

Applications of Animal Biotechnology

By

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Classical Biotechnology

- 19th and Early 20th century
- **Microbial-based fermentation process** such as alcohol, beer, wine, dairy products, various types of organic acids such as vinegar, citric acid, amino acids, and vitamins

Modern Biotechnology

- During the twentieth century
- Introduction of a large number of **new techniques** that has changed the face of classical biotechnology forever

Biotechnology definitions:

- ***Biotech*** in farm animals can be categorized as the biological, chemical and physical techniques that influence animal health (survival), nutrition, breeding, and reproduction.
- Controlled use of biological agents like cells or cellular components for beneficial use

Reproductive Biotechnology.....

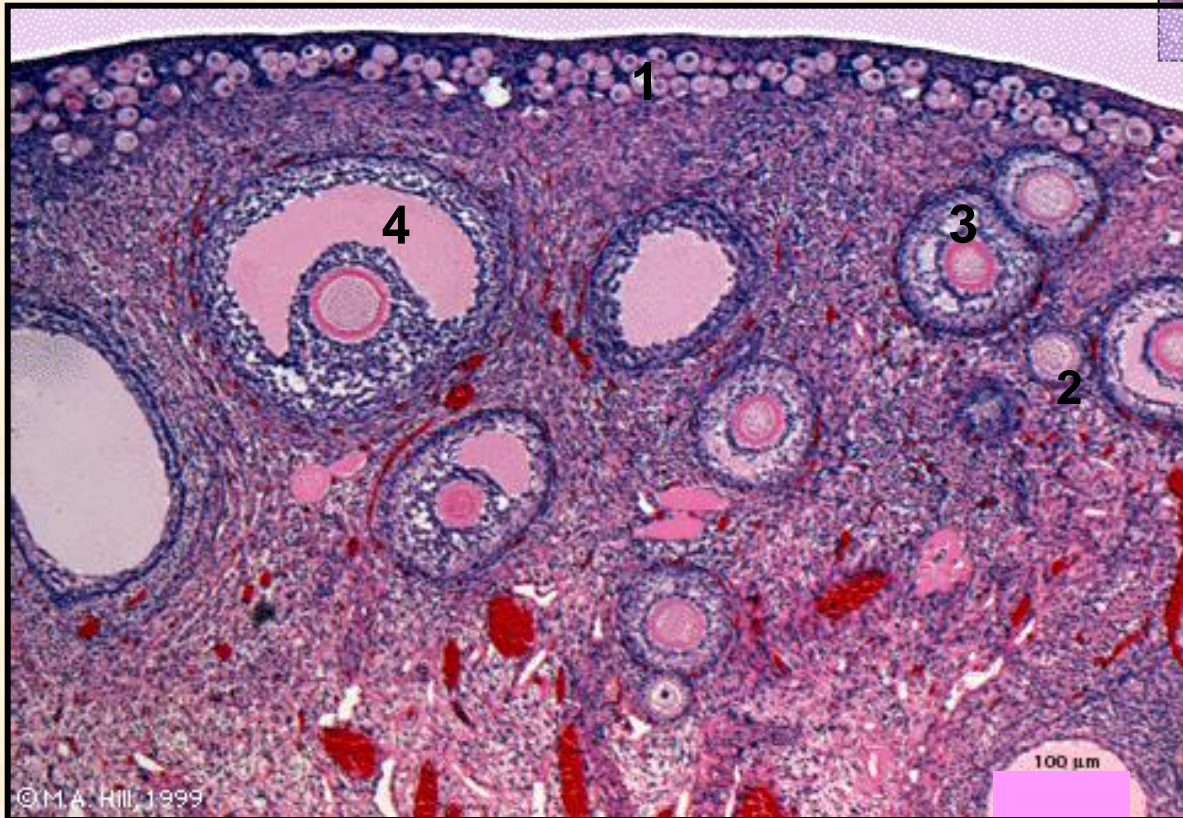
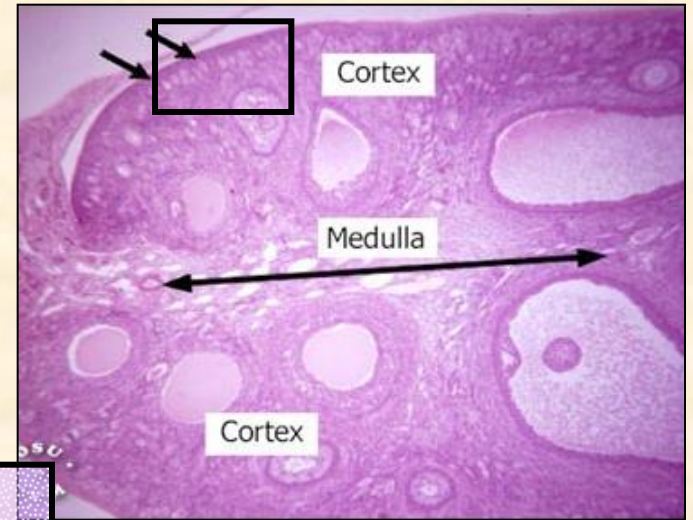
Why?

Reproductive Biotechnology.....

Why?

- To improve animal productivity and performance
- To overcome the increasing of human demands
- To increase the superior animal population
- To save endangered species
- To.....

