

Immune Defence Mechanisms

Specific Defence

Inflammation

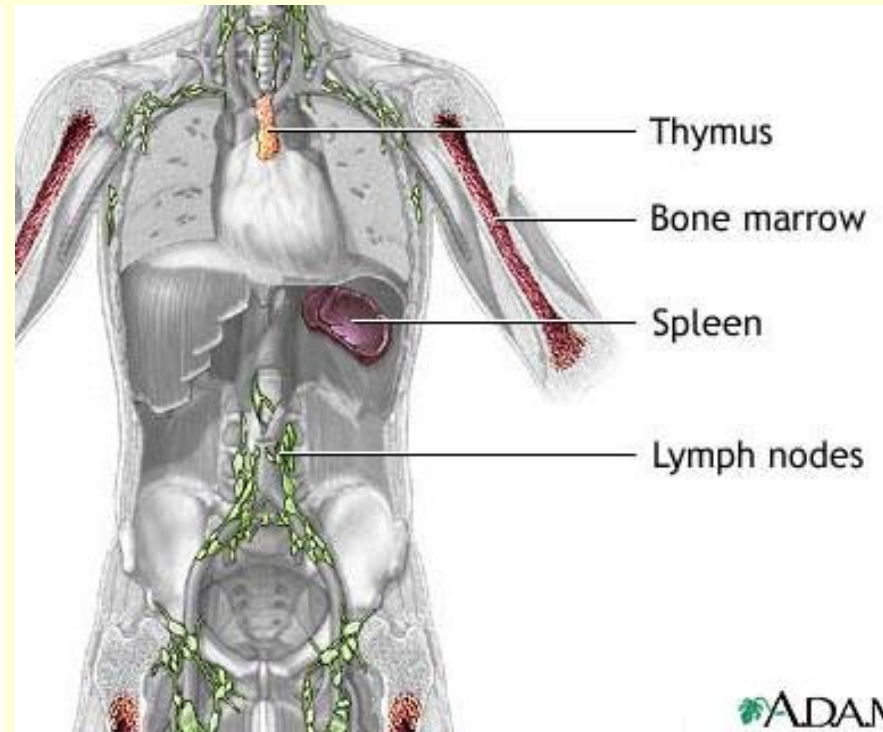
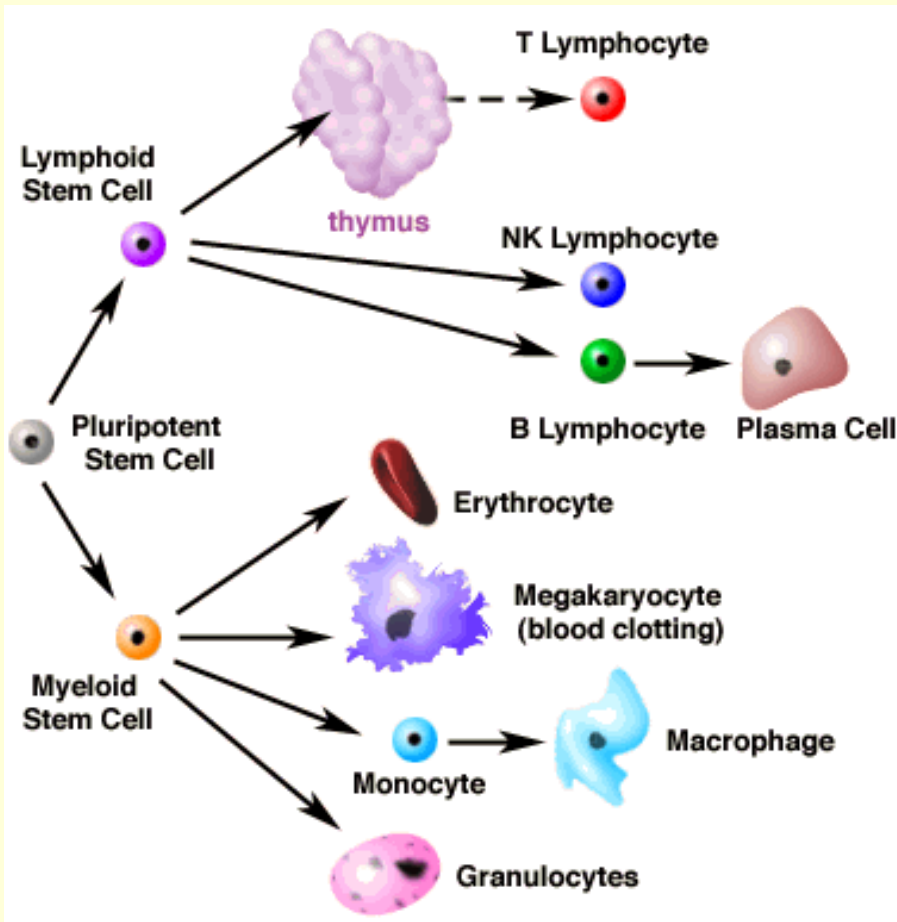
lecture from Physiology and Pathophysiology I

