

Science of Living System (BS20001)

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Genetic Engineering

- ❑ Manipulation of an organism's genome
- ❑ Can range from changing one base pair (A-T or C-G), deleting a whole region of DNA, or introducing an additional copy of a gene
- ❑ Extracting DNA from an organism's genome and combining it with the DNA of another individual
- ❑ Used to enhance or modify the characteristics of an individual organism

An Overview of Genetic Engineering

**Agricultural application (Bt Cotton,
Genetically
modified sweet corn)**

Bacillus thuringiensis

**Application in Dairy, Poultry Farms etc
(Growth hormones to increase production of
cow milk)**

**Environmental application (*Pseudomonas
putida*, an oil eating bacteria)**

**Therapeutic application (Human insulin,
many growth hormones, etc)**



Genetically modified Product

Flavr Savr (also known as CGN-89564-2; pronounced "flavor saver"), a genetically modified tomato



Glow Fish: Zebra fish with green fluorescent protein (GFP) and others

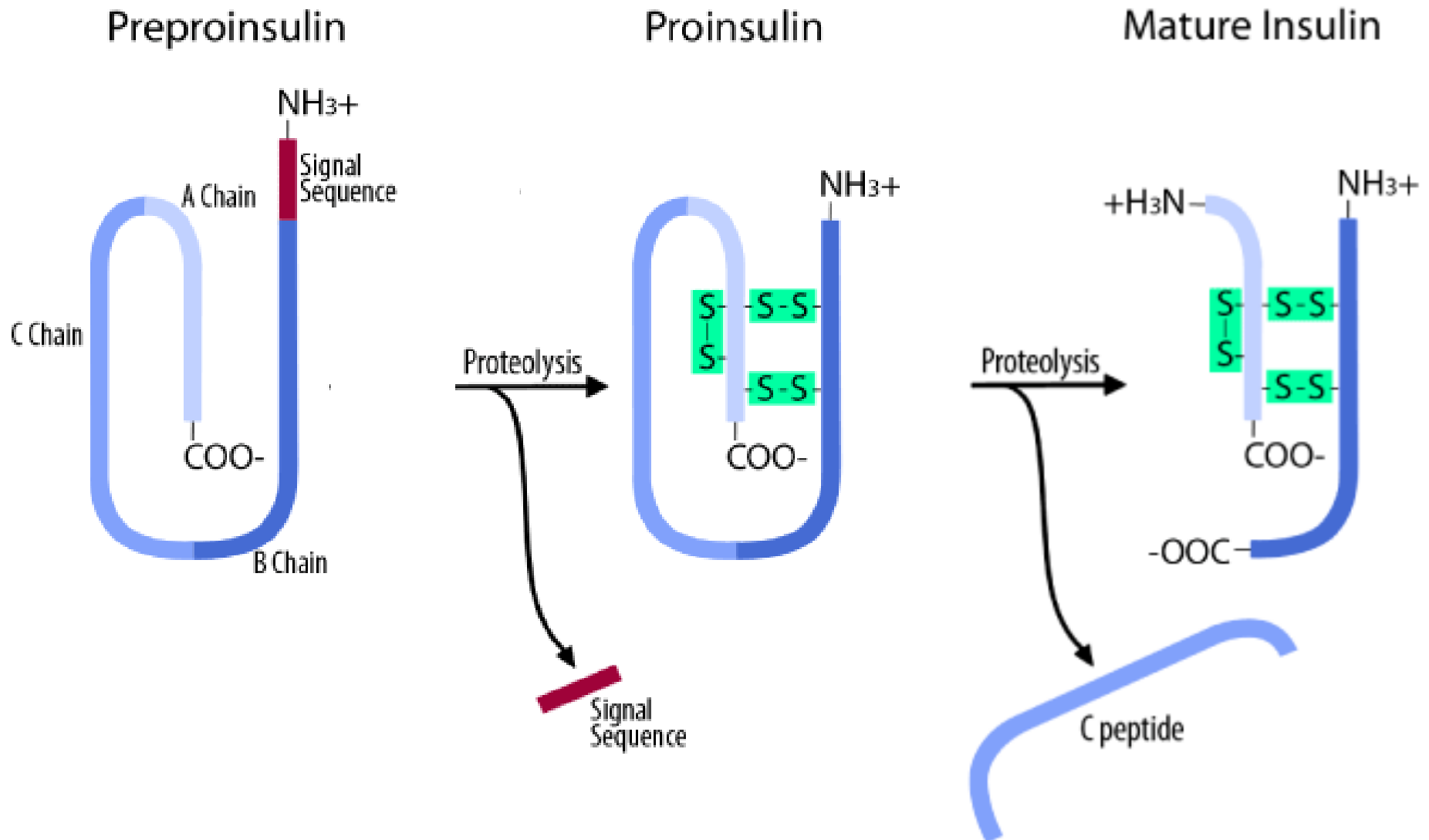


Wild type zebra fish



Glow fish (genetically modified fish)

Production of Human Insulin: The best example of Genetic Engineering/Recombinant DNA Technology



Insulin: an indispensable hormone in our body

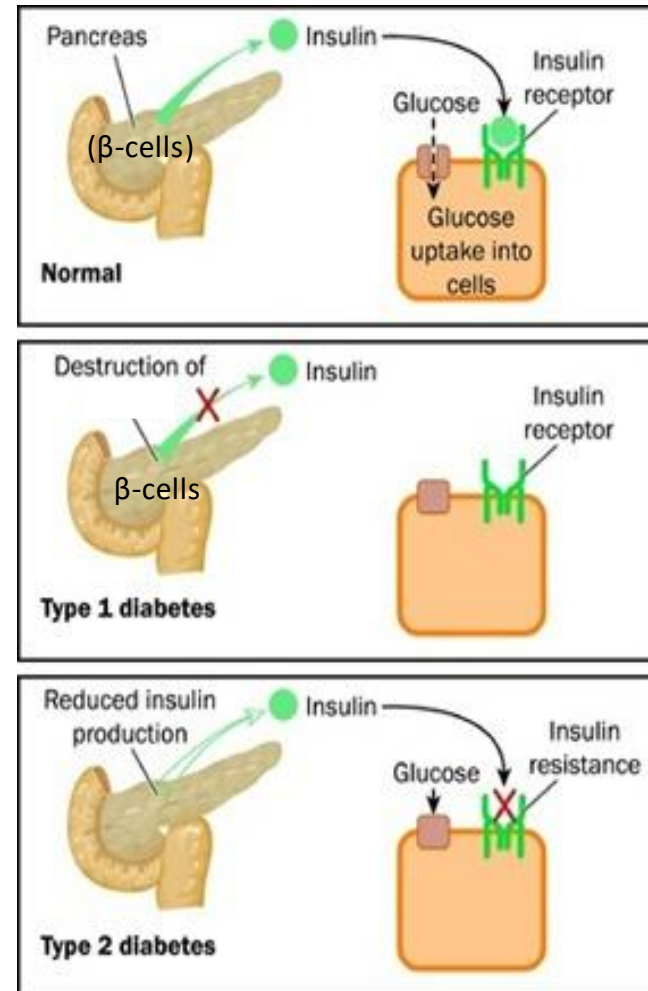
Insulin is a hormone that regulates the amount of glucose (sugar) in the blood and is required for the body to function normally.

Insulin is produced by cells in the pancreas, called the islets of Langerhans.

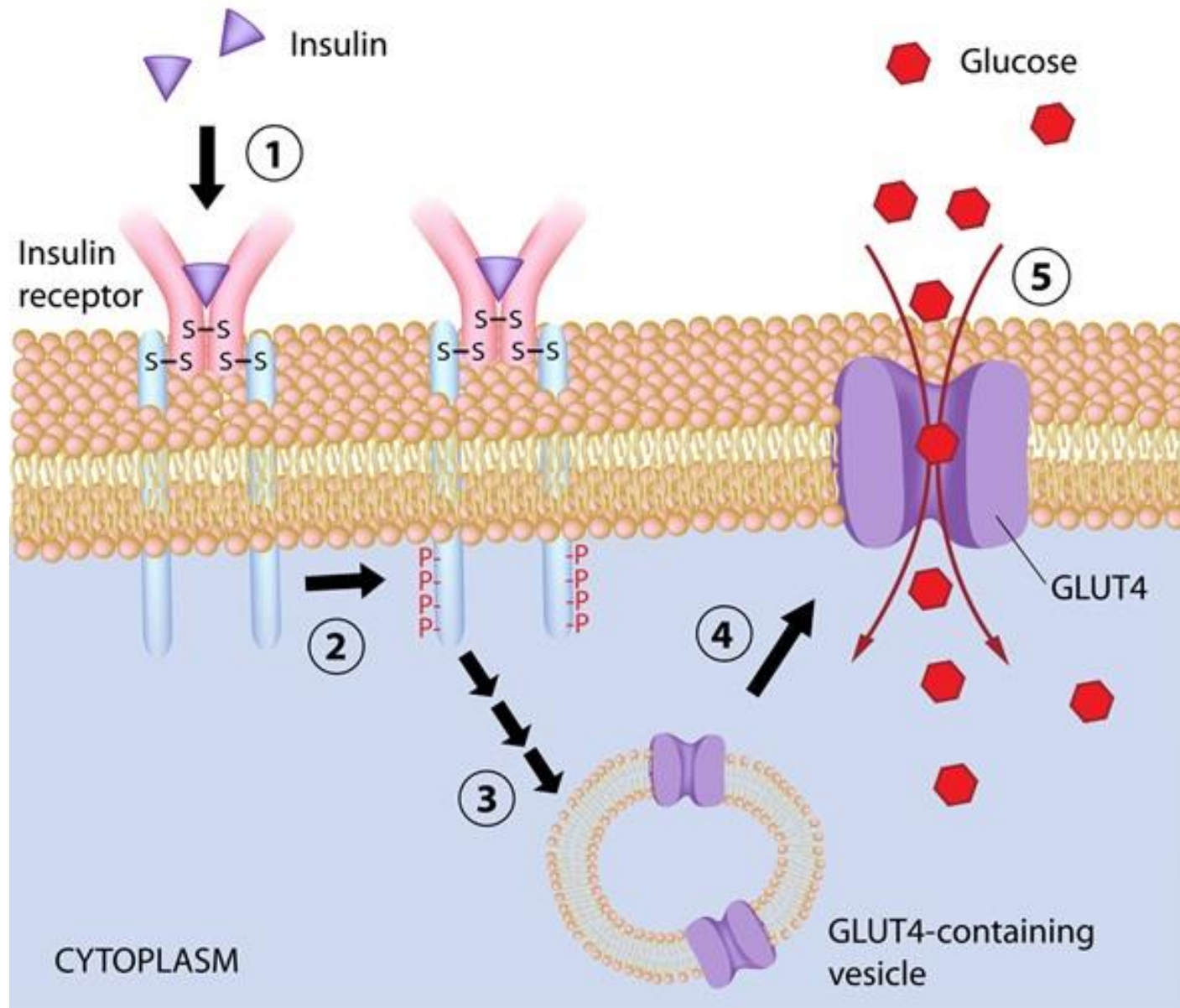
These cells continuously release a small amount of insulin into the body, but they release surges of the hormone in response to a rise in the blood glucose level.