Ovary

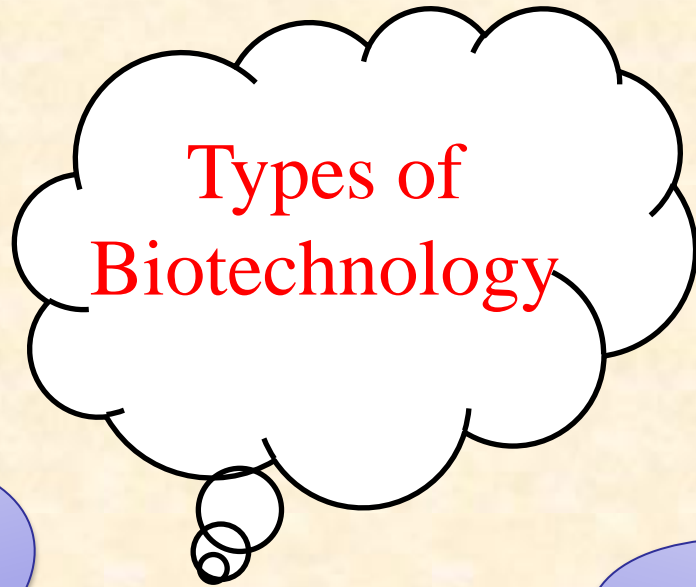


- 1 Premordial follicles
- 2 Primary follicles
- 3 Secondary follicles
- 4 Graafian follicles

Reproductive Biotechnology.....

Why?

The ovary, from birth, contains **a huge number** of follicles, the vast majority of these follicles become **atretic**, but they do represent a potential source of ova for embryo production.



AI
technique

Transgenesis

Cloning

Embryo
sexing

ICSI
technique

Cryopreservation

Superovulation

Embryo transfer

IVF
technique

Animal reproductive characteristics

Estrus cycle Producing one ovarian follicle (Graafian follicle) by the end of the cycle

- **Follicular phase**

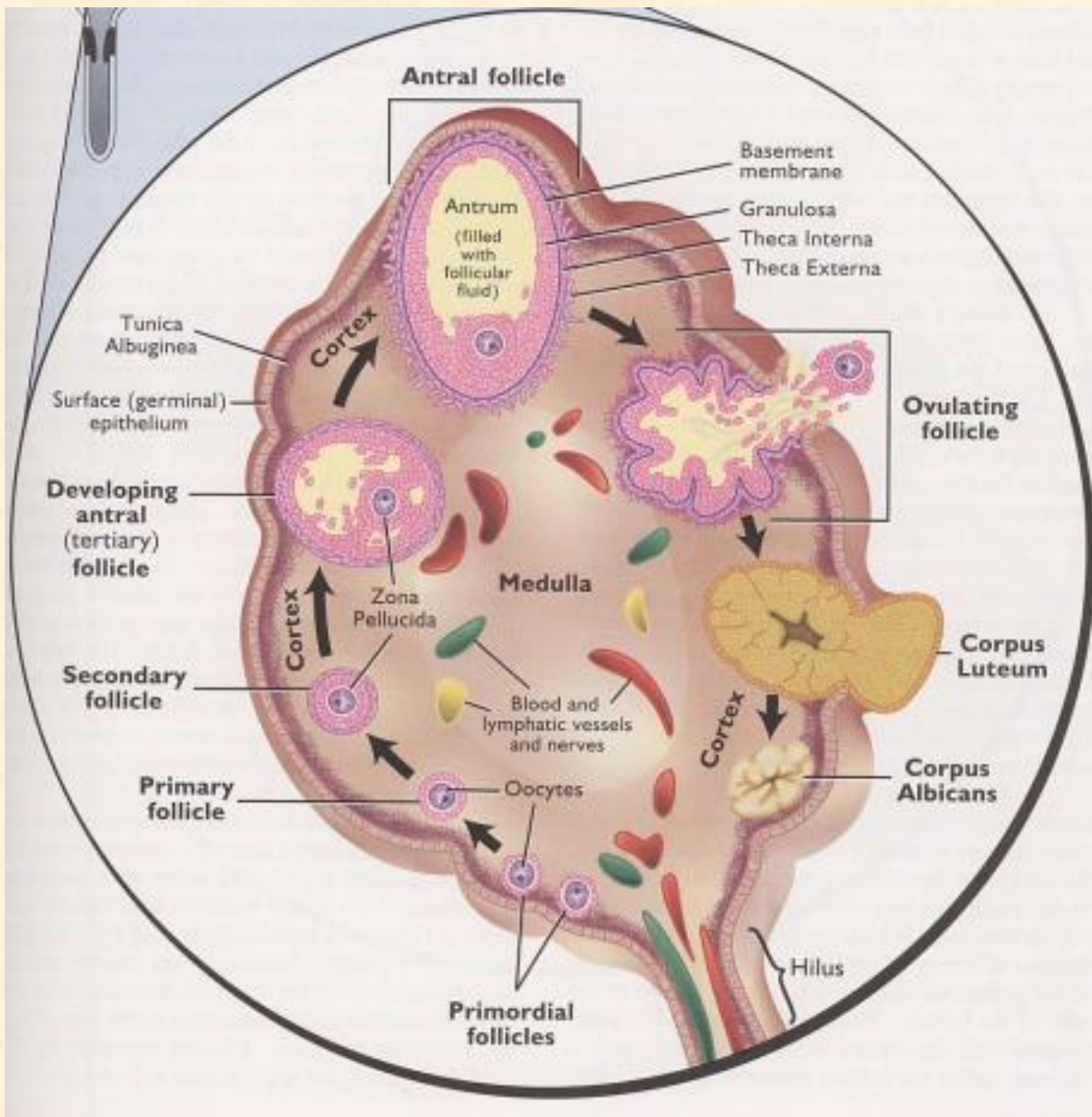
Ovarian follicles growth and development

