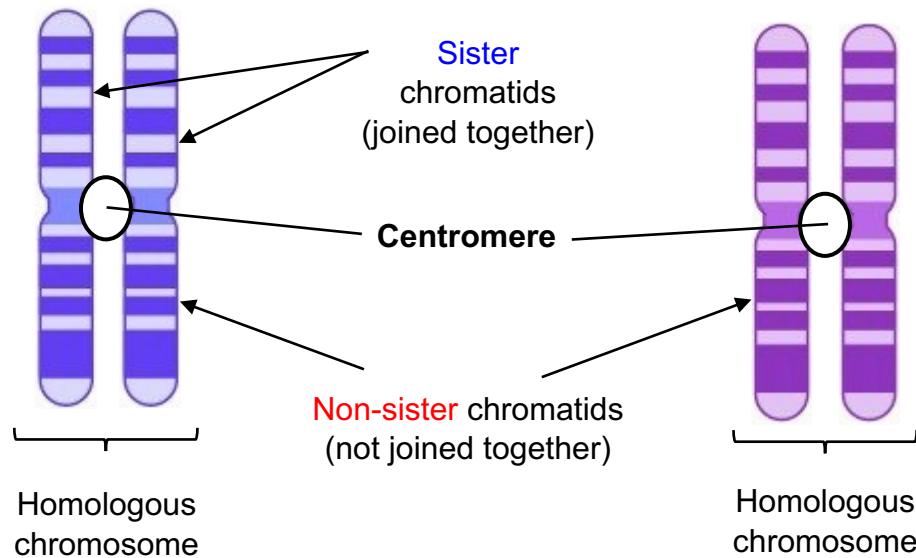
Why chromosomes look like an 'X'

Look at the **pair of homologous chromosomes** again:



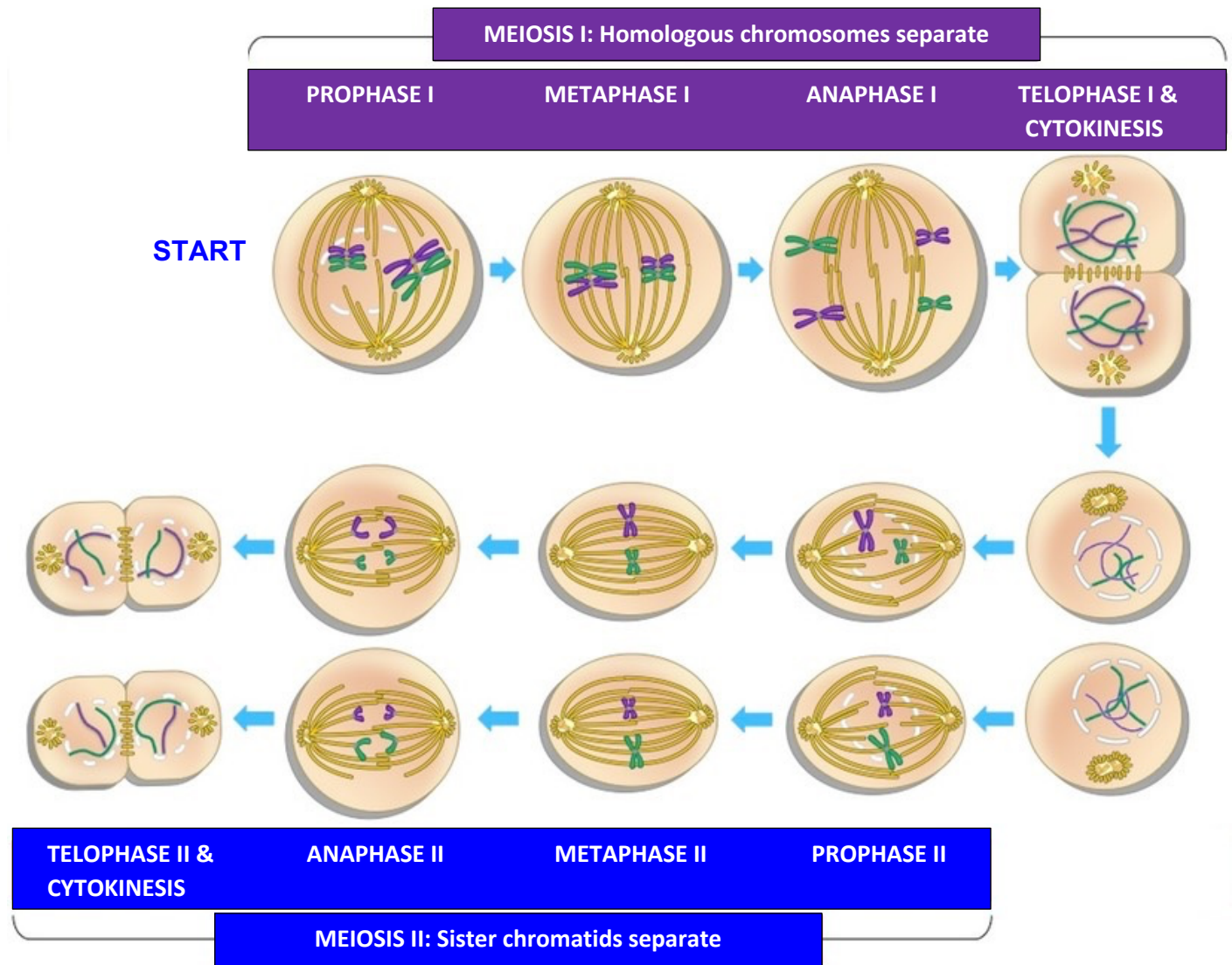
Homologous pair

- They look like this as they have **not been replicated** yet during **interphase**.
- **Interphase** happens **before cell division** (meiosis or mitosis)
- During **interphase**, a **copy** of each chromosome is **made** and it is **connected** to the **original chromosome** by a **centromere**, as shown below.



- Each **homologous chromosome** is made up of two **sister chromatids** (joined together).
- One **sister chromatid** is a **copy** of the **original** chromosome
- The **sister chromatids** are **held together** by a **centromere**
- It is these **X-shaped** chromosomes that undergo cell division (meiosis or mitosis).

C. THE PROCESS OF MEIOSIS



FIRST DIVISION

- Halves the number of chromosomes
- Increases genetic variation
- Separates homologous chromosomes

Prophase I	Homologous chromosomes pair up Crossing over happens where chromatids break and recombine
Metaphase I	Homologous chromosomes move to the equator of the cell
Anaphase I	Homologous chromosomes separate and move to opposite poles
Telophase I	Homologous chromosomes reach the poles and unwind

SECOND DIVISION

- Produces more cells
- Similar to mitosis
- Separates sister chromatids

Prophase II	Chromosomes condense and become visible
Metaphase II	Chromosomes move to the equator of the cell
Anaphase II	Sister chromatids separate and move to opposite poles
Telophase II	Chromosomes reach the poles and unwind

Contrast meiosis I and meiosis II

MEIOSIS I	MEIOSIS II
Homologous chromosomes pair up in prophase	Homologous chromosomes do not pair up in prophase
Crossing over occurs in prophase	Crossing over does not occur in prophase
Pairs of homologous chromosomes move/line up on equator in metaphase	Single chromosomes move/line up on equator in metaphase
Homologous chromosomes are separated in anaphase	Sister chromatids are separated in anaphase
Halves the chromosome number	Does not half the chromosome number
Produces two daughter cells	Produces four daughter cells
Increases genetic variation by crossing over and random orientation of homologous chromosomes	Increases genetic variation by random assortment of chromatids (into gametes)

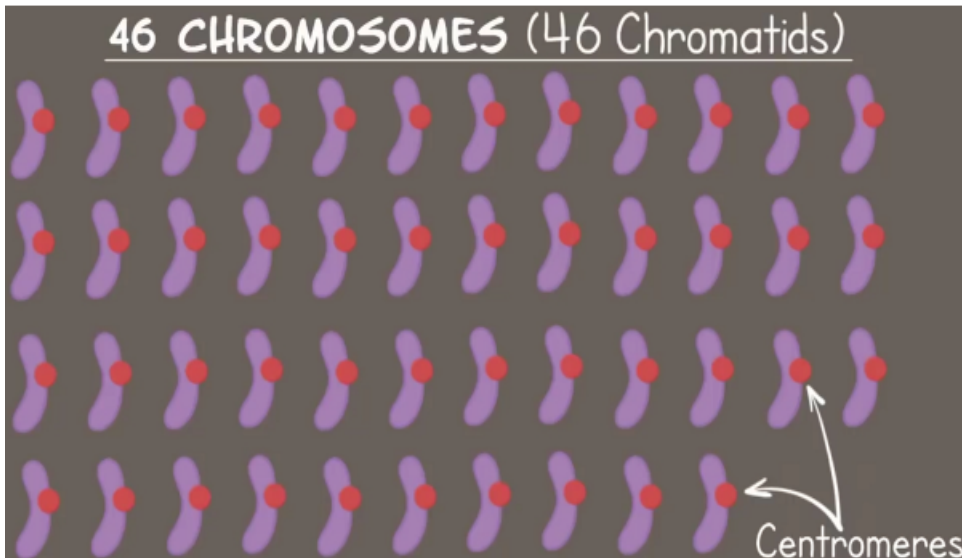
Comparing the number of cells, chromosomes and chromatids

	Number of cells	Number of chromosomes	Number of chromatids per chromosome
Before meiosis starts	1	2n	2
At the end of the first division	2	n	2
At the end of the second division	4	n	1

