

## Pathology Lec1 Summary ( Cell injury, death, and adaptations)

### General Overview

- Cells have internal compartments with specific functions that are regulated and maintained constant (homeostasis) to adapt with different stresses and demands from the surrounding environment.
- Cellular Adaptation is changes (increase/decrease) in the cell number, size, type, and accordingly the cell's function.
- Any stress would first cause cell adaptation, then it'll progress to reversible cell injury, further progression to irreversible cell injury meaning cell death which is either necrosis or apoptosis.

(Adaptation → reversible injury → irreversible injury (death))

- stress can directly cause cellular injury if the stimulus is injurious.

### Adaptation

- Types depending on the cause:
  1. physiologic (physiologic change).
  2. pathologic (disease).
- Forms/mechanisms of Adaptation:
  - 1-increase in cell size (hypertrophy).
  - 2-increase in cell number (hyperplasia).
  - 3-decreased in cell size (atrophy)
  - 4-change in cell type.

### Hyperplasia

- increased in cell **number** in tissues that have a proliferative ability.
- Can be pure or mixed.
- Types:
  - 1-Physiologic:
    - A. hormonal stimulation e.g. hyperplasia in breast glandular tissue (puberty or pregnancy).
    - B. compensatory e.g. liver hyperplasia after liver resection.

2-Pathologic: (give arise to cancer)

A. Excessive hormonal stimulation:

e.g. endometrial hyperplasia due to continuous **estrogen** stimulation, benign prostatic hyperplasia caused by hyper **androgenic** stimulation.

\*endometrial hyperplasia causes endometrial carcinoma (cancer)

B. Viral infections e.g. warts caused by HPV virus.

## Metaplasia

- change in cell **types** by reprogramming stem cells to differentiate into a new cell type (Reversible process)
- Examples:
  - A. changes in respiratory epithelium that lines the bronchi from ciliated pseudostratified into squamous.
  - B. epithelium that lines the esophagus changes from squamous to glandular in GERD.
- Causes:
  - 1-smoking. 2-vitamin A deficiency. 3-GERD (gastroesophageal reflux disease)

\*esophageal metaplasia causes esophageal carcinoma (cancer)

### Hypertrophy

vs.

### Atrophy

<b>increase</b> in cell size and functional capacity by increasing the production of structural proteins and organelles.	<b>decrease</b> in cell size and function by <ul style="list-style-type: none"><li>1)decreasing in protein synthesis</li><li>2)increase in protein degradation and</li><li>3)autophagy.</li></ul>
1.physiologic (hormonal, growth factor stimulation or increased functional demand).  2.pathologic	1.physiologic: e.g. endometrial atrophy caused by loss in hormone stimulation in menopause. 2.pathologic: caused by denervation injury, or chronic ischemia resulted from arteriosclerosis or diabetes.
Can be:  1.pure (happens in cells that can't divide like cardiac, skeletal muscles)  2.Mixed (accompanied with hyperplasia)	Causes: 1.decrease workload. 2. loss of innervation. 3.diminished blood supply (ischemia). 4.inadequate nutrition. 5.loss of endocrine stimulation. 6.aging.

## Hypertrophy Examples

Heart (cardiac muscles)		Uterus (smooth muscles)	Skeletal muscles
Pure		Mixed (accompanied with hyperplasia)	Pure
<b>Types</b>	Pathologic: Hypertension aortic valve stenosis	Physiologic: Estrogenic stimulation during pregnancy	Physiologic
<b>impact</b>	Can cause irreversible cell injury if untreated, resulting in myocardium degeneration due to lack of blood supply which leads to heart failure, ischemia, or myocardial infarction.	Reversible cell injury: Once the stress/demand is relieved	
<b>Side notes</b>	Purpose of hypertrophy: Increase the force of contraction.	-	-

## Cell Injury Causes

### 1. Oxygen deprivation (Hypoxia) by:

A. Ischemia (lack of blood due to artery blockage by a clot)

B. Pulmonary diseases: pulmonary hypertension, emphysema (obstructive pulmonary disease).

ischemia leads to hypoxia  
→ lack of blood leads to lack of O<sub>2</sub>

**2. Chemical Agents** (sugar, drugs, pesticides, and insecticides)

**3. Infectious Agents** (viruses, bacteria, protozoa parasites, worms)

**4. Immunologic Reactions** (Autoimmune, microbes, allergic rxns: rhinitis, conjunctivitis, eczema)

**5. Genetic factors** (chromosomal and gene mutations)

**6. Nutritional Imbalances** (excess nutrition or malnutrition)

**7. Physical Agents** (trauma, extreme temp., electric shock)

**8. Age**