FSH- Follicular Stimulating Hormone

- **Luteal phase**

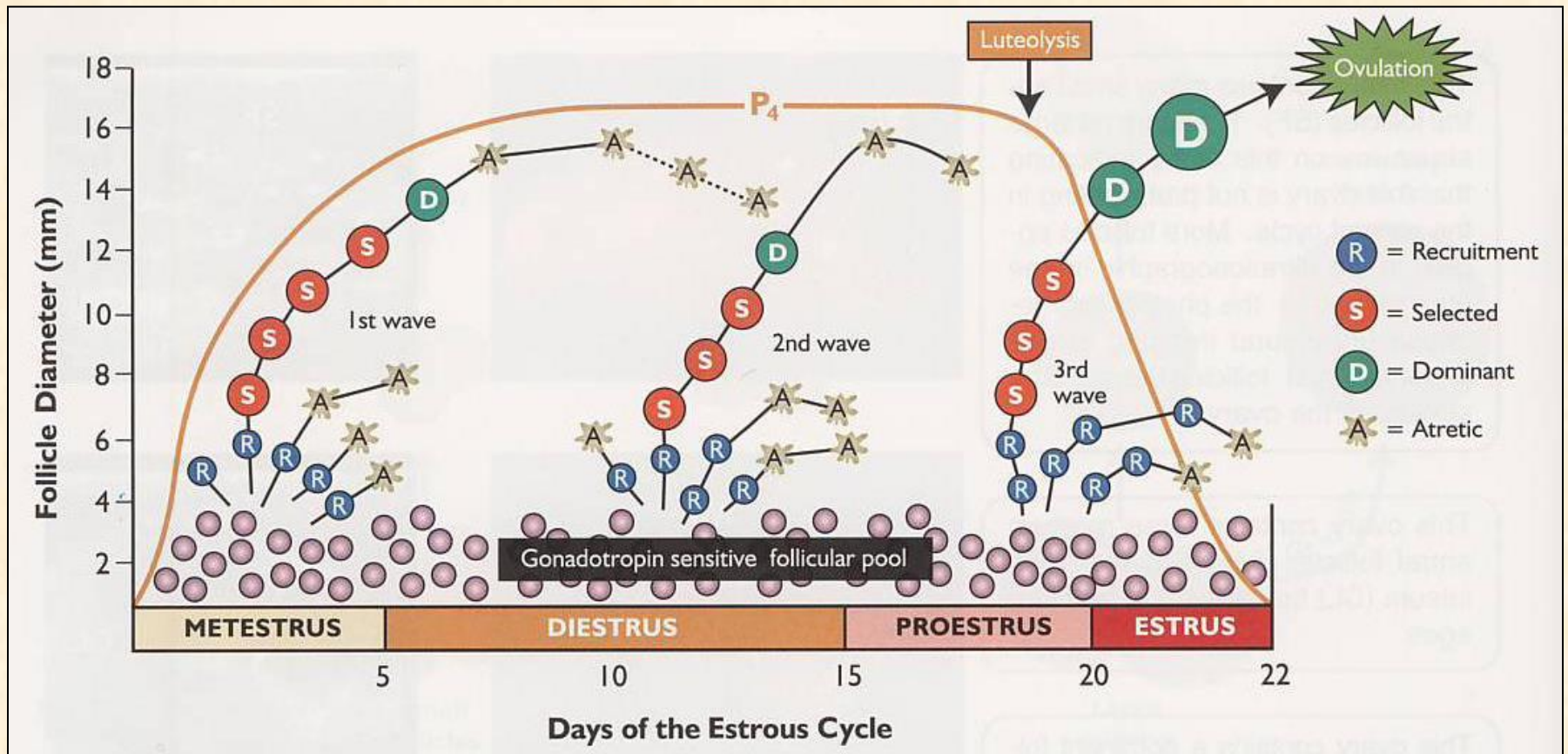
Corpus Luteum growth and development

LH- Luteinizing Hormone

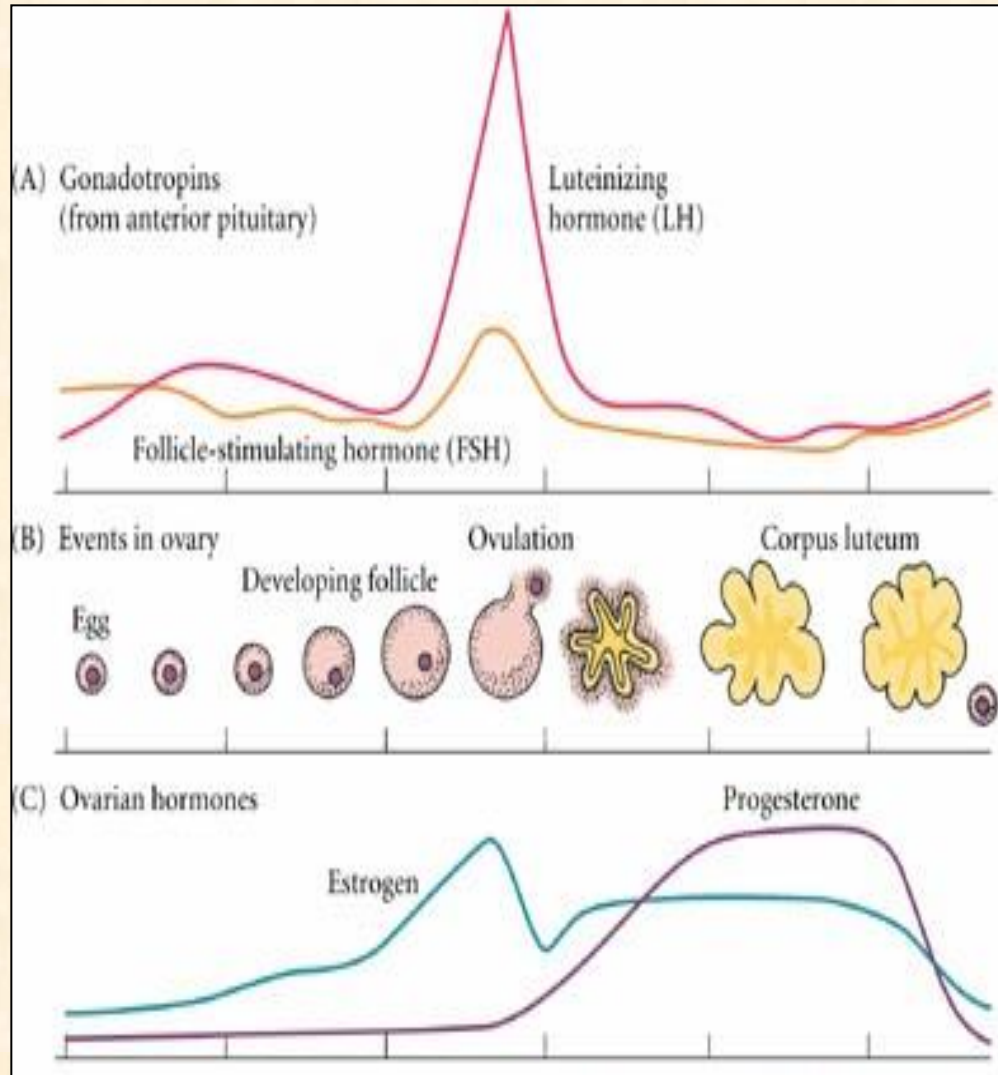


1- Superovulation:

The natural mechanism normally allow only one follicle to become dominant and ovulate.



Superovulation:



FSH

- The half-life of FSH is relatively short (0.5–1 hour)
- Repeated injections are required

PMSG (Pregnant Mare Serum Gonadotropins)

- FSH-like and LH-like properties
- longer half life in the body (50 hours)

Prostaglandins (PGF₂α)

The timing of ovulation
can be controlled!!!

Superovulation



Up to 20 oocytes



Fertilization

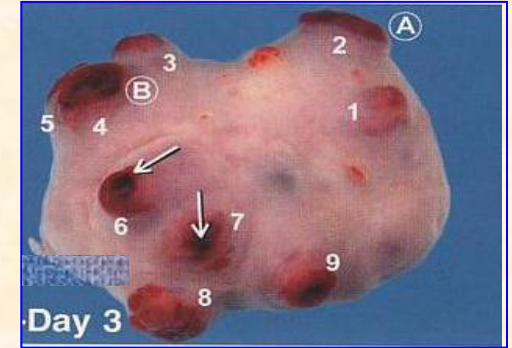
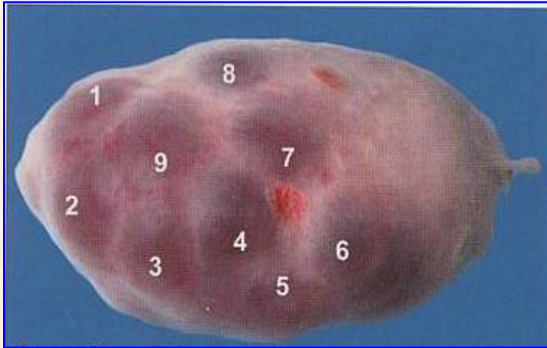