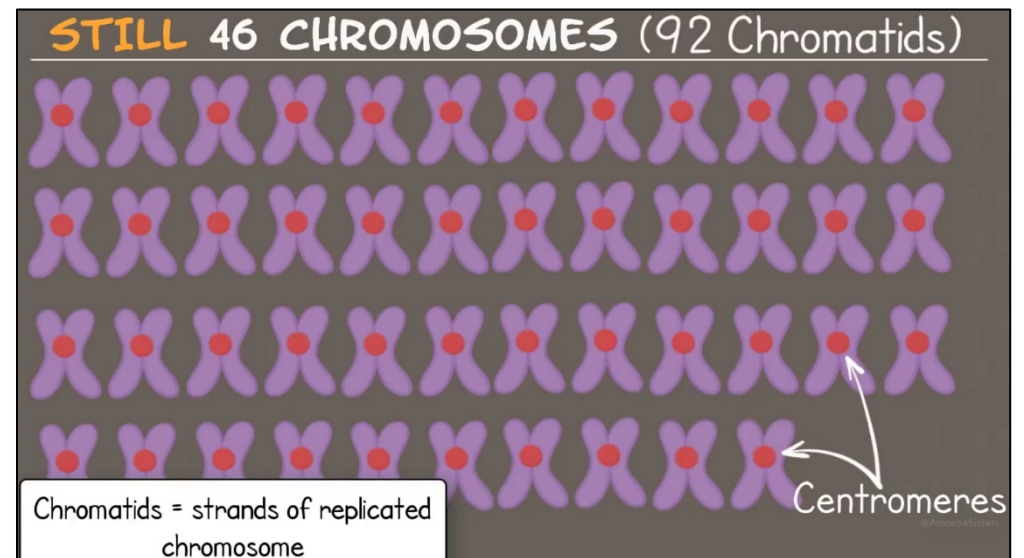
C. NUMBER OF CHROMOSOMES AND CHROMATIDS IN DIFFERENT STAGES OF MEIOSIS

- By convention, we **count the number of centromeres** to give us the **number of chromosomes**.
- This confuses many students!