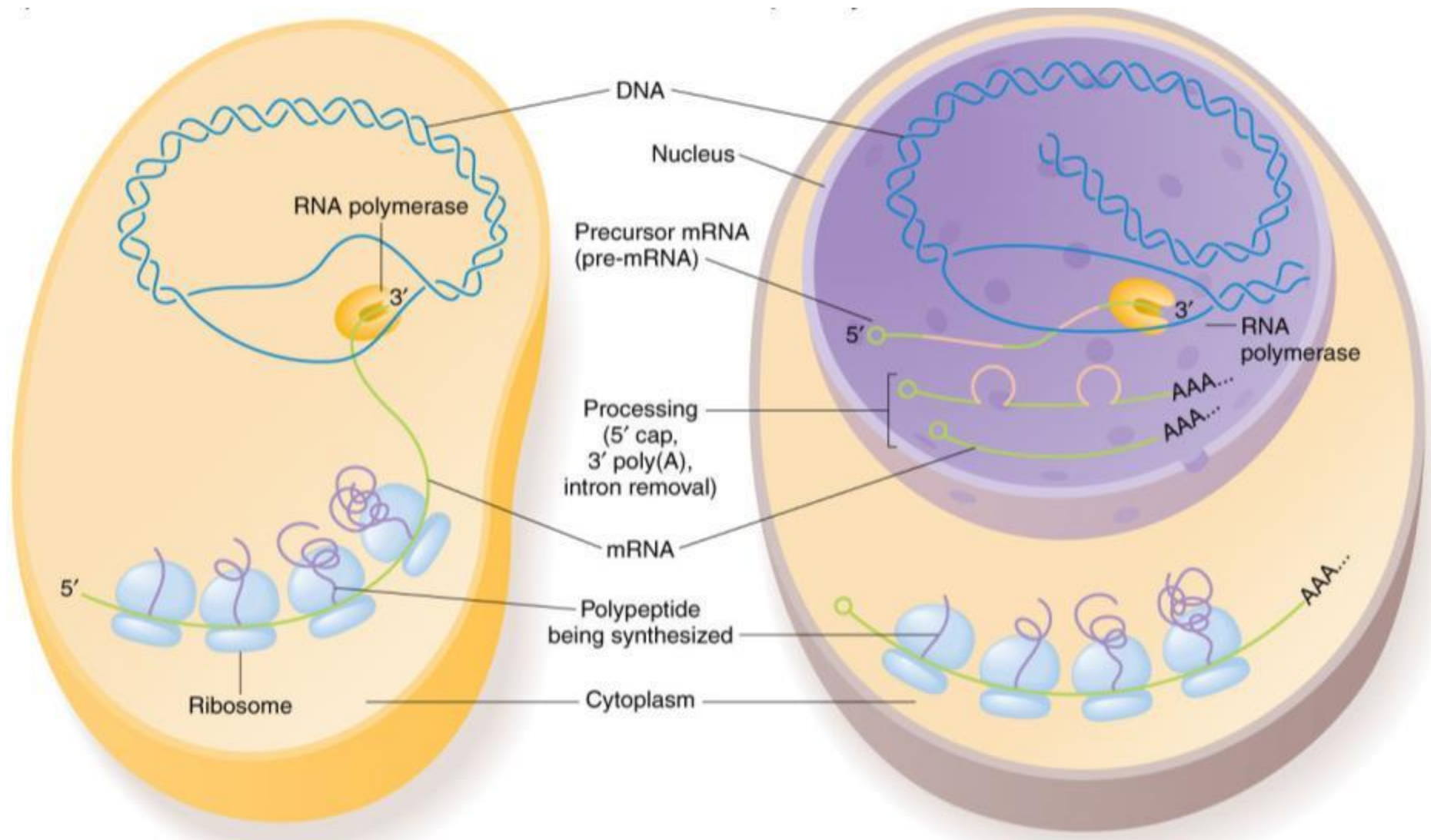
Every time a person eats, the blood glucose rises. Raised blood glucose triggers the cells in the islets of Langerhans to release the necessary amount of insulin. Insulin allows the blood glucose to be transported from the blood into the cells.

Once transported into the cell, the blood glucose level is returned to normal within hours.



Effect of Insulin on Glucose Uptake





Bacteria

Eukaryote

Production of Insulin

1921: Frederick Banting and Charles Best successfully purified insulin from a dog's pancreas. Over the years scientists made continual improvements in producing insulin.

In the 1980s, researchers used genetic engineering to manufacture a human insulin.

In 1982, the Eli Lilly Corporation produced a human insulin that became the first approved genetically engineered pharmaceutical product.

At present, researchers could produce genetically engineered insulin in unlimited supplies without depending on animals.

HOW??