Embryos



Cryopreservation

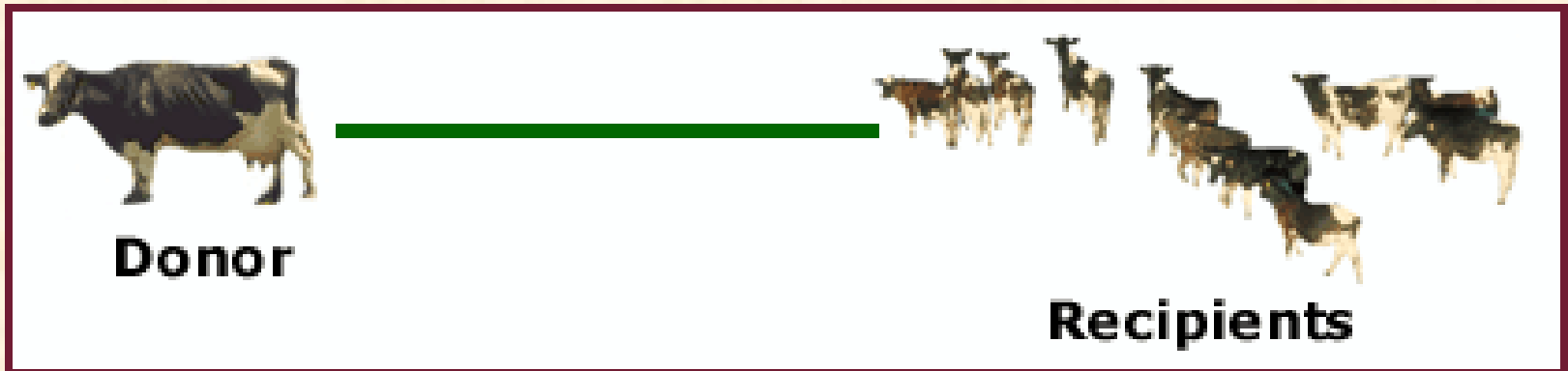
In Vitro
Fertilization

Embryo Transfer



2- Embryo Transfer (ET):

ET definition: is a process by which an embryo is collected from a donor female and then transferred into a recipient female where the embryo completes its development



2- Embryo Transfer (ET):

ET benefits:

1. Through ET a genetically superior female produces more offspring than she could by natural reproduction
2. ET offers more rapid genetic gain
3. Prevents disease transmission
4. ET preserves superior genetics for future generations due to embryo freezing
5. Frozen embryos can be shipped almost anywhere
6. Extend the productive life of some older cows

2- Embryo Transfer Steps:

