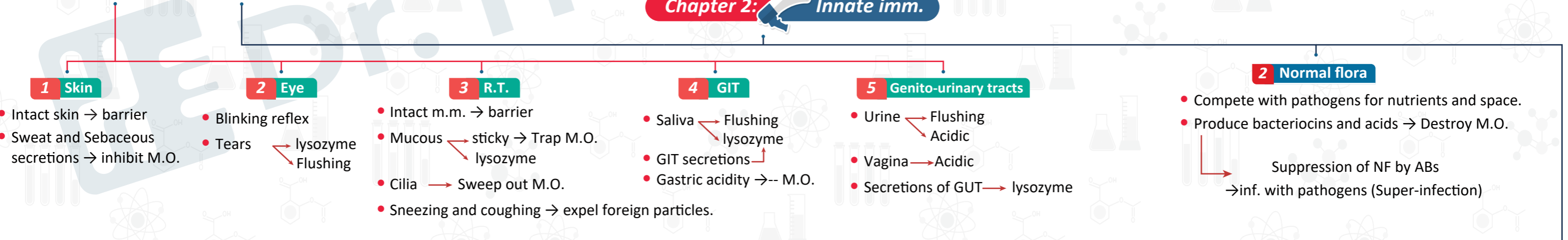


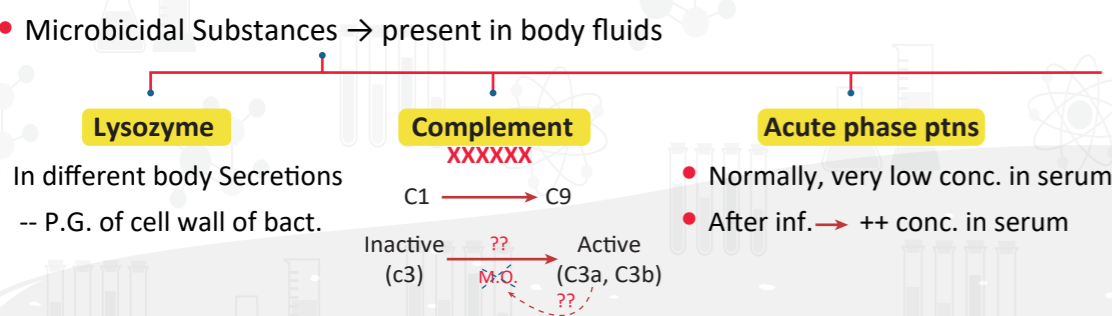
Innate imm. الجلد وما شابه	Acquired (Adaptive) imm. مرورة ← الحصبة ← مناعة ضد الحصبة
• Present since birth	• acquired during life time of the person
• 1st line of Defense → immediately available after infection	• 2nd line of Defense → Delay for few days until it acts
• Non-specific → in all individual # all M.Os	• Specific → in specific individual # specific M.O. <small>مرورة الحصبة</small>
• Less efficient in eliminating M.Os	• Very effective
• Doesn't require previous exposure to M.O.	• Require previous exposure to M.O.
• Doesn't increase with repeated exposure to the same M.O. (NO memory cells)	• Increase with repeated exposure to M.O. (Memory cells) → life long protection # the same pathogen
• Depends on:- 1) Granulocytes → Neutrophils, Eosinophils, basophils 2) Monocytes/Macrophages 3) N.K cells	• Depends on:- 1) B-lymphocytes. 2) T-lymphocytes