BEFORE DNA REPLICATION AND MEIOSIS

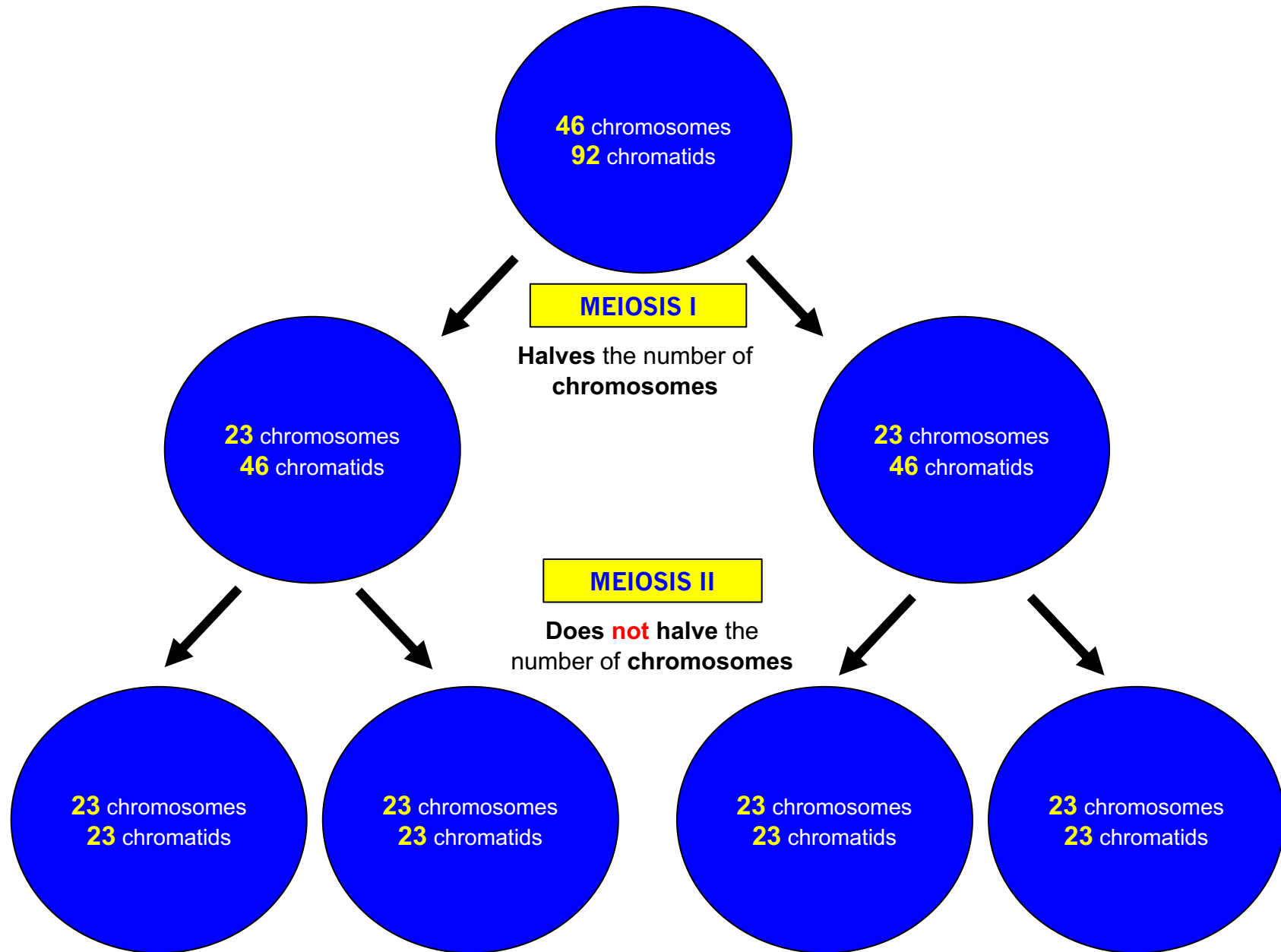


AFTER DNA REPLICATION AND AT THE START OF MEIOSIS



- Both diagrams show the **same number of chromosomes** but the one on the right has **double the number of chromatids**.
- We measure the **amount of DNA** in a cell by the **number of chromatids**.

IN HUMANS:



Amended Exam Question

A **sex cell** from an organism contains **14 chromosomes**.

- (a) Complete the table to show how many chromosomes or chromatids would have been present in **this cell** at different stages of meiosis.

Description	Number
Number of chromosomes at the start of meiosis	28
Number of chromatids at the start of meiosis	56
Number of chromatids in metaphase I	56 (cell has not yet split)
Number of chromatids in metaphase II	28 (not 14 until the cell splits)

- (b) The amount of DNA in this **sex cell** is **x**.

What would be the amount of DNA present in this cell at the start of meiosis?

Explain your answer.

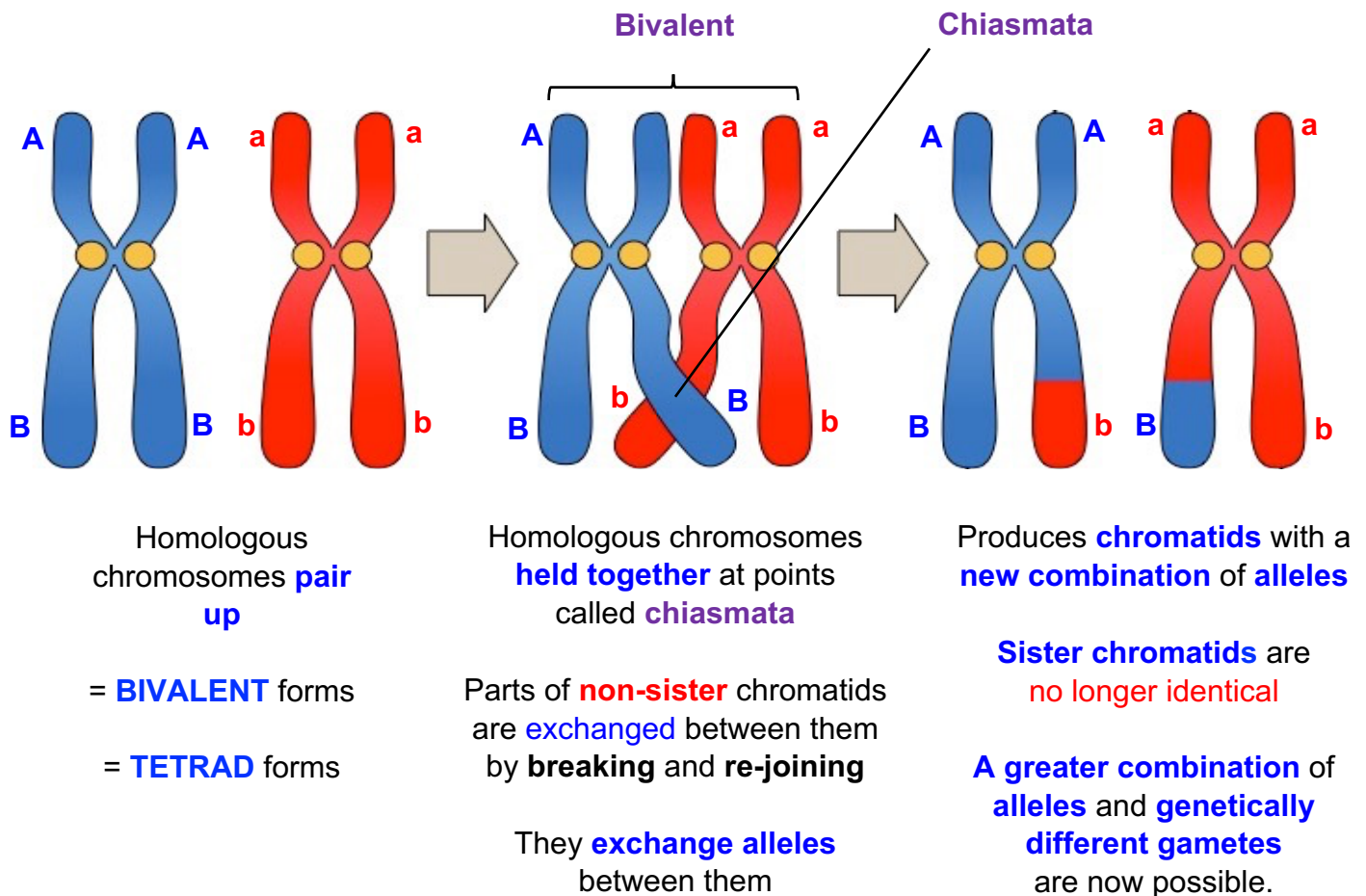
Amount of DNA (at start of meiosis) = **4x (1)**;