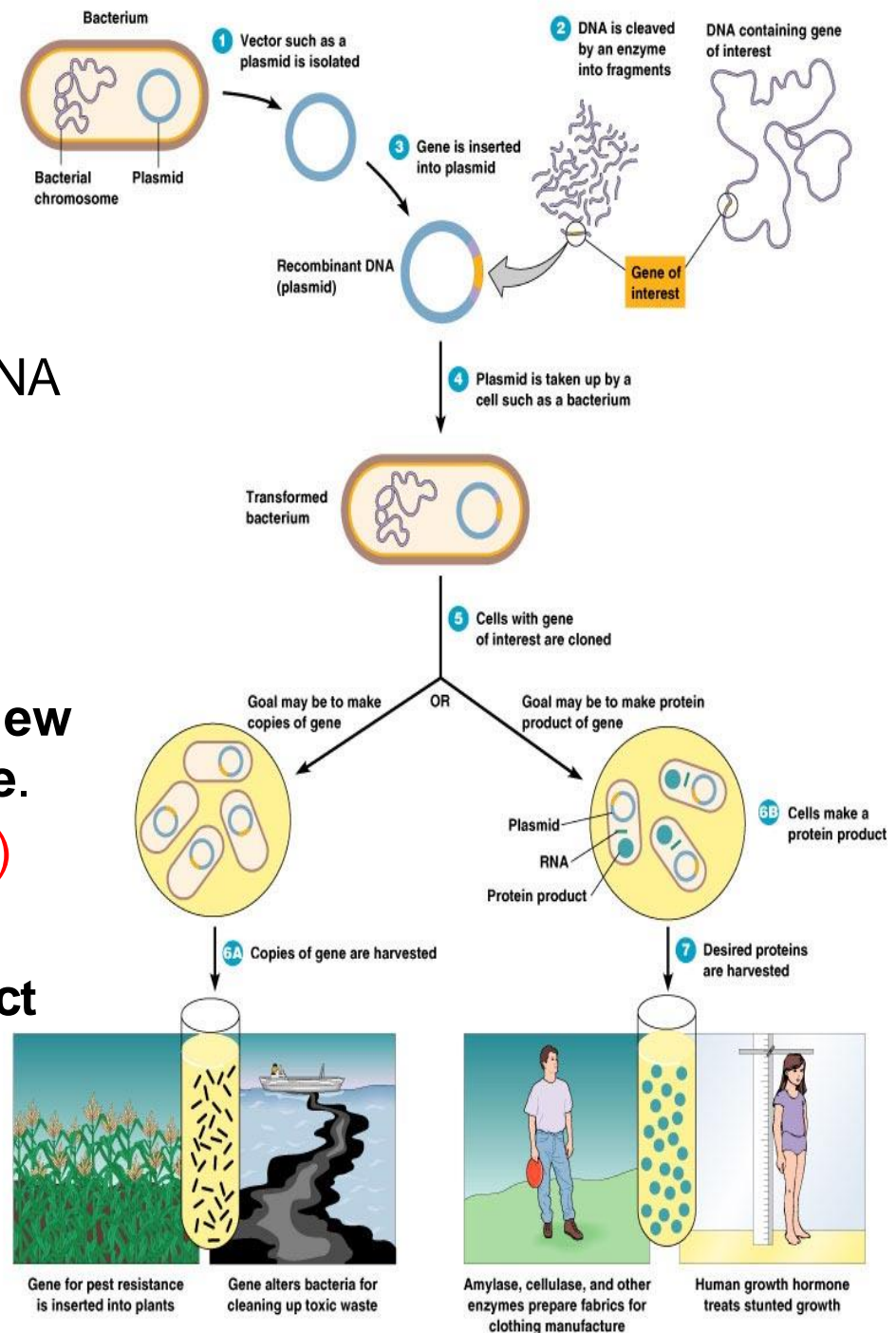
Genetic Engineering

Key Words in Genetic Engineering

- ❑ Gene of interest: DNA segment that is to be inserted or deleted
- ❑ Plasmid: a small, circular, double-stranded DNA molecule that is distinct from a cell's chromosomal DNA (commonly found in bacteria and may provide antibiotic resistance)
- ❑ Vector: DNA molecule (such as plasmid) used as a vehicle to carry gene of interest (or foreign genetic material) into another cell where it can be expressed
- ❑ Transformation: Transfer of gene of interest in to a host cell (may be bacteria) where it can be maintained as well as expressed.
- ❑ Clone: Organisms carrying identical genes

An Overview of Genetic Engineering

1. **Gene of interest (DNA)** is isolated
(DNA fragment)
2. **A desired gene** is inserted into a DNA molecule - **vector**
(plasmid, bacteriophage or a viral genome)
3. **The vector** inserts the DNA into a **new cell**, which is grown to form a **clone**.
(bacteria, yeast, plant or animal cell)
4. Large quantities of the **gene product** can be harvested from the clone.



Tools for making a collage



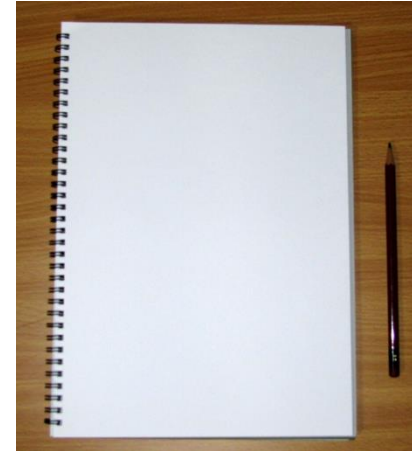
Colourful papers



Scissors

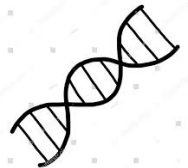


Glue

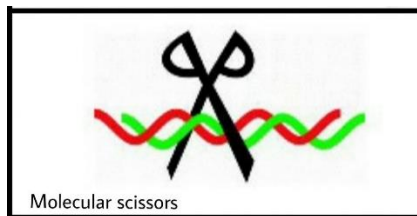


Drawing paper

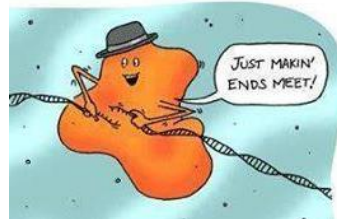
DNA of interest



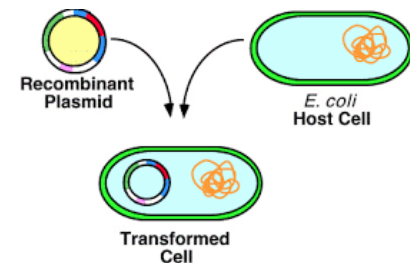
Restriction enzymes



Ligase enzyme



Host organism



Tools for making a Recombinant DNA

Tools for Genetic engineering

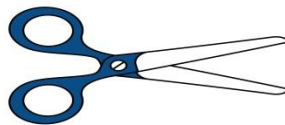
1. Restriction Enzymes

- **Naturally produced by bacteria – restriction endonucleases**
 - **Natural function** - destroy bacteriophage DNA in bacterial cells
 - Cannot digest host DNA with methylated C (cytosine)
- **A restriction enzyme**
 - **Substrate –DNA** -recognizes one particular nucleotide sequence in DNA and **cuts** the DNA molecule (breaks down the bond between two nucleotides)

sticky ends

G|AATTC
CTTAAG|

Palindrome



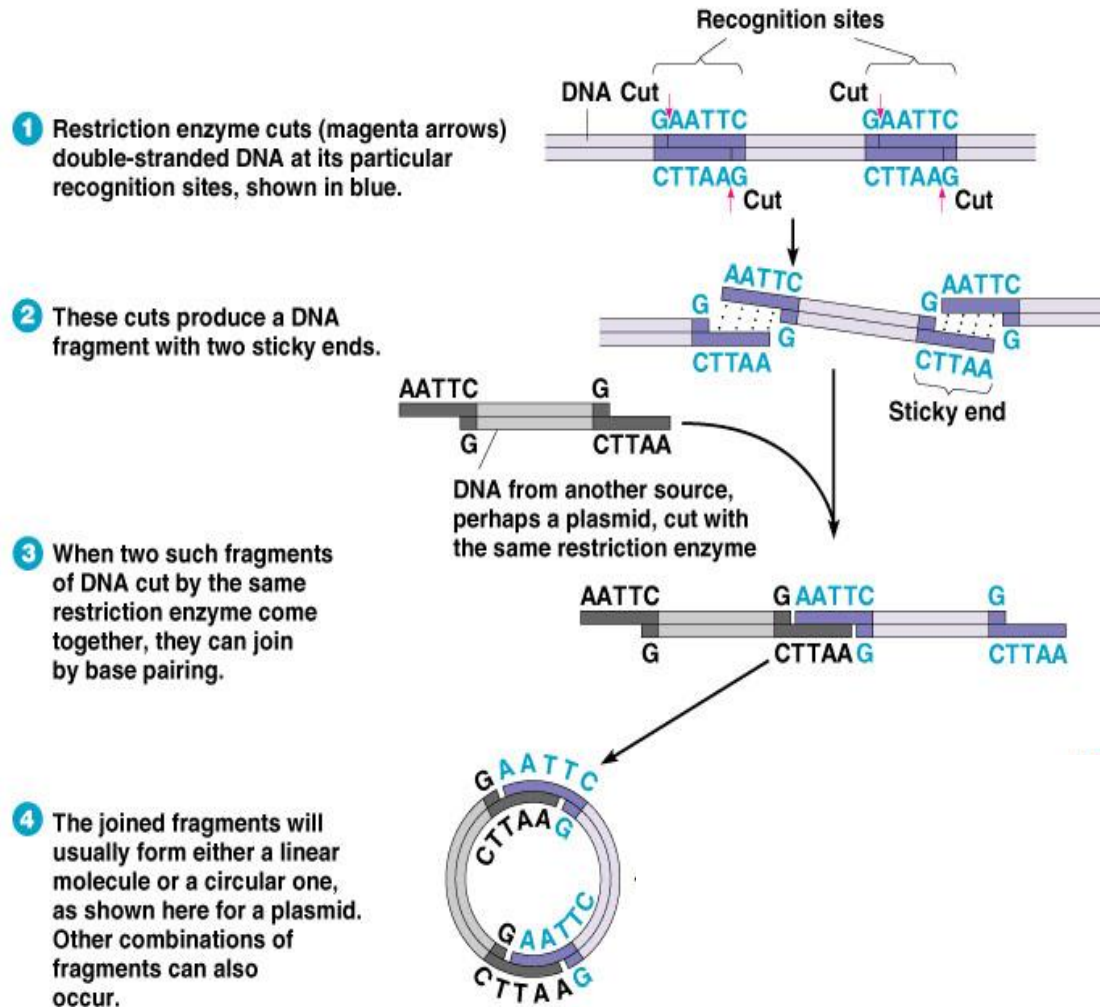
blunt ends

CCCGGG
GGGCCC

- Prepackaged kits are available for rDNA techniques

Restriction Enzymes: How it Works?