1 Mechanical Barriers and Surface Secretions

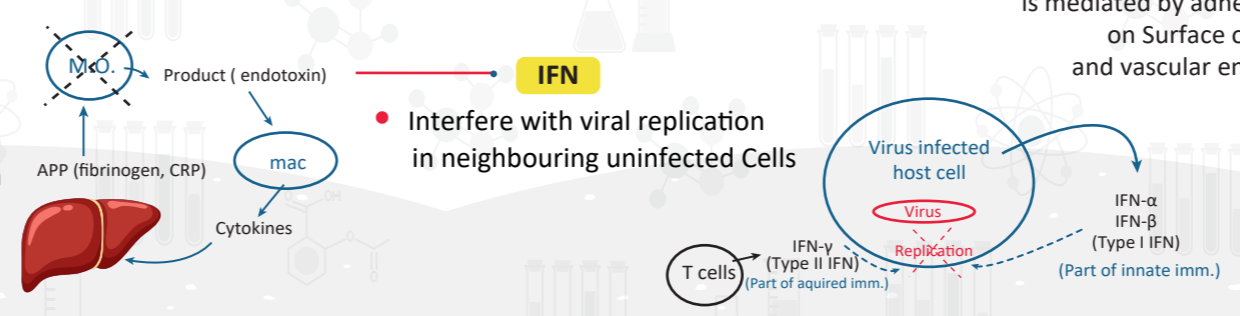


Chapter 2: Innate imm.

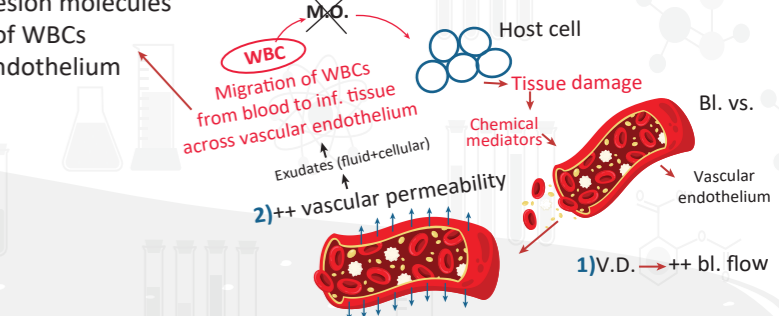
2 Soluble Defence factors

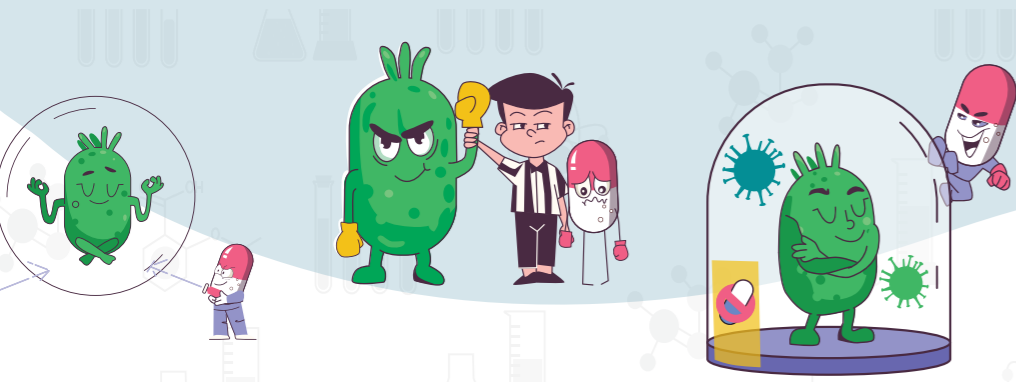


3 Cellular Defence factors



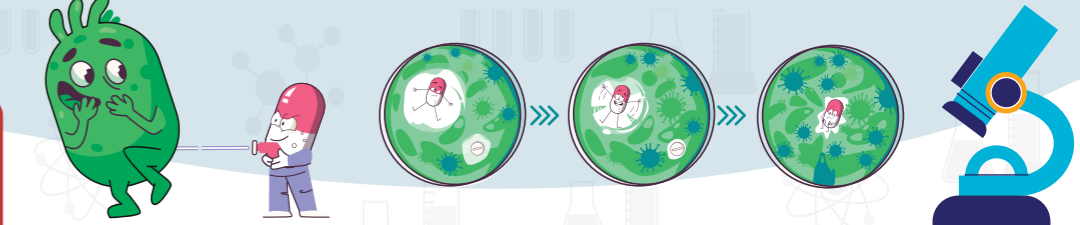
4 Inflammation





There is cooperation between innate and acquired imm.