(As) there are 4x more chromatids (at the start) **(1)**;

E. HOW MEIOSIS INCREASES GENETIC VARIATION

1. Crossing over of alleles in Prophase I

- Consider two genes, **A** and **B**, on a homologous chromosome.
- In **prophase I**, homologous chromosomes undergo a process called **synapsis**, whereby they **pair up** to form a **bivalent** (or **tetrad** = **four sister chromatids**).



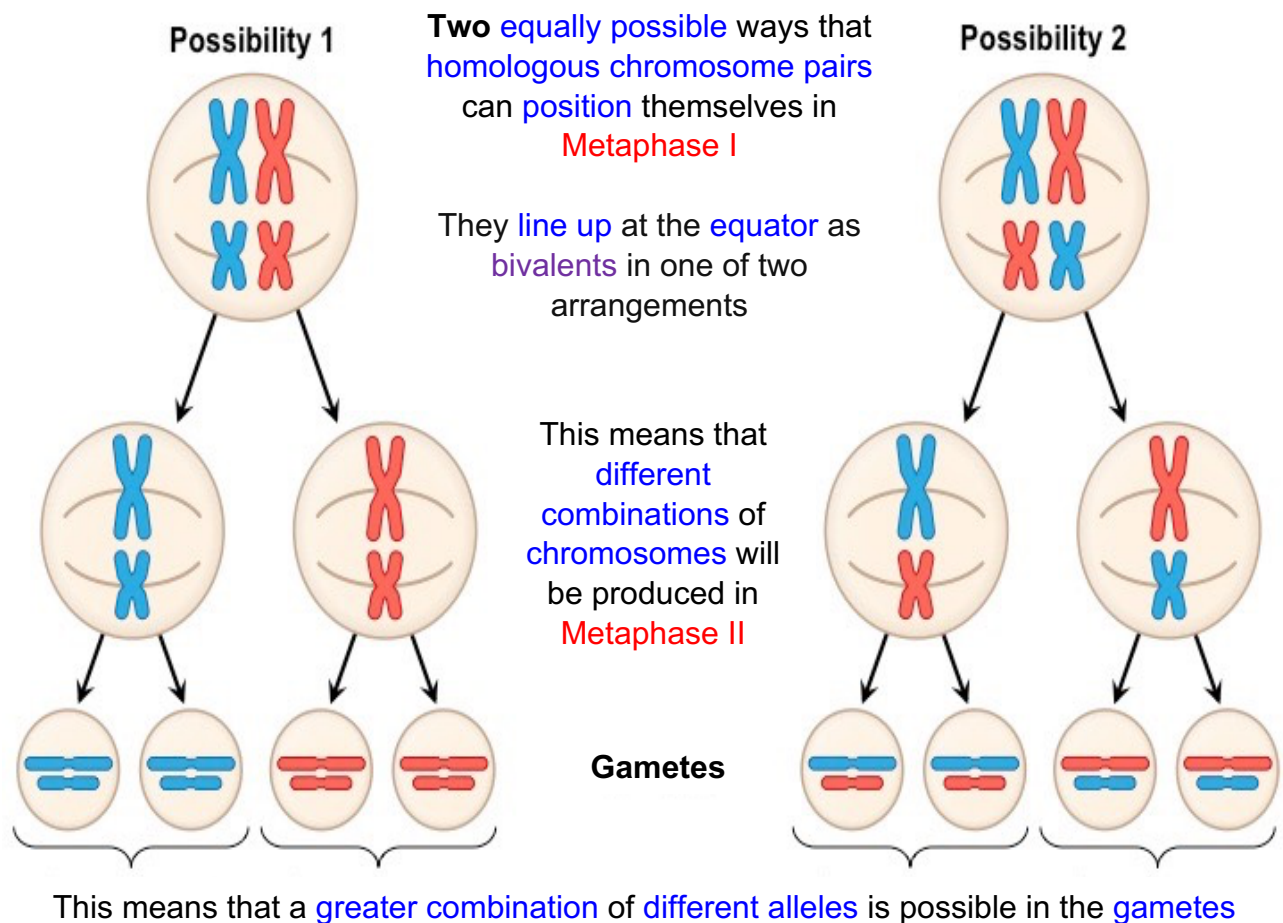
It is **random** where along the length of the chromosome **crossing over** occurs