12. 10. 2021

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Immune System

- system of defence against pathogens (bacteria, viruses, fungi, parasites) and tumor cells



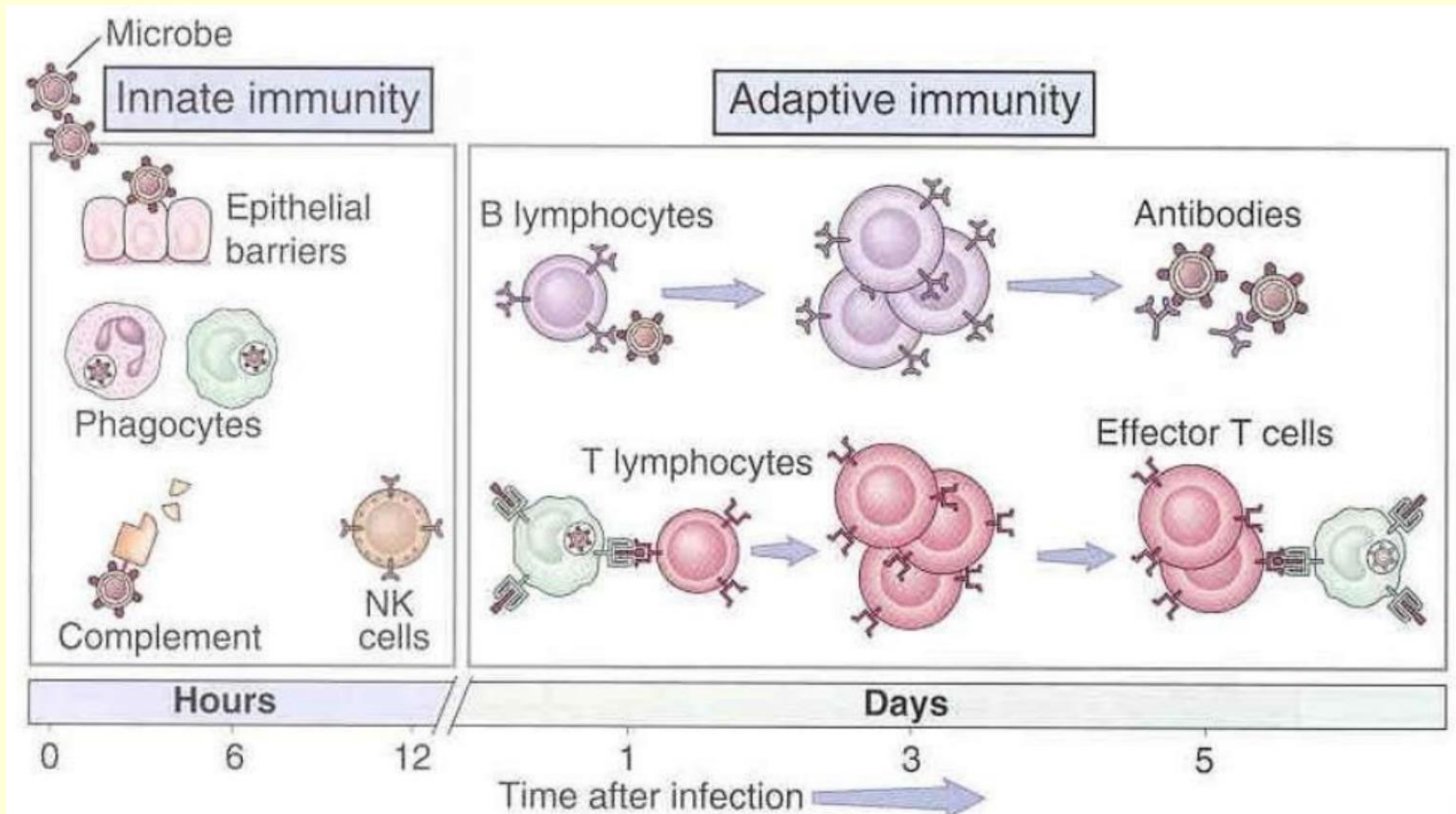
Nonspecific vs. Specific Defence

	Nonspecific/congenital	Specific/acquired
Cellular	polymorphonuclears monocytes–macrophages NK-cells	T-lymphocytes
Humoral	complement acute-phase proteins (CRP, ceruloplasmin...)	antibodies B-lymphocytes