- Fragments of DNA produced by the same restriction enzyme will spontaneously join by **base pairing**.
- Each of the DNA strands will have a break



Frequency of occurrence of restriction sites

If DNA sequence has equal amounts of each base

If bases are distributed randomly

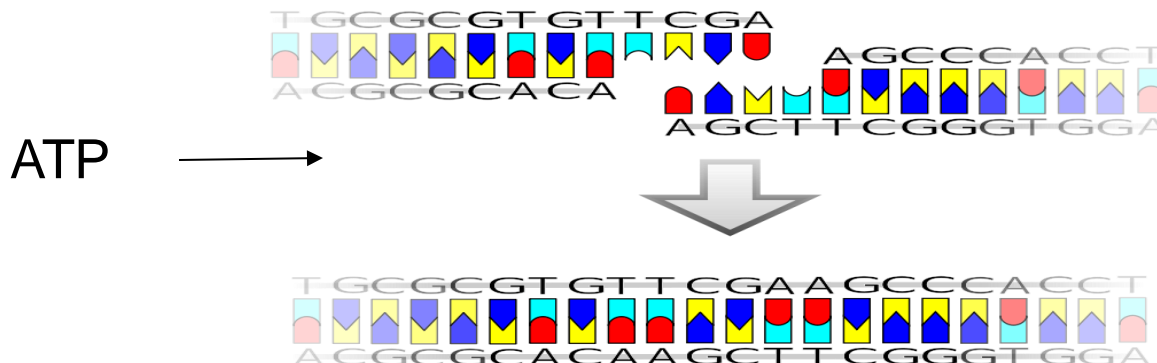
6 base cutter $(1/4)^6 = 1$ site in ~4000 bp

4 base cutter $(1/4)^4 = 1$ site in 256 bp

Tools for Genetic engineering

2. Ligase

- **DNA ligase** is an enzyme that can link together DNA strands that have double-strand breaks (a break in both complementary strands of DNA).
 - Naturally DNA ligase has applications in both **DNA replication** and **DNA repair**.
 - Needs ATP
- DNA ligase has extensive use in molecular biology laboratories for **genetic recombination experiments**



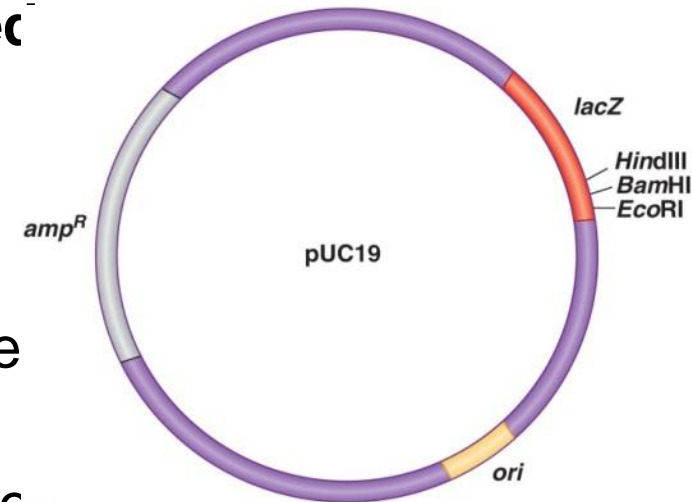
Tools for Genetic engineering

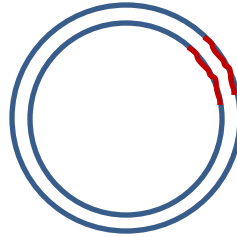
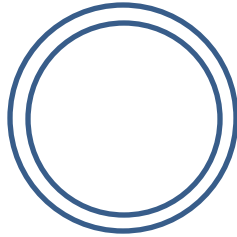
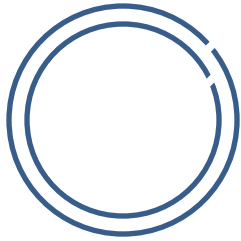
3. Plasmids

Vectors - Small pieces of circular DNA used for cloning

Requirements of the Vector

1. **Self-replication** - able to replicate in the host (independent origin of replication)
2. **Cloning site** - (region containing multiple restriction sites)
3. **Promoter** (and operator) - to support the expression of insert DNA (i.e. gene of interest) in the host.
4. **Selectable marker** – antibiotic resistance (Ampicillin resistant)
5. **Proper size**- for easy handling





Hosts for Recombinant DNA Technology

1. Bacteria

- *E. coli* - used because is easily grown and its genomics are well understood.
- Gene product is purified from host cells

2. Yeasts - *Saccharomyces cerevisiae*

- Used because it is easily grown and its genomics are known
- May express eukaryotic genes easily
- Continuously secrete the gene product.
- Easily collected and purified

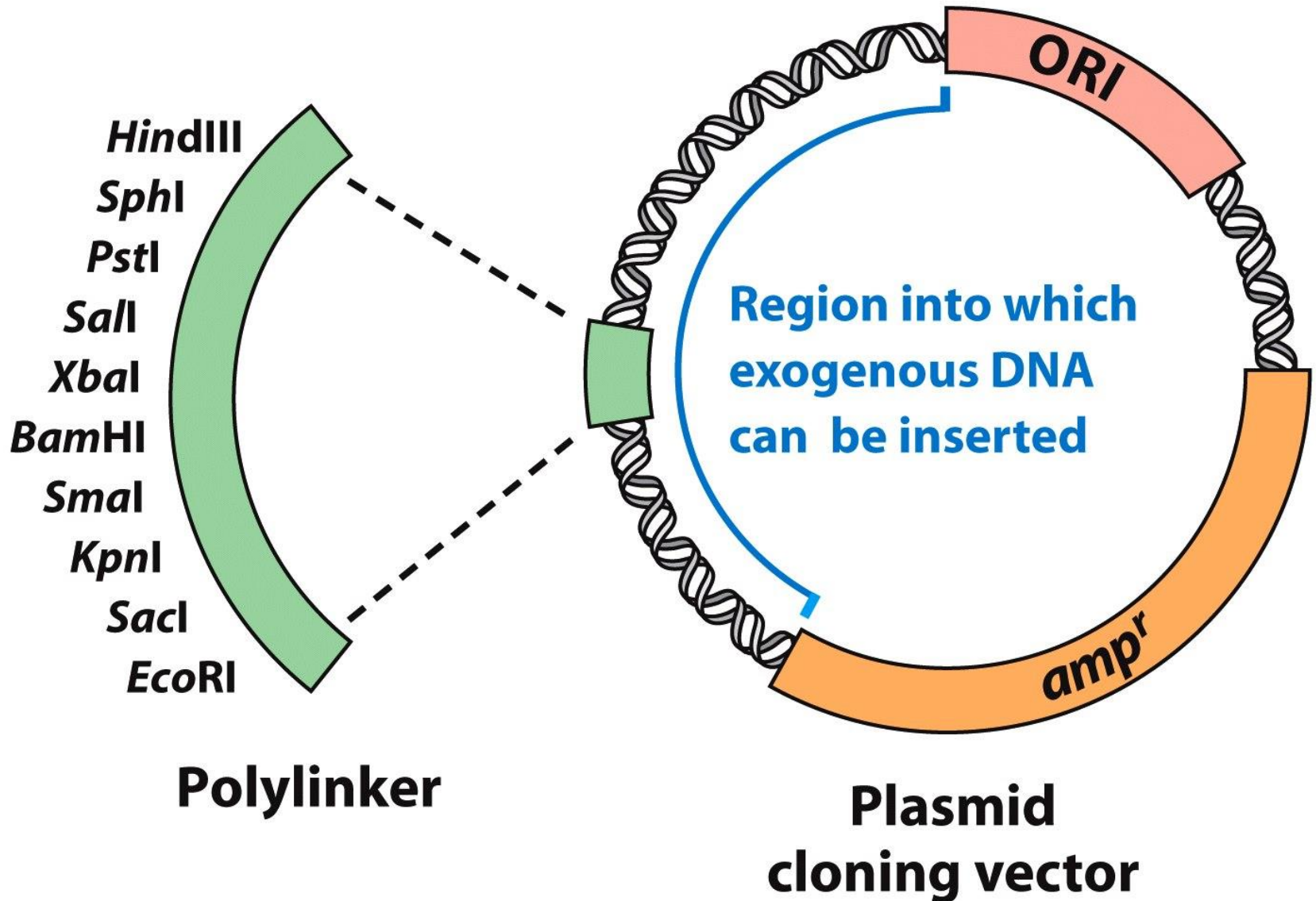
3. Plant cells and whole plants

- May express eukaryotic genes easily
- Plants are easily grown - produce plants with new properties.