Crossing over is also a **rare event** – it occurs at a **low frequency**

This further **increases genetic variation**

2. Random orientation of homologous chromosomes in Metaphase I

- This is also known as **independent assortment/segregation**.
- The **orientation** of **pairs of homologous chromosomes** in **metaphase I** is **random**.



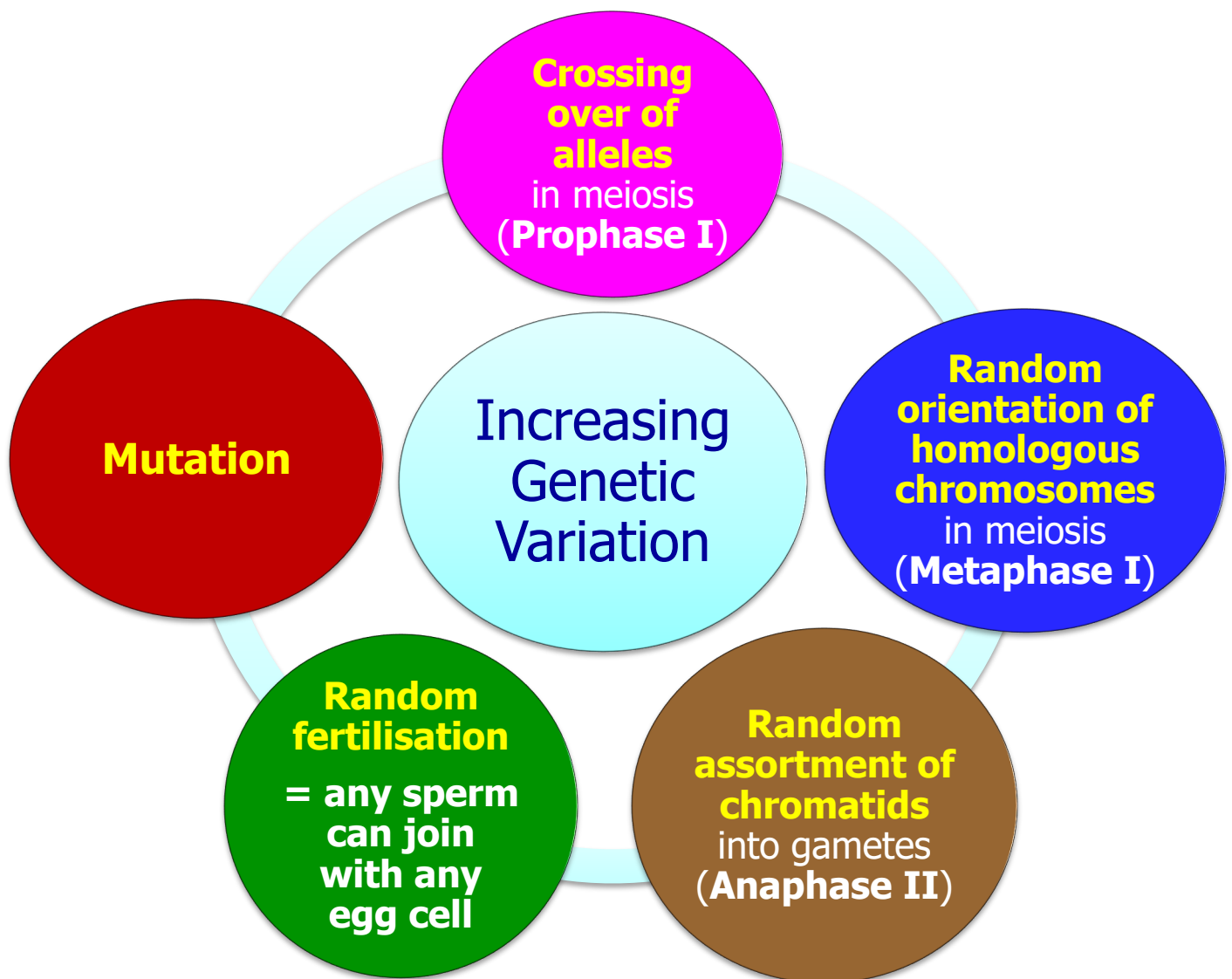
- Different gametes have different combinations of **maternal** and **paternal** chromosomes.