Congenital vs. Acquired Immunity

Attribute	Congenital/Nonspecific Immunity	Acquired/Specific Immunity
Response time	Minutes/hours	Days
Specificity	Specific for molecules and molecular patterns associated with pathogens	Highly specific; discriminates even minor differences in molecular structure; details of microbial or nonmicrobial structure recognized with high specificity
Diversity	A limited number of germ line–encoded receptors	Highly diverse; a very large number of receptors arising from genetic recombination of receptor genes
Memory responses	None	Persistent memory, with faster response of greater magnitude on subsequent infection
Self/nonsel self discrimination	Perfect; no microbe-specific patterns in host	Very good; occasional failures of self/nonsel self discrimination result in autoimmune disease
Soluble components of blood or tissue fluids	Many antimicrobial peptides and proteins	Antibodies
Major cell types	Phagocytes (monocytes, macrophages, neutrophils), natural killer (NK) cells, dendritic cells	T cells, B cells, antigen-presenting cells

Congenital vs. Acquired Immunity



Surface Coverage

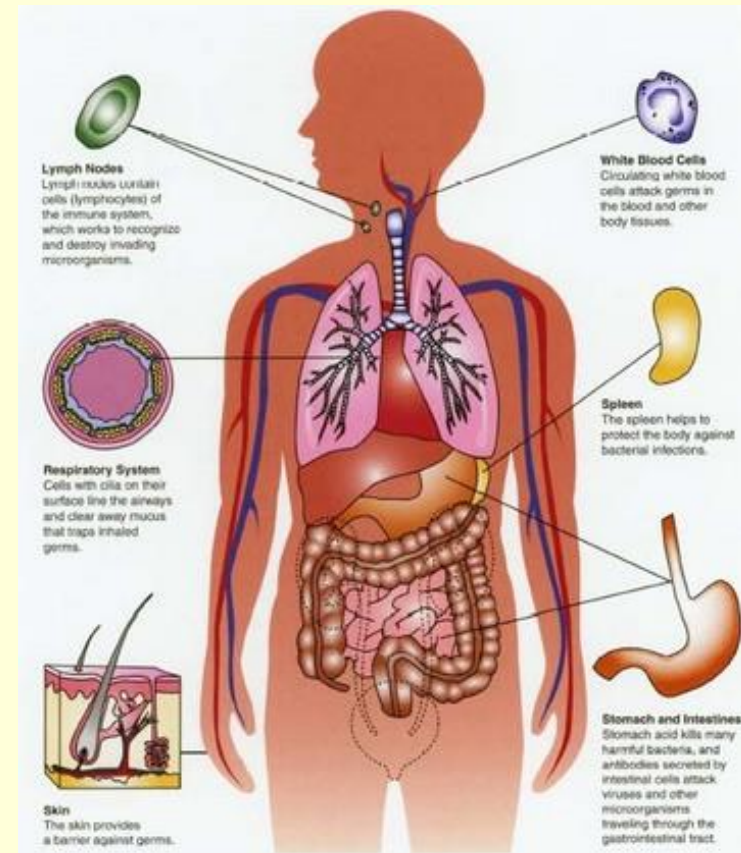
First Line of Defense

PHYSICAL FACTORS

- skin
- mucous membranes with lysozymes
- lacrimal apparatus
- salivary glands
- vaginal secretions
- flow of urine

CHEMICAL FACTORS

- sebum
- lysozyme
- gastric juice
- vaginal secretions
- urine



The human body has several lines of defense against infection, which work to prevent germs from invading the body or to destroy them once they find their way in.