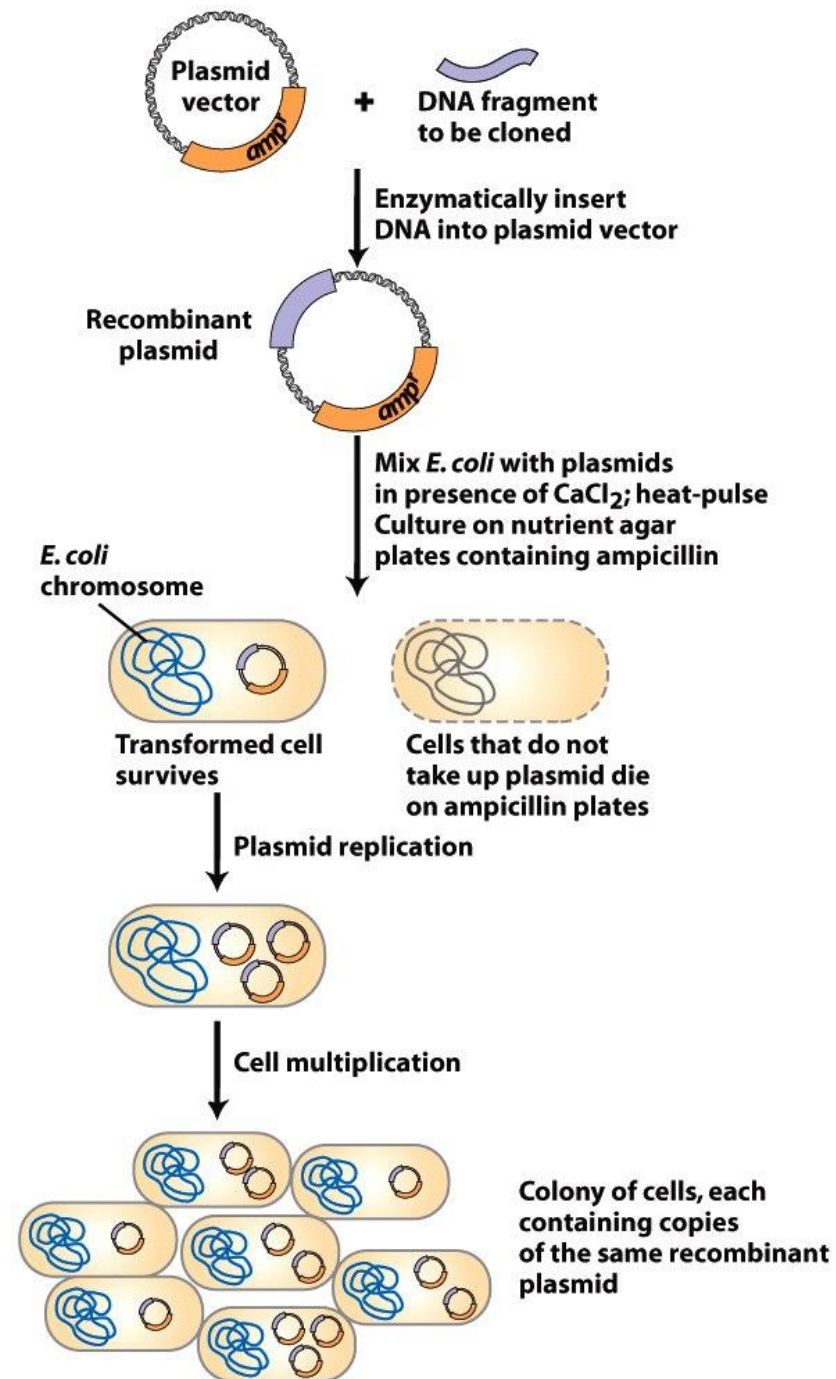
4. Mammalian cells

- May express eukaryotic genes easily
- Harder to grow
- Medical use.

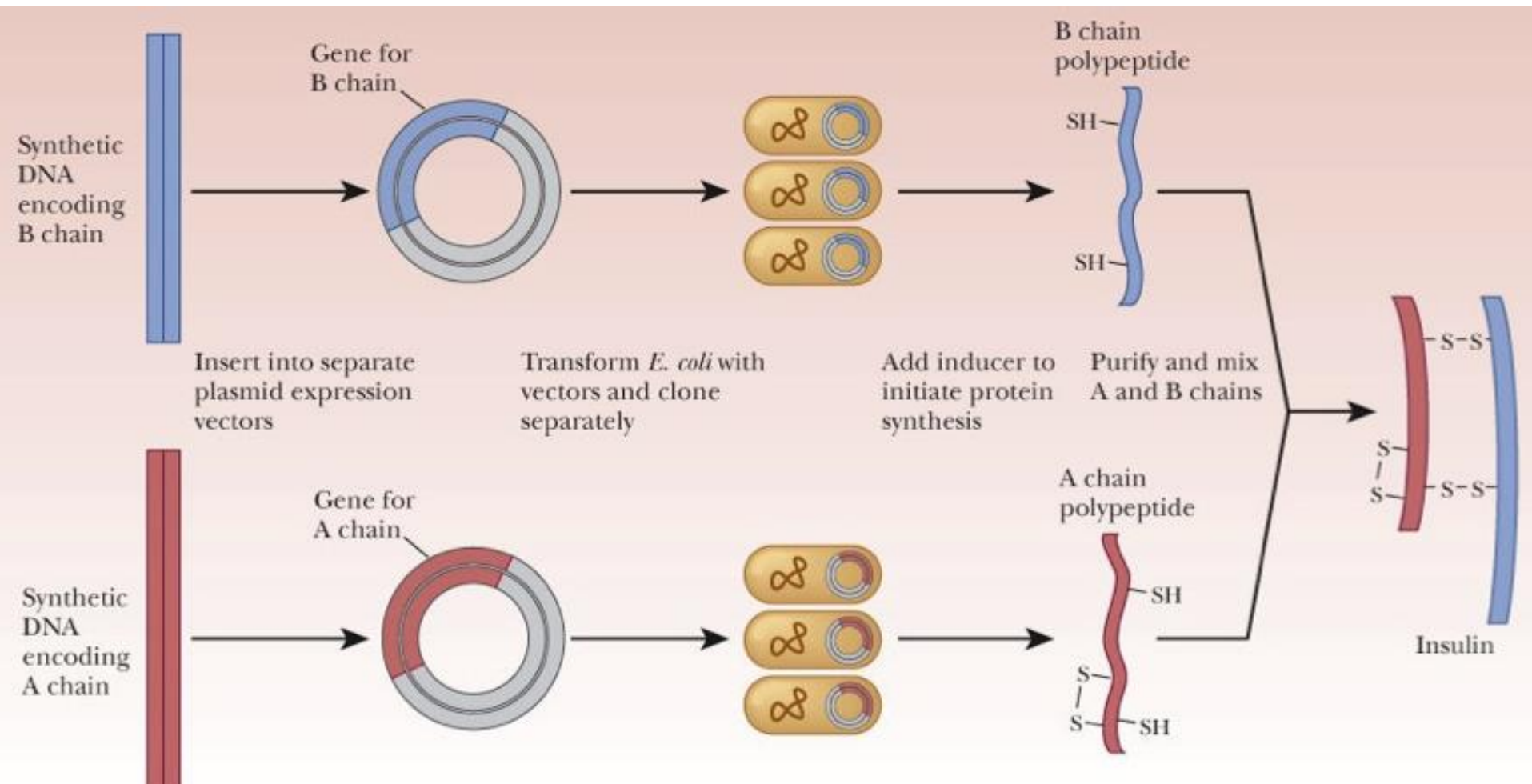
Cloning of a Gene



Cloning and Transformation of a Gene



Production of Insulin through Genetic Engineering Approach



Cloning Whole Animals

Two techniques

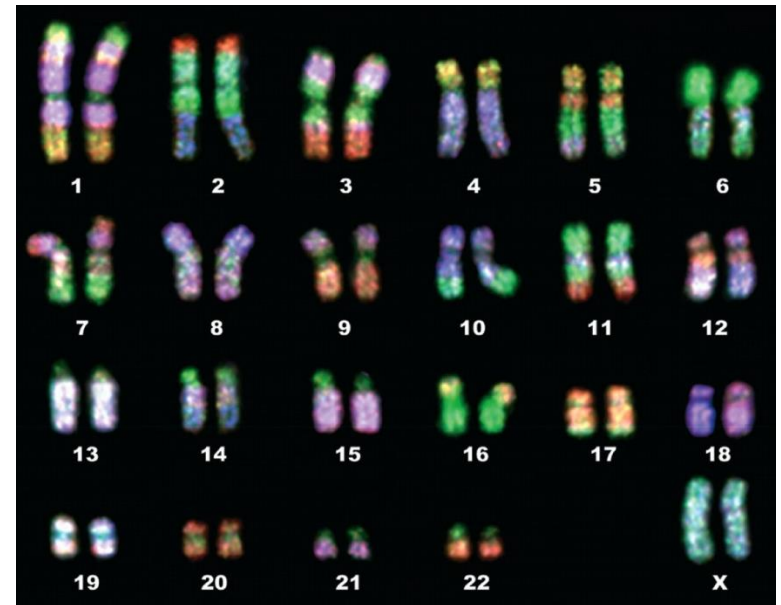
- Embryo splitting**
- Nuclear transfer**

Somatic cells are **diploid (2n)** and Gametes (sperm/ovum) are **haploid (n)**

- ❖ Most higher eukaryotes are **diploid (2n)**; i.e. their body (somatic) cells contain **two copies of the genome set** (two sets of homologous chromosomes)
- ❖ Their sex cells (gametes) are **haploid (n)** i.e. these cells contain **one copy of the genome set** (one set of each chromosomes)

