

A \* Types of cells + ECM

B \* Types of tissues

C \* Histological optical instruments.

Cell. biology  
1st lecture

Cell: Structural & functional Unit of living body.

It can be Uni-cellular like "Amoeba"

Eu-karyotic cell

Pro-karyotic cell

Its genetic material is located in "Genophore" Nucleoid

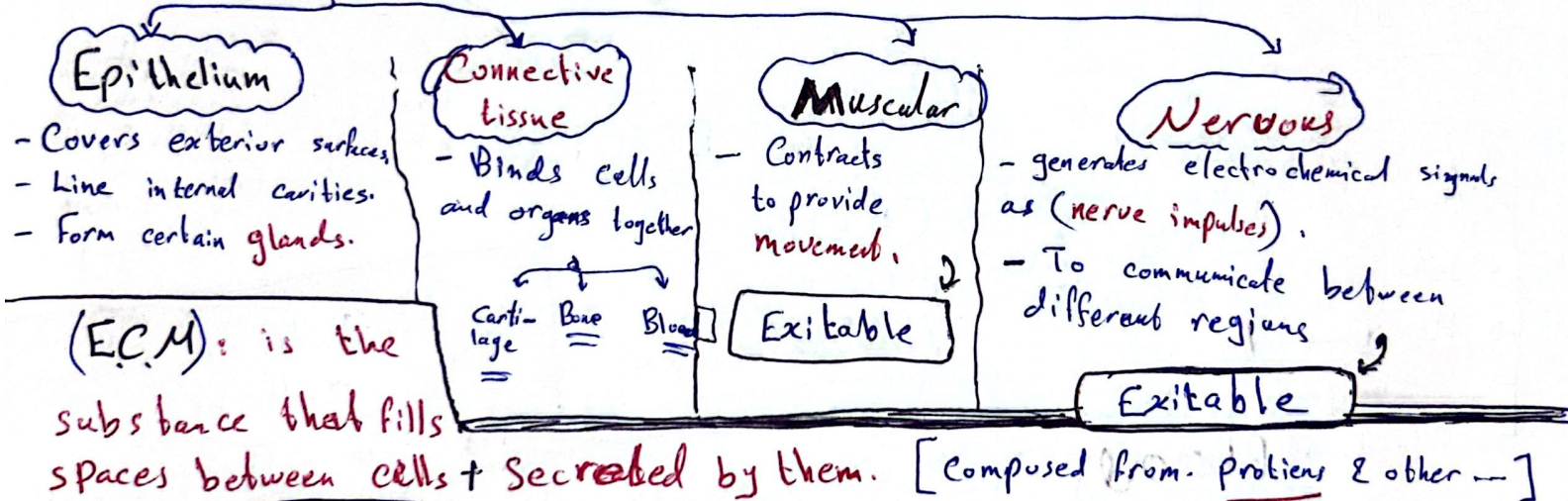
Reproductive mechanism: "Binary" "asexual" = fission

Features	Eu-karyote	Pro-karyote
Genetic mater.	✓ (genophore)	✓
Ribosomes	✓	✓
Cell membrane	✓	✓
Cytoplasm	✓	✓
membrane bounded cell organelles	✗	✓