In humans, the number of different combinations of chromosomes possible in gametes is:

$$2^{23} = \text{over 8 million}$$

F. INCREASING GENETIC VARIATION

- **Three** of these ways involve **meiosis**

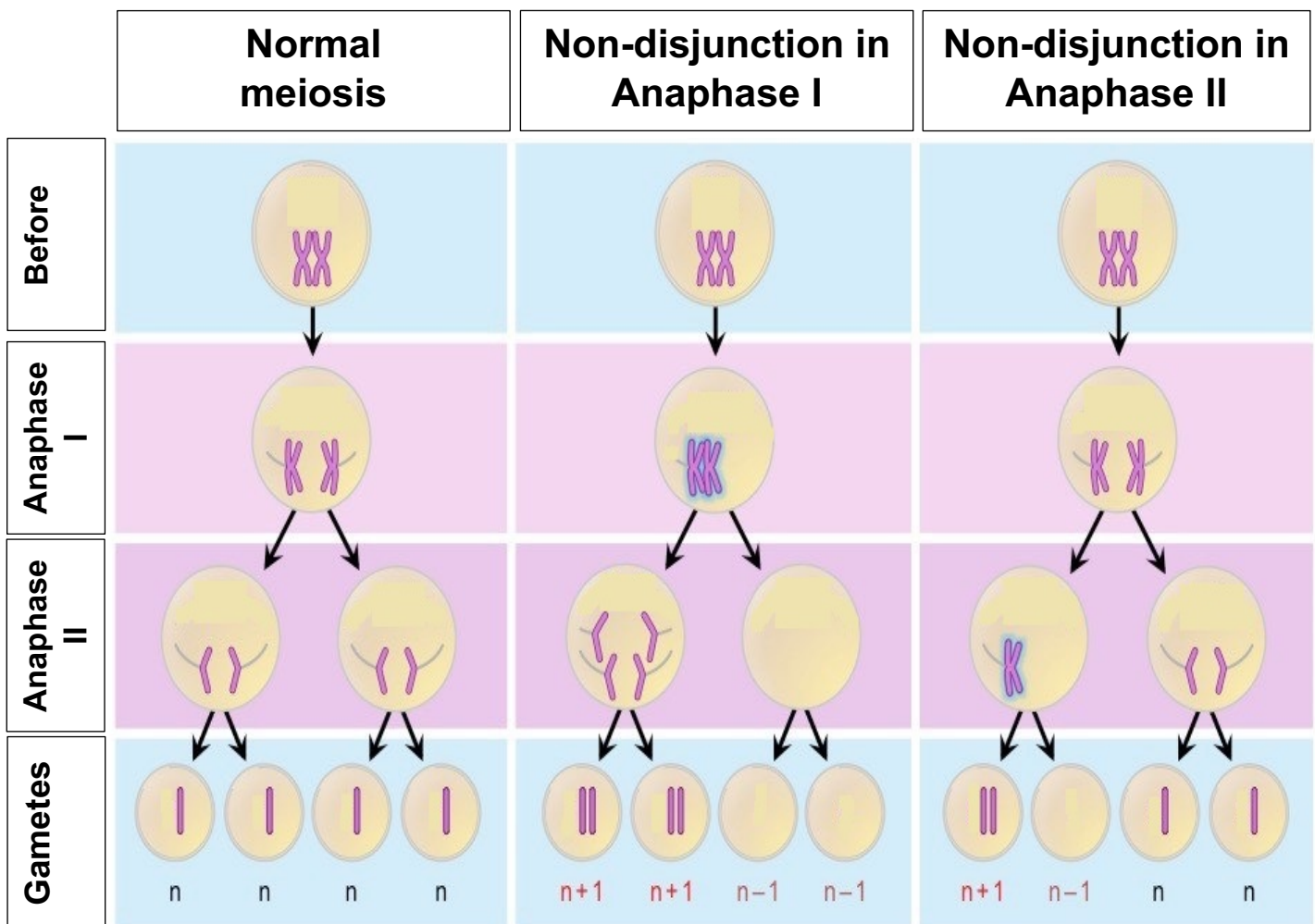


Natural selection **DECREASES** genetic variation

G. DOWN SYNDROME

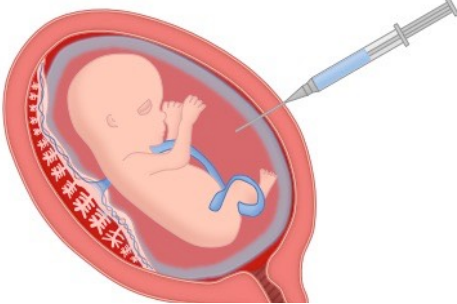
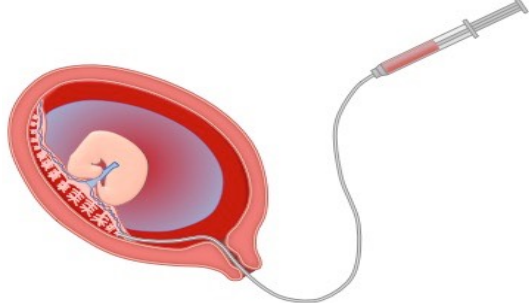
How it occurs