Chromosomes

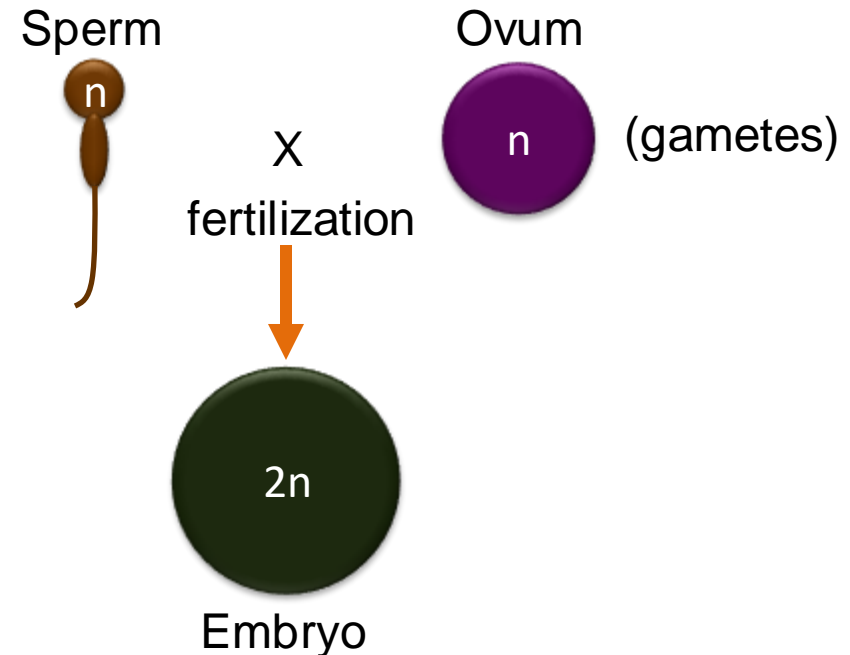
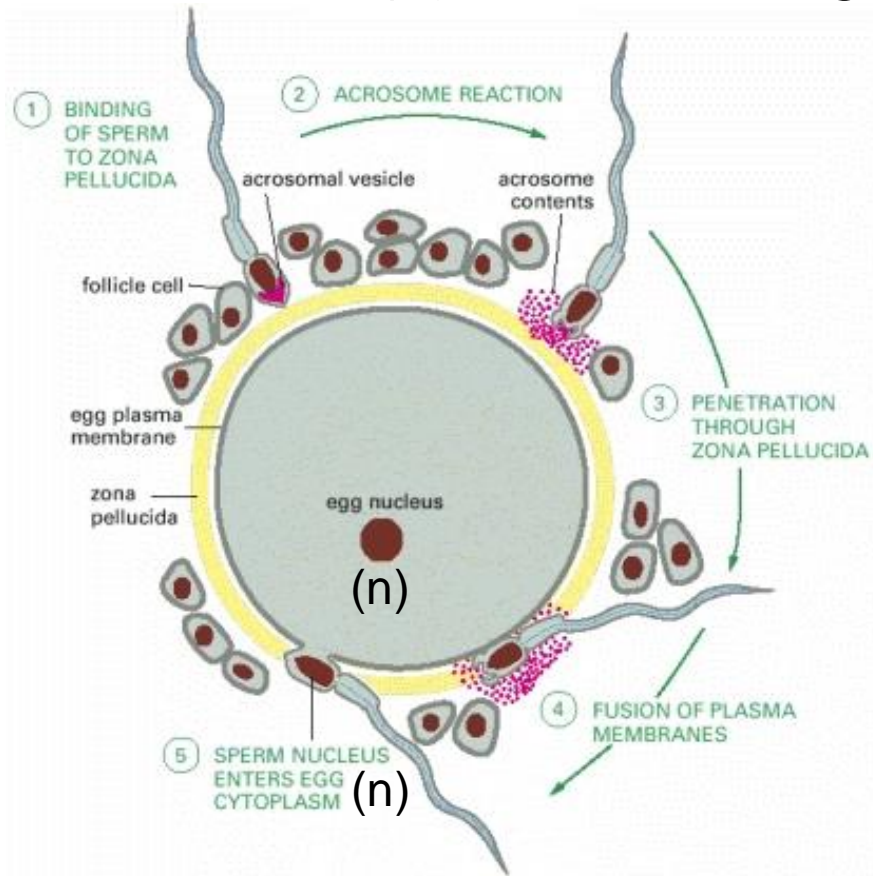
haploid	1 set
diploid	2 sets
triploid	3 sets
tetraploid	4 sets
hexaploid	6 sets
octaploid	8 sets



Diploid set of human chromosomes

How does the '2n' genome arise in embryo?

Through **fertilization** of two sex cells (gametes): one basic genome set (n) from male gamete (father's sperm) and another set (n) from female gamete (mother's egg).



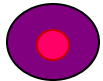
❖ So gametes need to be haploid (n)

➤ How do we get haploid gametes in a diploid organism?

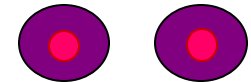
In eukaryotic organism, **two different types** of cell divisions occur

Mitosis (equal division): When the somatic (body) cells just increase in number.

One cell $\xrightarrow{\text{(genome duplication \& segregation into two)}}$ Two cells



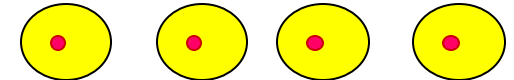
$$2n \xrightarrow{\text{(genome duplication \& segregation into two)}} (4n) \xrightarrow{\text{(genome duplication \& segregation into two)}} 2n + 2n$$



$$n \xrightarrow{\text{(genome duplication \& segregation into two)}} (2n) \xrightarrow{\text{(genome duplication \& segregation into two)}} n + n$$

Meiosis (reduction division): For sexually reproducing diploid organism specialized diploid cells (meiocytes) undergo two sequential nuclear divisions to form four haploid cells.

One cell $\xrightarrow{\text{(genome duplication \& segregation into four)}}$ Four cells



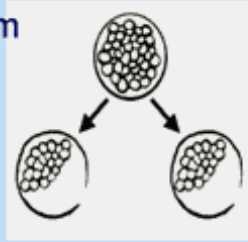
$$2n \xrightarrow{\text{(genome duplication \& segregation into four)}} (4n) \xrightarrow{\text{(genome duplication \& segregation into four)}} (2n) + (2n) \xrightarrow{\text{(genome duplication \& segregation into four)}} n + n + n + n$$

These haploid cells are called gametes (sperms and eggs in plants, animals) or spores (fungi, algae).