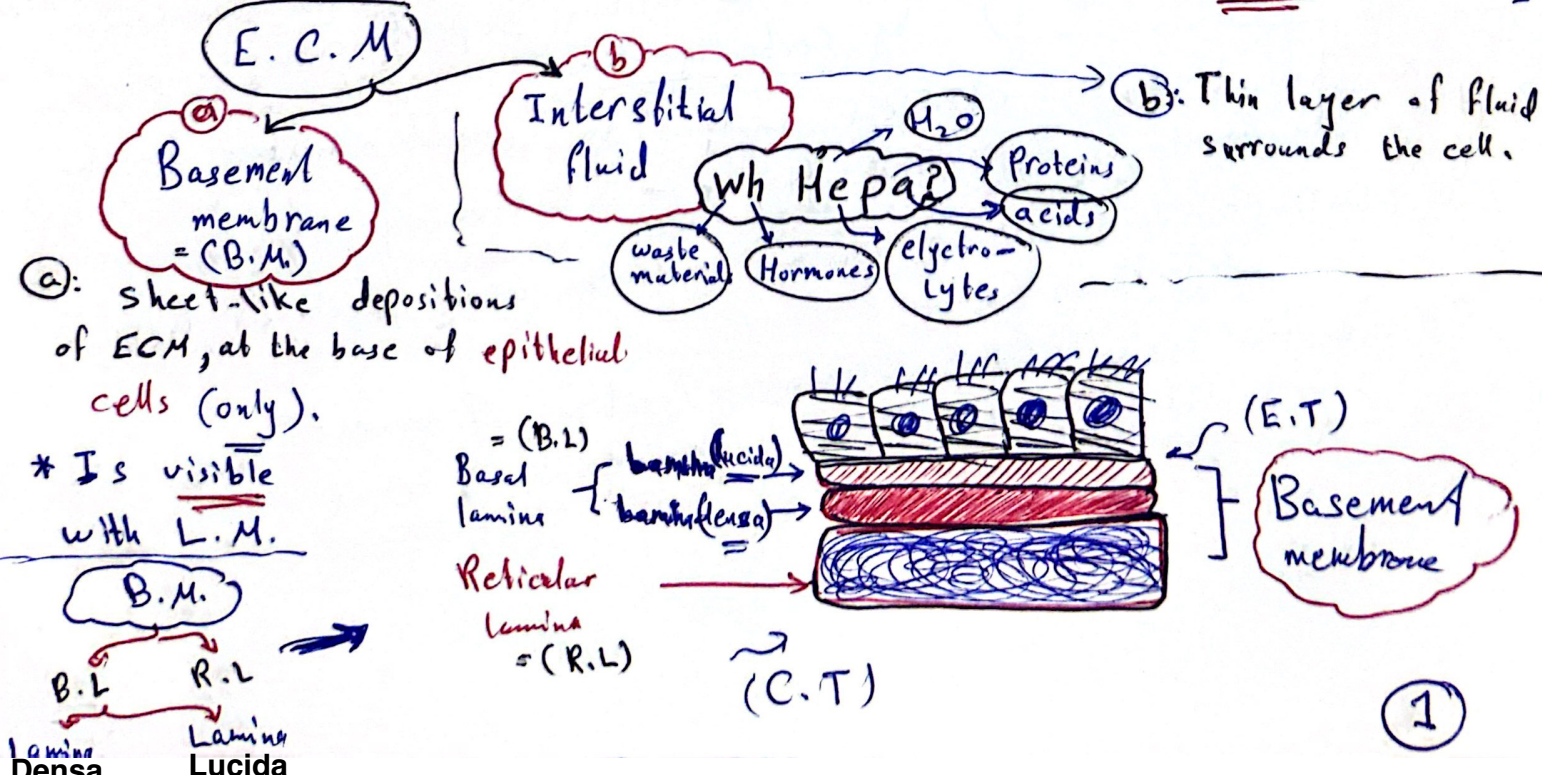
(4) only

\* There are 200 types.

Basic tissues (Types):



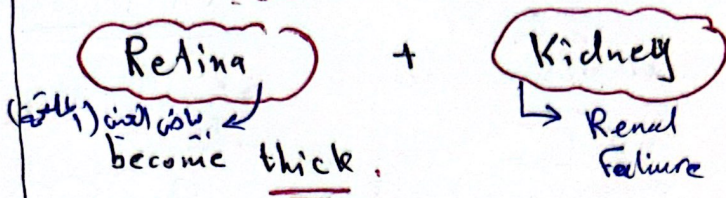
(ECM): is the substance that fills spaces between cells + secreted by them. [composed from proteins & other --]



■ The basal lamina:

- ① - visible with (E.M.)  
[ 20-100 nm ]
- ② - Its major component is: Collagen (IV)

\* In D.M. basement membrane of:



\* Epithelial tissue is:

avascular [but] Innervated

↓  
 (X) nourishment from C.T. (Connective tissue)

■ The reticular lamina:

- ① - Formed by (reticular fibers)  
 Secreted by C.T. (fibroblasts)
- ② - Thicker than basal lamina.