1. Selection of the donor – Identifying the recipients cows
2. Synchronization of estrus cycle
3. Superovulation
4. Insemination – A.I
5. Flushing of the embryos – followed by embryos evaluation
6. Fresh embryos transfer or can be frozen

Donor
COW

Valuable donor



Recipients



Recipients
COW

PGF2 α



Synchronization
of oestrous cycles

PGF2 α



FSH
(PMSG)



Superovulation

A.I



Breeding
at oestrus

Oestrus

Embryo
evaluation



Recovery
of embryos



Transfer
of embryos

Normal breeding
or return after
two to three months
for another transfer
operation

Pregnancy



Valuable offspring

After two
days the
animal will
be in heat
(estrus)

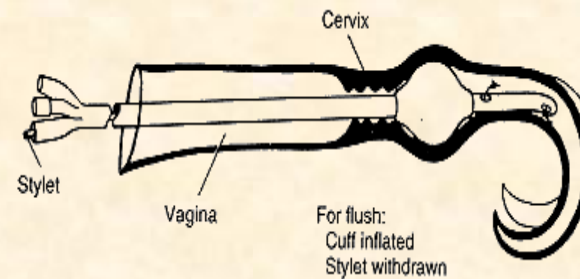
Donor cow:

- Insemination – A.I



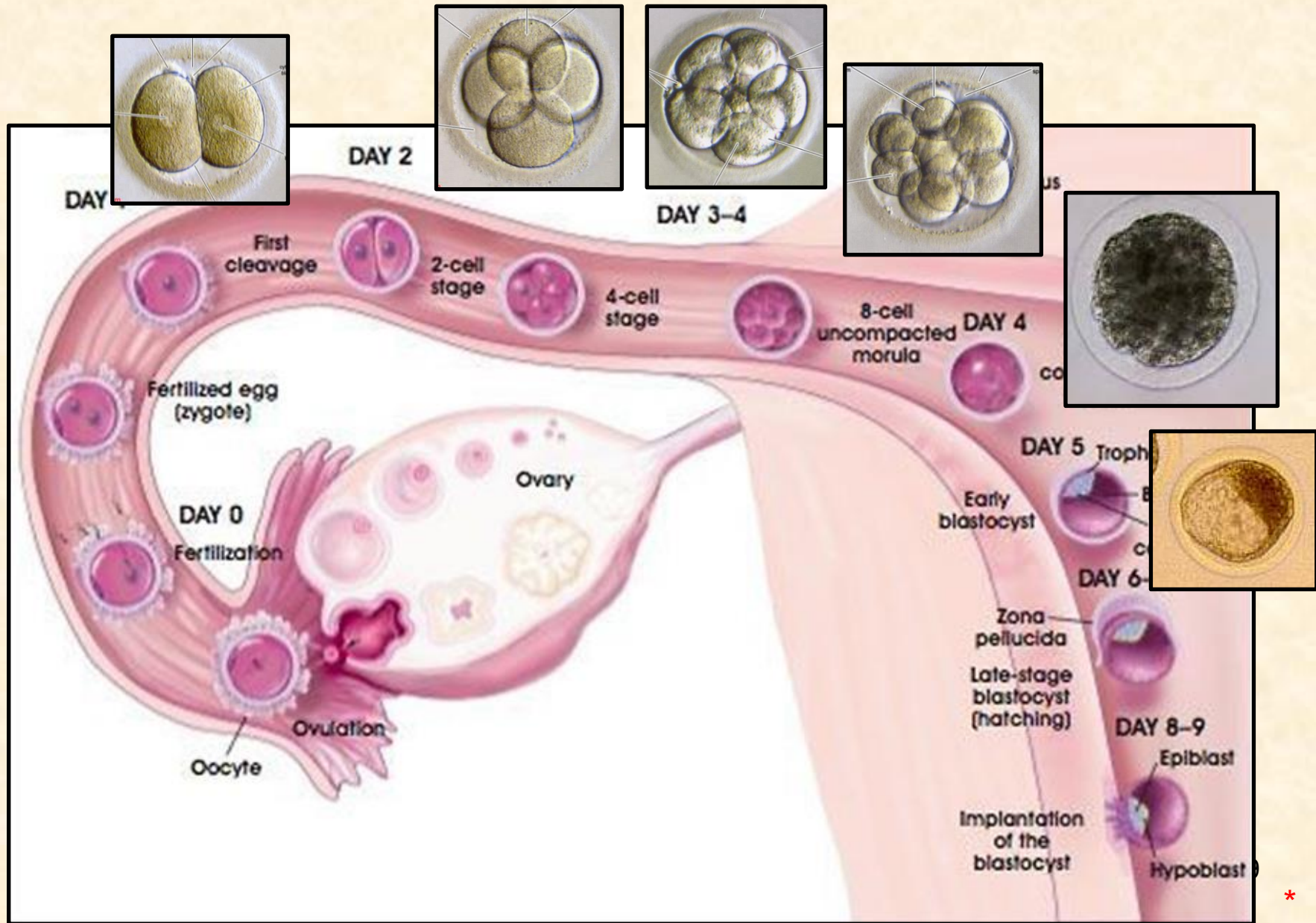
**Artificial
insemination gun**

- Embryo flushing



Foley catheter

Embryo development stages:



Evaluation of embryos

Embryos are evaluated according to their **stage of development:**

Stage 1: **Unfertilized**

Stage 2: **2 to 12 cells**

Stage 3: **Early morula**

Stage 4: **Morula**

Stage 5: **Early blastocyst**

Stage 6: **Blastocyst**

Stage 7: **Expanded blastocyst**

Stage 8: **Hatched blastocyst**

Stage 9: **Expanding hatched blastocyst**

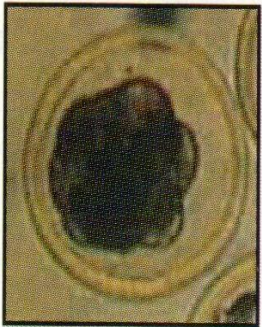
Embryos classified according to their **quality:**