- 1- Innate imm. helps acquired imm. → as in Ag presentation
- 2- Acquired imm helps innate imm. → as in opsonization



4 Cellular Defence factors

1-Phagocytes

- Neutrophils=microphages = PMNLs=polymorphs
- Monocytes (in blood) → macrophages (in tissues)
- Dendritic cells

Phagocytic cells

Steps of phagocytosis

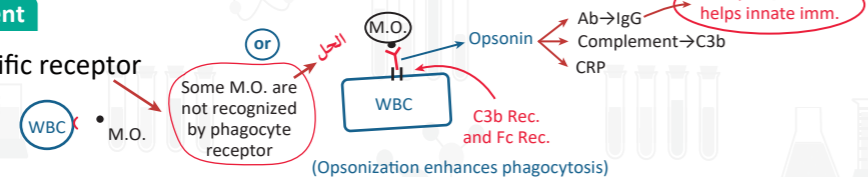
1 Migration=chemotaxis:

- Movement of WBCs towards M.O. in response to chemotactic factors
- from M.O. or from damaged tissue

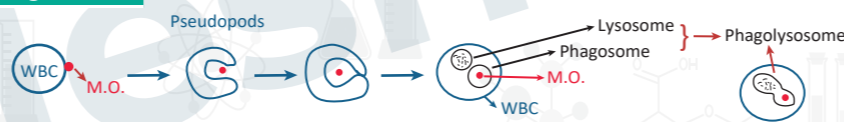
E.g. 1) Some cytokines
2) Complement

2 Attachment

- By non-specific receptor



3 Engulfment

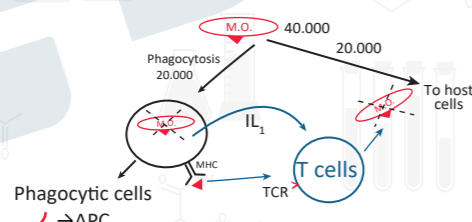


4 Killing

- O₂ dependent Killing: → by H₂O₂, O₂⁻, OH⁻ → (FR) → lethal to bact.
- O₂ independent Killing: → by lysosomal enz. (lysozyme, elastase,.....)

5 Ag presentation

Innate imm. helps acquired imm.



6 Other functions of monocytes and macrophages

- 1- Ag presentation
- 2- Secretion of cytokines (Interleukins) e.g. IL1
- 3- Direct cytotoxicity → Direct killing without engulfment

Helminthic parasites

Tumor cells

(Too large to be engulfed) → killed by releasing macrophage toxic contents onto them

2-Eosinophils

- Granulocytes present mainly in Tissues
- In blood → 1:3% of TLC

- 1- Defence # helminthic Parasites (too large to be phagocytosed → Direct cytotoxicity)
- 2- Role in allergy (histaminase enz.)
- 3- Phagocytic

3-Basophils & mast cells

- Granulocytes:
- 1- Basophils in blood 0-2% of TLC
- 2- Mast cells → Present either around bl. vs. or in submucosa of bl. vs.

Functions

- Their granules Contain imp. mediators (as histamine)
- 1- ++ inflammation 2- Role in allergy

4-NK cells

- Large granular lymphocytes (non-B, non-T cell)
- 10-15% of blood Lymphocytes

- 1- Non Specific killing of tumor cells. and virus infected Cells (as T_c Cells → They differ in the way they recognize their targets)

Immunity (2)

	Neutrophils	Monocytes / Macrophages
Content	• 60 : 80 % TLC	• 1 : 5 % TLC
Presence	• Absent from normal tissues → with inf., they migrate to site of inf. in response to chemotactic factors	• Continuously leave blood to tissues → mature into mac. • e.g. 1- Kupffer cells of liver 2- Alveolar mac. in lung
Size	• Smaller	• Larger
Life span	• Shorter → die after phagocytosis → form pus cells	• longer → survive after phagocytosis
Response during inflammation	• Rapid ++ in production • Rapidly form pus	• Slight ++ in bl. vs. • Slowly form granuloma
Importance	• Most numerous and most imp. cells of innate imm. → phagocytosis	• Phagocytosis • Ag presentation • Cytokines secretion • Direct cytotoxicity