Embryo Splitting

Cloning by Embryo Splitting

Embryo is split to form two half-embryos



Embryos are transferred to an unrelated surrogate mother

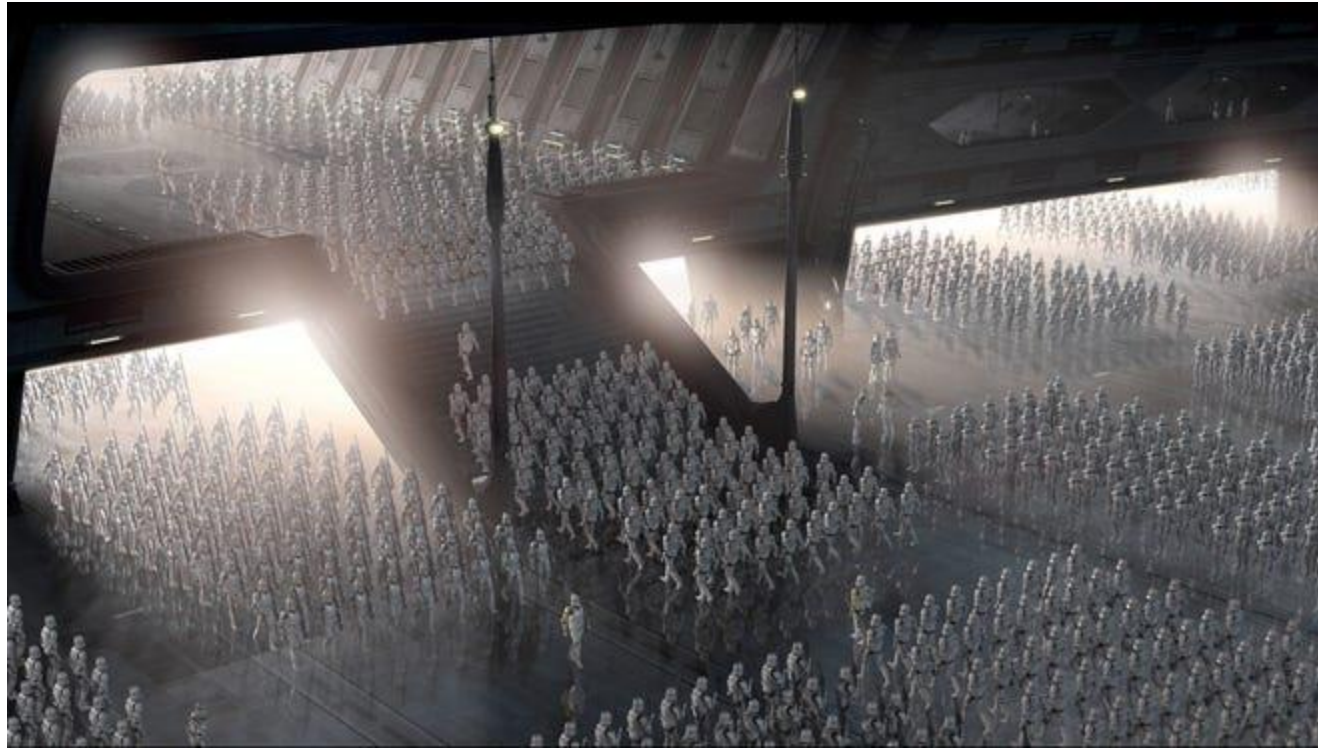


Pregnancy is monitored by ultrasound

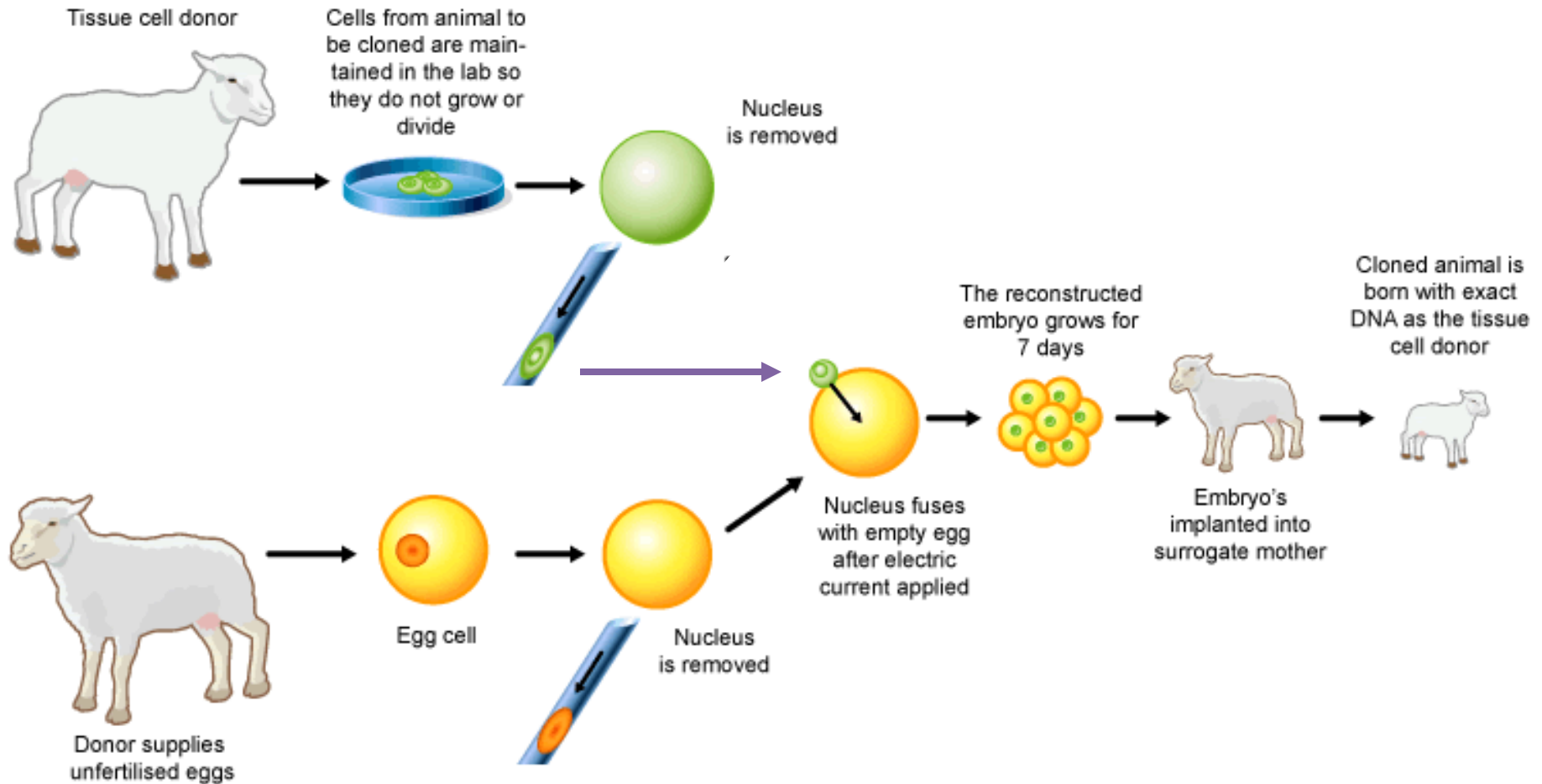


Sheep gives birth to identical twins

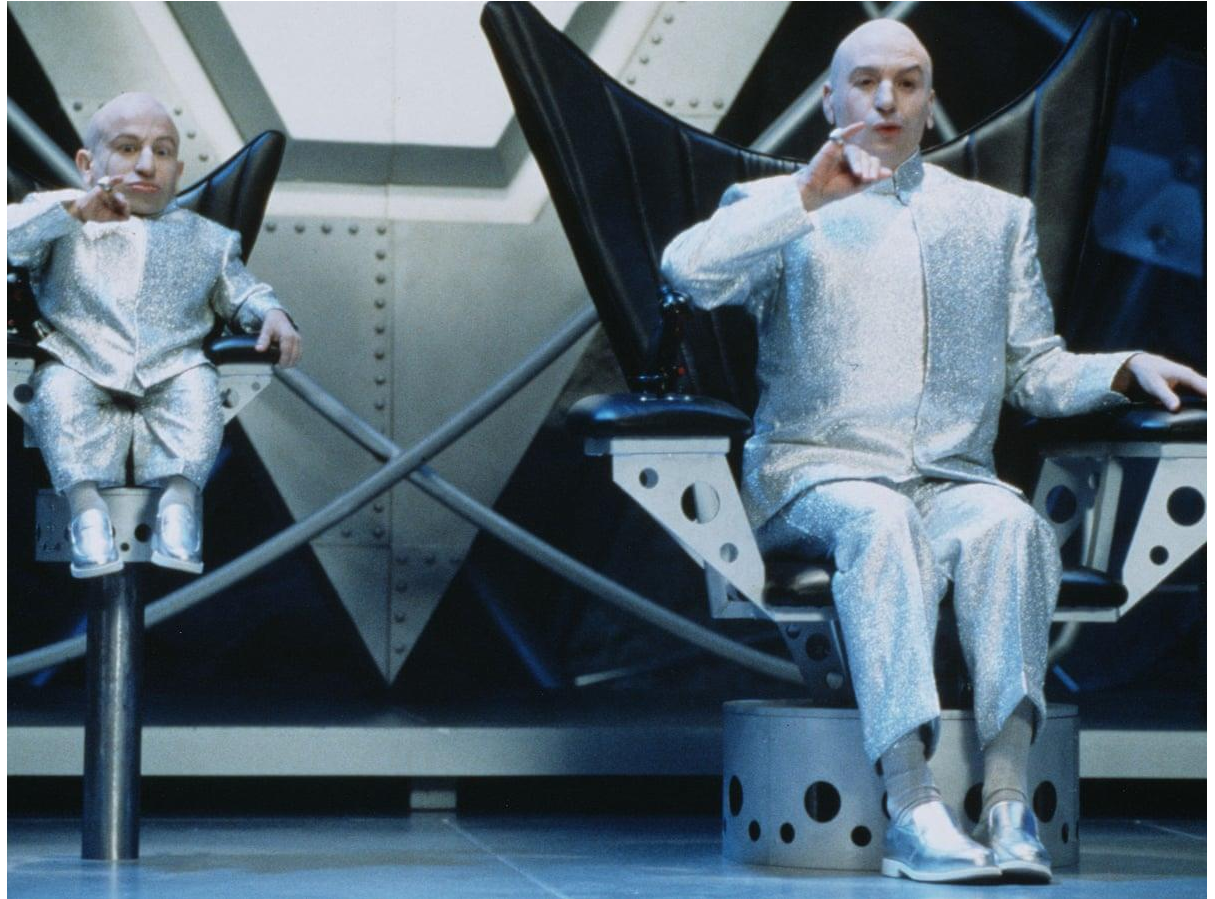
- Egg collected
- Fertilized by *in vitro* fertilization (IVF)
- Embryo is grown to 8–16 cells
- Cells are separated
- Separated cells grown into separate embryos
- Embryos transplanted into surrogate mothers