Grade 1: **Excellent or good**

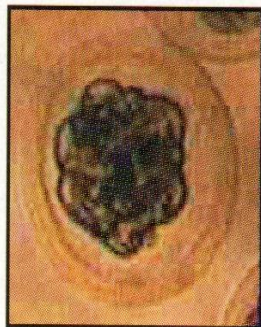
Grade 2: **Fair**

Grade 3: **Poor**

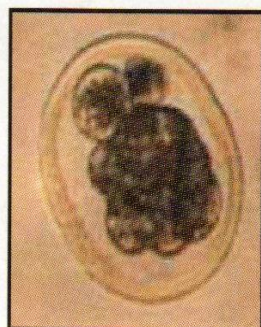
Grade 4: **Dead or degenerating**



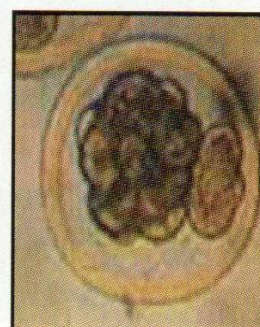
Cycle day: 6
Stage Code: 3
Quality Code: 1



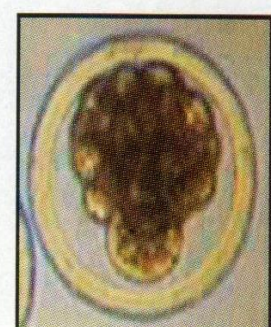
Cycle day: 6.5
Stage Code: 3
Quality Code: 1



Cycle day: 6.5
Stage Code: 3
Quality Code: 2



Cycle day: 6.5
Stage Code: 3
Quality Code: 2



Cycle day: 6.5
Stage Code: 4
Quality Code: 1



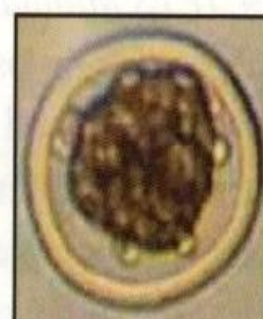
Cycle day: 7
Stage Code: 4
Quality Code: 1



Cycle day: 7
Stage Code: 4
Quality Code: 1



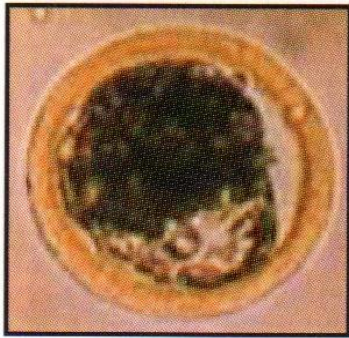
Cycle day: 7
Stage Code: 4
Quality Code: 1



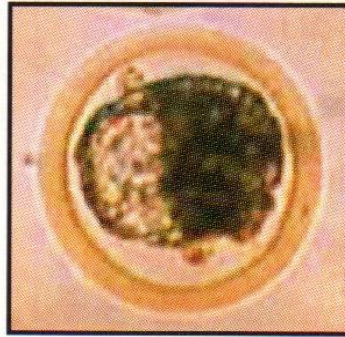
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Stage Code: 4
Quality Code: 1



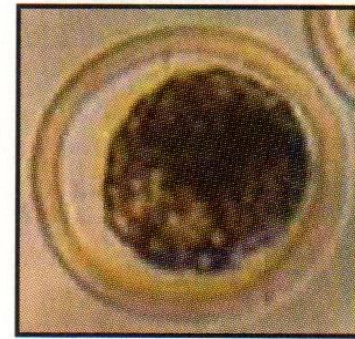
Cycle day: 7
Stage Code: 4
Quality Code: 1



Cycle day: 7
Stage Code: 5
Quality Code: 1



Cycle day: 7
Stage Code: 5
Quality Code: 1



Cycle day: 7
Stage Code: 5
Quality Code: 1



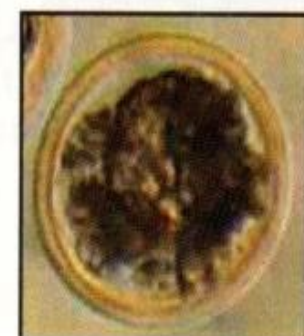
Cycle day: 7
Stage Code: 5
Quality Code: 2