- Individuals with Down syndrome have **three copies** of **chromosome 21** (= 'trisomy 21').
- It is caused by **non-disjunction**.
- This is where chromosomes **do not separate** in **Anaphase I** or **II** of **meiosis**, so they **move to the same pole** of the cell.
- This causes the **gametes** to have either **one extra chromosome** or **one chromosome less**.

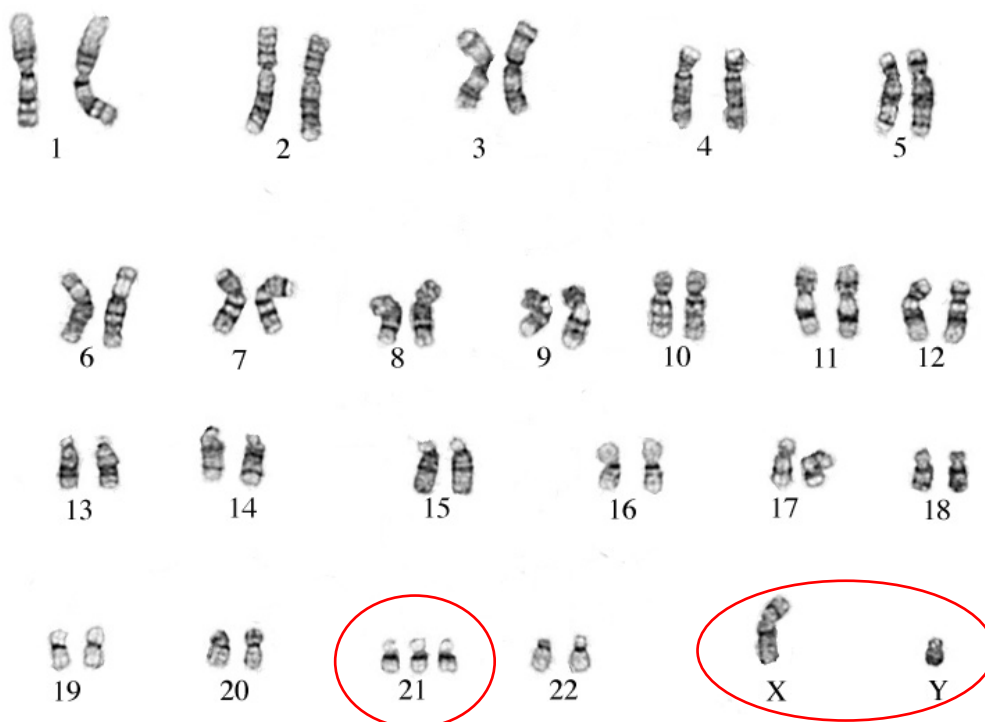


- Non-disjunction** of **chromosome 21**
- Homologous chromosomes** **do not separate** in **Anaphase I**
- Due to **incorrect attachment** to the **spindle**
- OR
- Sister chromatids** **do not separate** in **Anaphase II**
- Due to **centromeres not dividing/splitting**
- (So) they **move** to the **same pole**
- Gametes can have **one extra** chromosome or **one less** chromosome
- Gamete** with **2 copies** of chromosome 21 **fertilises** a **normal gamete**
- Zygote** produced has **3 copies** of chromosome 21
- Risk** of Down syndrome **increases** as the **mother's age increases**

Two methods used to obtain cells from an unborn child for chromosome testing

Amniocentesis	Chorionic villus sampling (CVS)
	
Needle used to withdraw foetal cells from the amniotic fluid	Suction tube used to withdraw foetal cells from the chorion (where the placenta will develop from)
Inserted through the abdominal wall and uterus wall	Inserted through the cervix (neck of the uterus)
Usually done later in pregnancy (16 weeks)	Can be done earlier in pregnancy (11 weeks)
Slight risk of inducing miscarriage (~1%)	Slightly lower risk of miscarriage (~0.5%)

- The chromosomes can then be arranged in **homologous pairs** to produce a **karyogram**.
- Doctors can then check the baby's **sex** and look for any **chromosome abnormalities**.



This is a **male** with **Down syndrome**