Functions of basement membrane

- ① Achoring
- ② Pathway of cell migration
- ③ Wound healing
- ④ Barrier between E.T. & C.T.
- ⑤ Blood filtration in kidney
- ⑥ Carcinoma in situ (Cancer early stage)

E.C.M

↓ ↓ in epithelium (E.T.)  
 ↑ ↑ in Connective tissue (C.T.)

- Jelly like → C.T. proper → loose/dense + adipose
- Hard → Bone
- Rubbery → Cartilage
- Fluid → ~~Blood~~

Methods of studying cell bio.

- 1\* Cell culture: Isolating cells & make them grow under controlled conditions.
- 2\* Cell fractionation: separate cellular components by centrifugation.

3\* Chromatography: chemical analysis to separate a mixture to its components. (2) phases: Mobile - Stationary

4\* Electrophoresis: separating charged molecules using electrical field depends on (size & charge)

5\* Genetic tech.: Study structure + function of genes:  
 - Isolating gene / cloning (copy) / determine unknown DNA sequences

# Microscopy: is the standard optical instrument for magnifying images.

## ① Light microscope (L.M.)

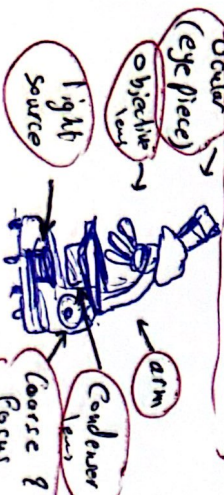
- \* Illuminating system: Light source + Condenser lens
- \* Imaging/Magnifying system:

- a - Ocular lens x 10x
- b - Objective // 4, 5, 90x

Microscope's capacity factors:

- ① Resolution power: Is the smallest distance between 2 particles that can still be seen as 2 separated.
- ② Magnification Power: The power to enlarge objects.

Healthy eye = 0.2 mm  
 L.M. = 0.2 μm  
 E.M. = 0.2 nm



## ② Electron microscope (E.M.)

- \* Section thickness: (0.01) μm
- \* Illuminating system: Electron gun (electron beam) - Condenser lens.
- \* Imaging system: Electromagnetic lenses 2-3
- Screen (Fluorescence)
- \* Lens are, Electro-magnetic
- \* Beam passes through Vacuum tube.
- Lenses → Objective (a) + Projecting (b)
- Refocuses electrons → Form image



## ③ DIC (3D) Phase Contrast

- Produces images for unstained (Transparent objects)
- Depends on the "Different refractive indices"
- Useful for: Living cells, Tissue cultures.

Fluorescence: Some substances absorb ultraviolet → Reflect it as visible long wave length (Fluorescence)

Fluorescence: Useful for: Blood cells, Sperm, DNA/RNA

DIC → Differential interference contrast microscope.

\* 3D images

\* Utilize 2 separate beams of light

## ④ Fluorescence

\* IDEA: Some substances absorb ultraviolet → Reflect it as visible long wave length (Fluorescence)

Illuminating sys: special lamp emits ultraviolet rays

Useful for: Proteins, antigen antibody complex

Antibodies are labeled with Fluorescence.

Uses ↑ optical resolution & contrast.

\* A computer system Reconstruct full image.

## ⑤ Confocal Laser

- \* Illuminating sys: Laser source
- \* Specimen should be labeled with Fluorescence.
- \* LASER passes through small hole? A. to avoid photo bleaching (fine details)

Heavy Salts used in E.M. Lead Nitrate, Uranium Acetate

③