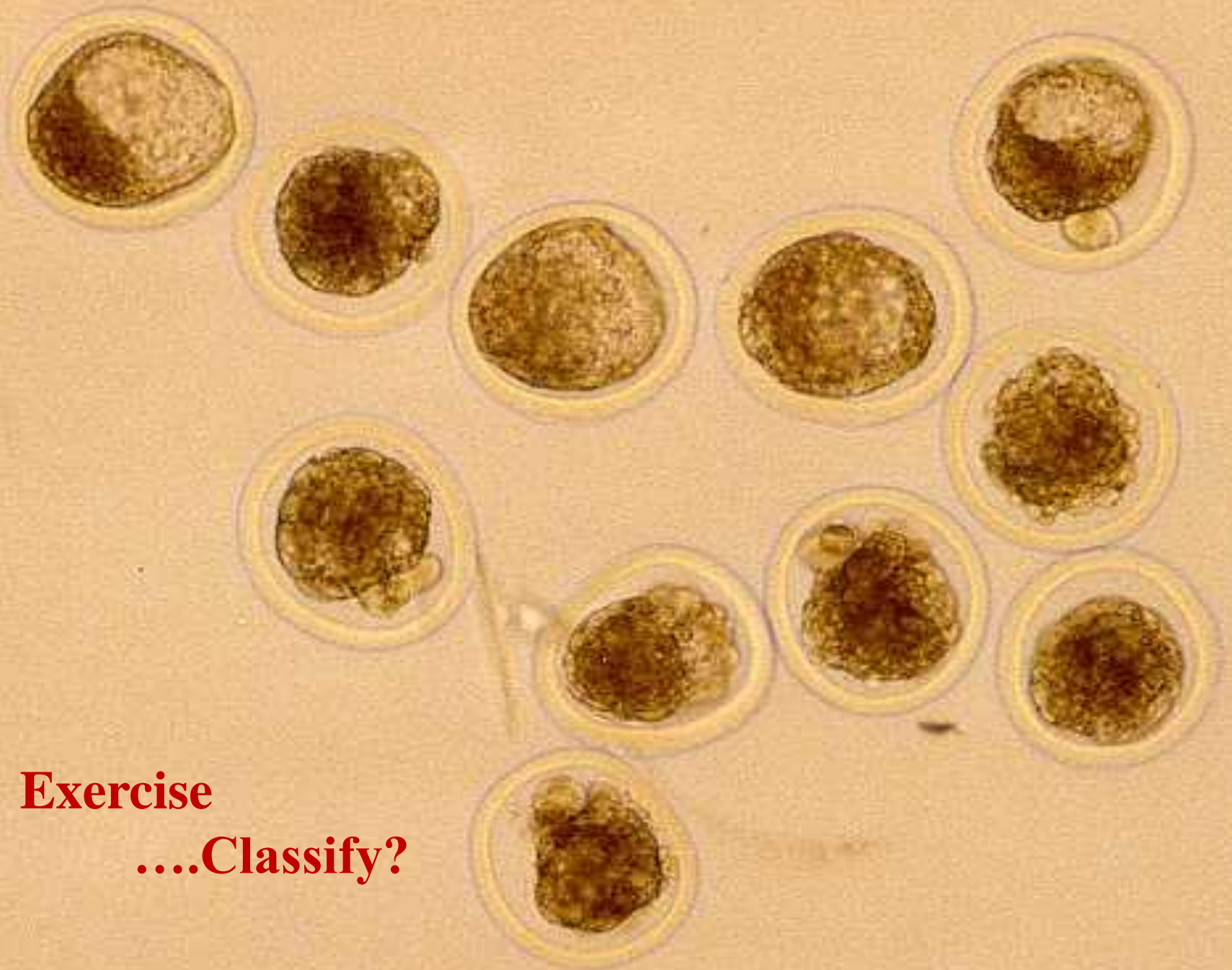
Cycle day: 7
Stage Code: 5
Quality Code: 2



Cycle day: 7
Stage Code: 5
Quality Code: 2



Cycle day: 7
Stage Code: 5
Quality Code: 3



Exercise
....Classify?

Embryo Transfer (ET):

•Limitations

Embryo transfer has failed to achieve the increased rates of genetic progress that had been predicted.



100 freezable embryos per donor per year

Because,

- *Not all potential donors respond positively to treatment*
- *On average about 60% of the embryos transferred result in pregnancies*

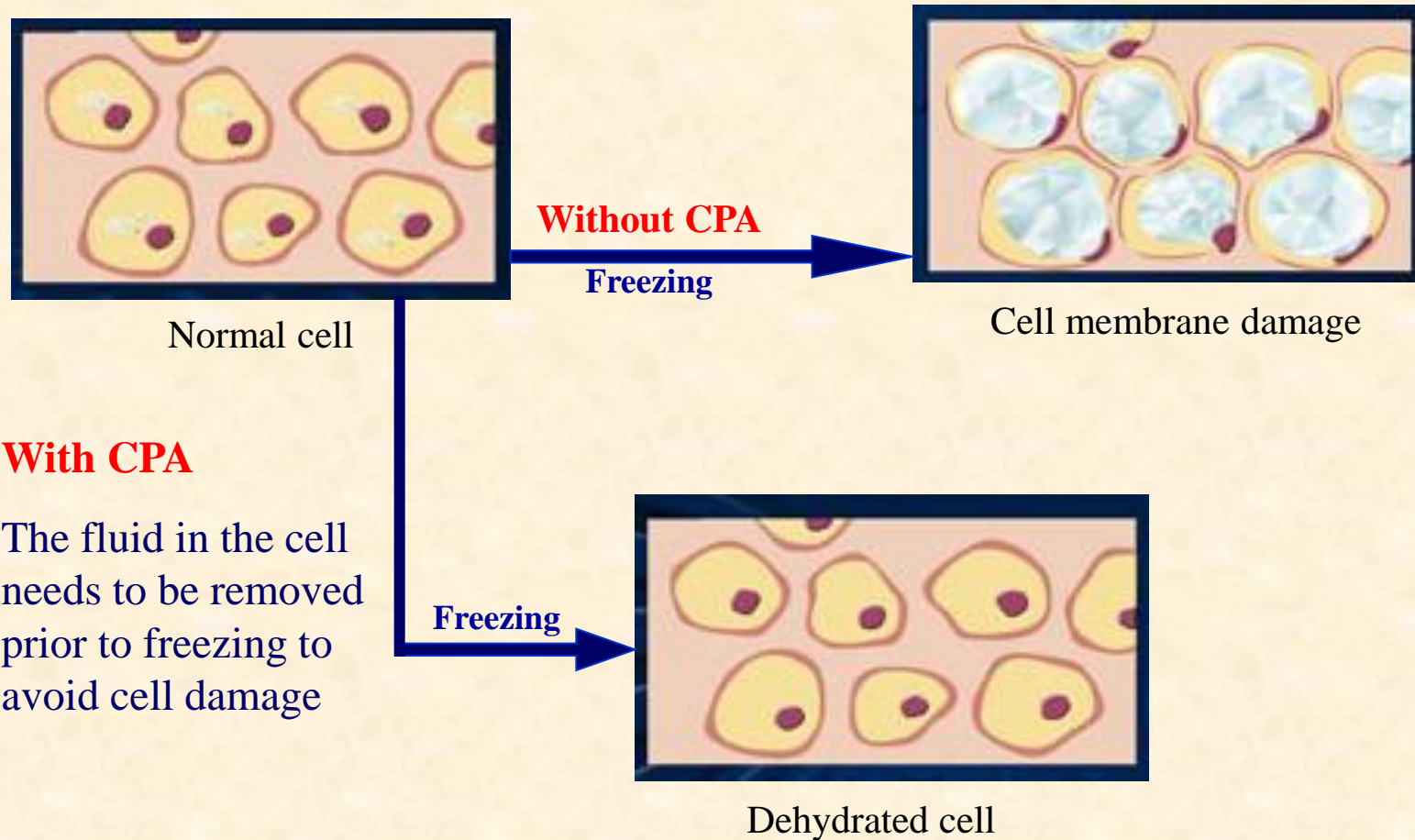
3- Cryopreservation:

- Is a process where cells or tissues are preserved by cooling to low sub-zero temperatures (-80°C or -196°C). At these low temperatures, any biological activity, including the biochemical reactions that would lead to **cell death is effectively stopped.**

Cryopreservation methods:

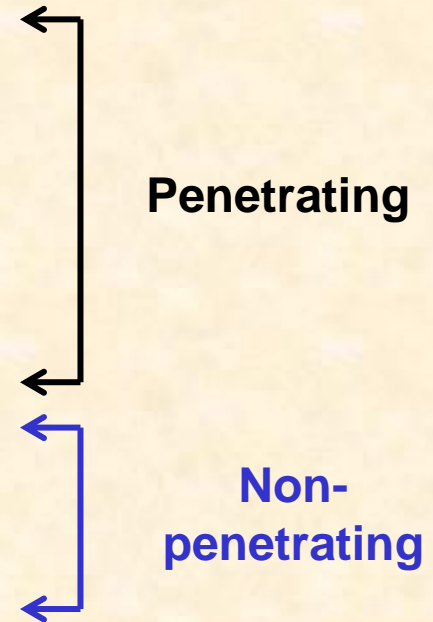
- Freezing (Slow and Rapid)

Role of Cryoprotectants :



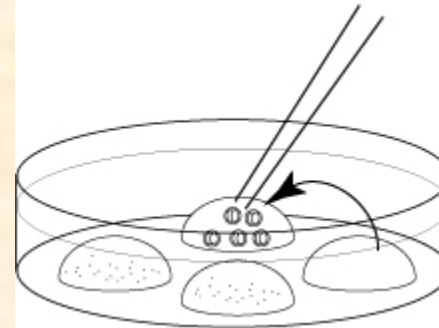
Types of Cryoprotectants :

- Glycerol
- Propanediol
- Dimethyl sulfoxide (DMSO)
- Ethylene glycol
- Sucrose
- Trehalose

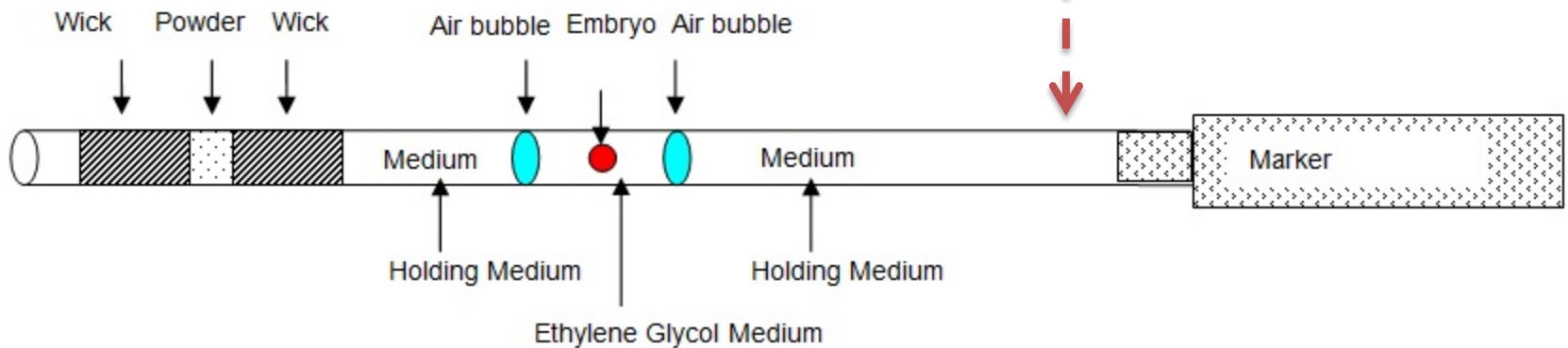


Embryo handling and freezing

- **Holding medium**
- **Freezing medium**



(Washing dish)
10 times



References:

- Ball, P.J.H. and Peters, A.R. (2004). ***Reproduction in cattle***. Blackwell Publishing Ltd, 9600 Garsington Road, Oxford OX4 2DQ, UK. 3rd edition
- Louis-Marie Houdebine (2003). ***Animal Transgenesis and Cloning***. John Wiley & Sons, Ltd. ISBNs: 0-470-84827-8 (HB); 0-470-84828-6 (PB).
- Senger, P.L. (2005). ***Pathways to pregnancy and parturition***. Washington State University Research & Technology Park. 2nd edition.