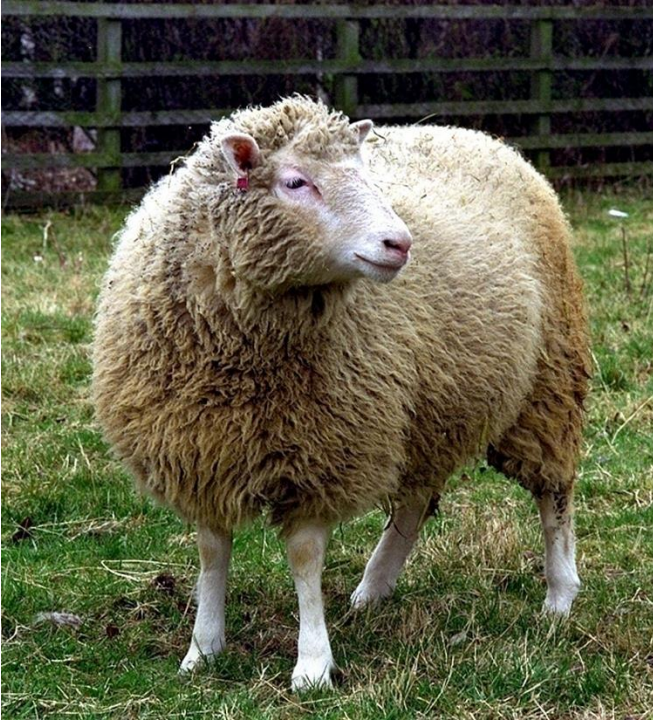
Cloning Whole Animals



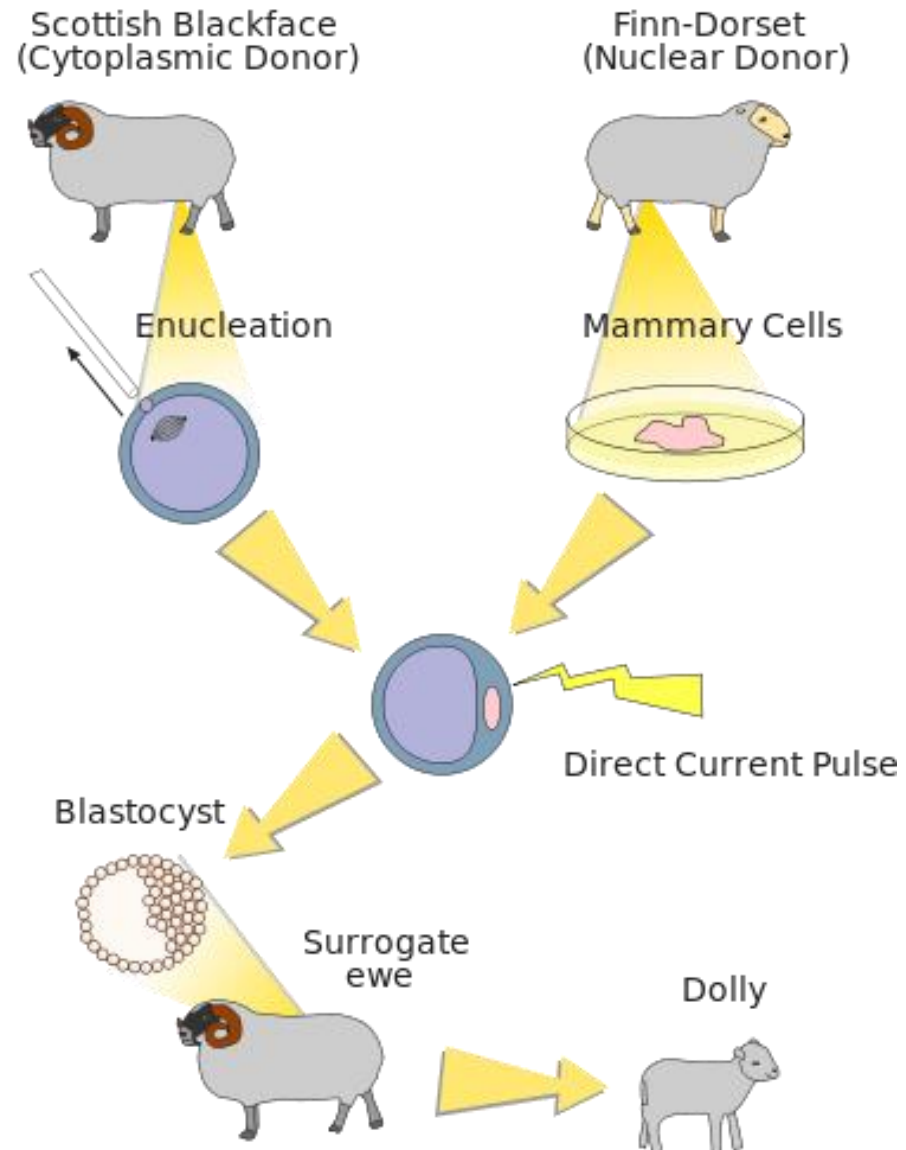
Nuclear Transfer



The First Cloned Mammal



**Born on 5th July, 1996
at Roslin Institute, Edinburgh,
Scotland**



The First Cloned Cat



In 2001, scientists at [Texas A&M University](#) created the first cloned cat known as Copy Cat/Cloned Cat (CC).

Selecting Best Possible Features from Parents: Science Fiction Film

	Directed by	Andrew Niccol
	Produced by	Danny DeVito Michael Shamberg Stacey Sher Gail Lyon
	Written by	Andrew Niccol
	Starring	• Ethan Hawke • Uma Thurman
	Music by	Michael Nyman
	Cinematography	Sławomir Idziak
	Edited by	Lisa Zeno Churgin
	Production company	Jersey Films
	Distributed by	Columbia Pictures
	Release date	•October 24, 1997
	Running time	106 minutes
	Country	United States
	Language	English
	Budget	\$36 million
	Box office	\$12.5 million ^[1]

The first cloned primate



Zhong Zhong and Hua Hua born November 2017

TECH

Experiments to gene-edit babies are ‘criminally reckless,’ says Stanford bio-ethicist

PUBLISHED MON, NOV 26 2018 • 3:27 PM EST | UPDATED TUE, NOV 27 2018 • 9:19 AM EST



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Gene-editing babies: Call to pause humanity-altering research

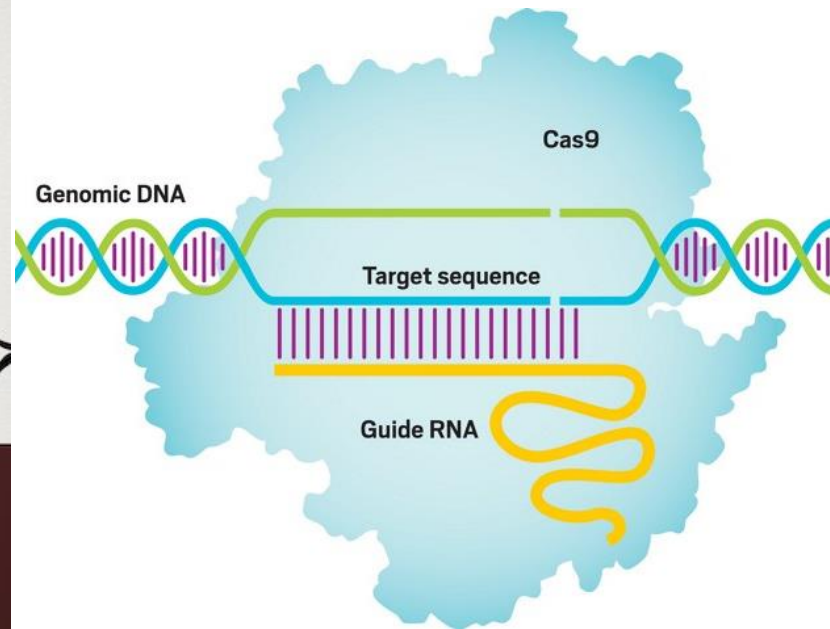
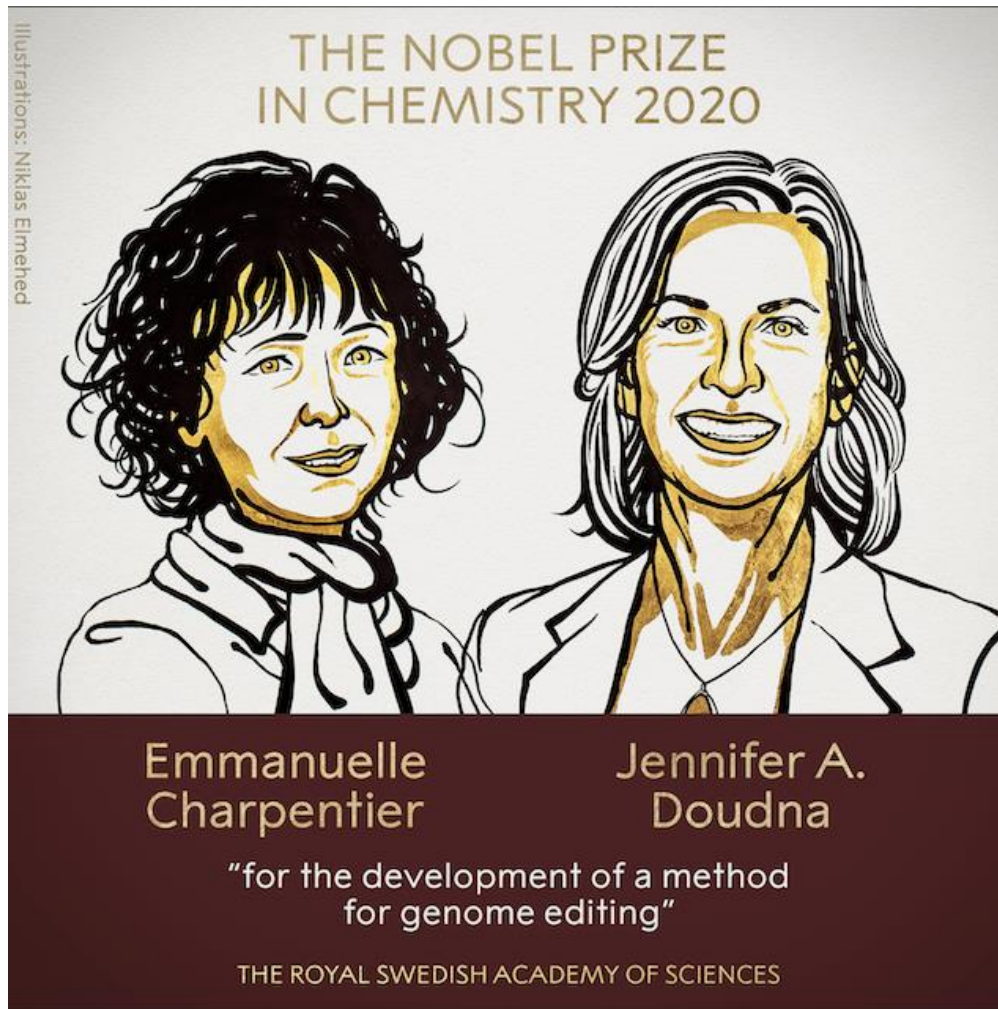
By James Gallagher
Health and science correspondent, BBC News

🕒 13 March 2019



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Nobel Prize for CRISPR-based Genome Editing Method



Thank You