Nonspecific Immune System

Second Line of Defense

- generalized responses to pathogen infection – do not target a specific cell type

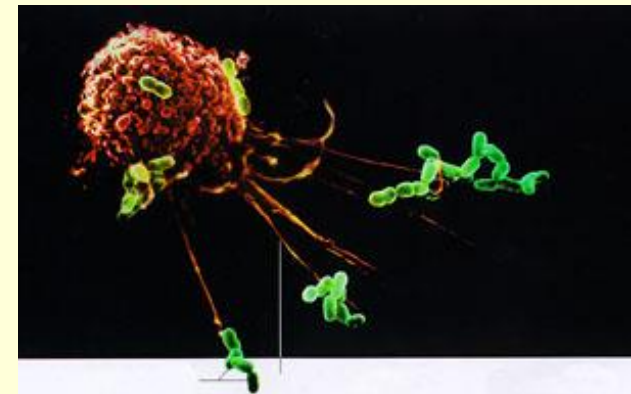
Phagocytes

- neutrophils
- eosinophils
- macrophages

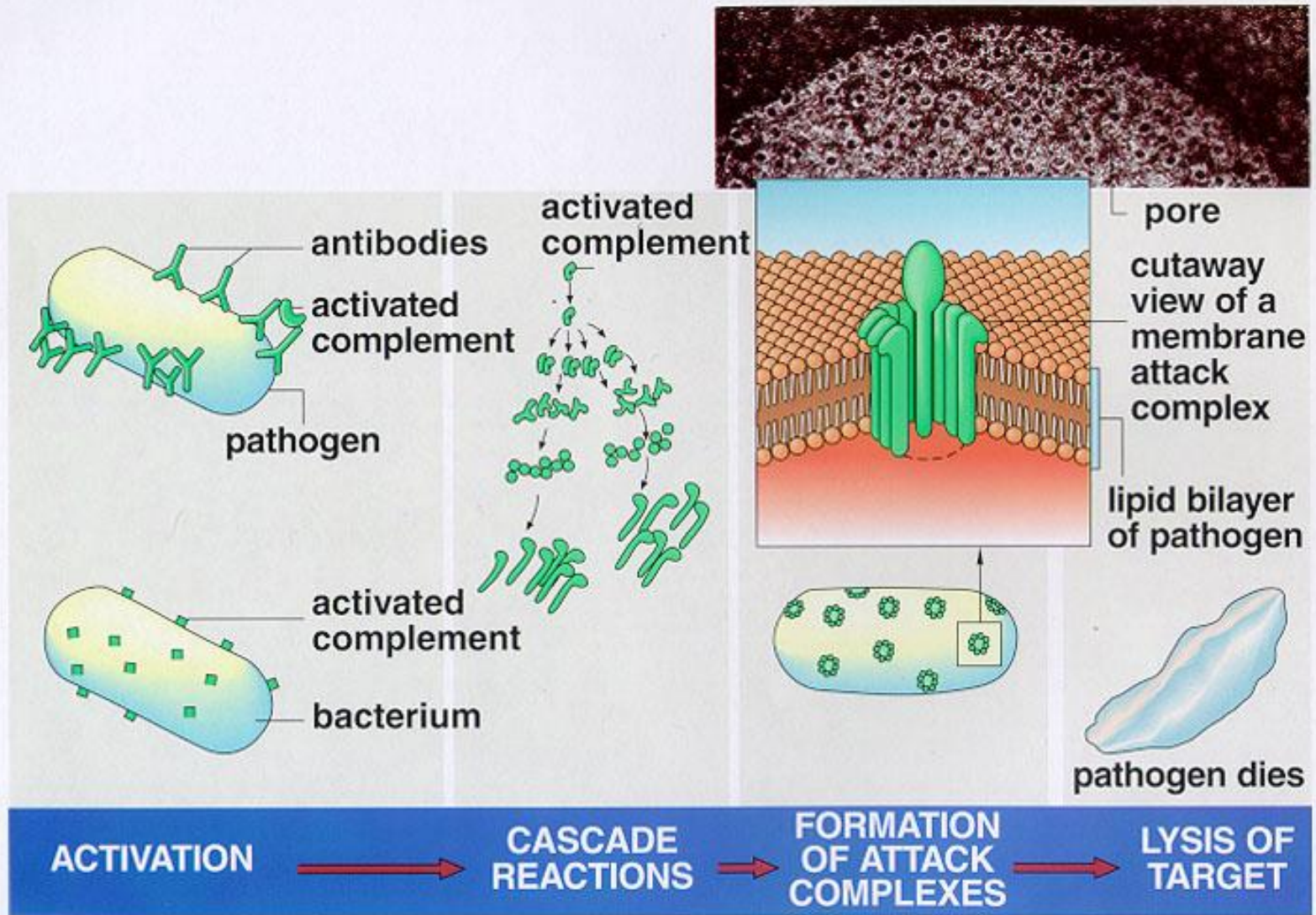
Non-phagocytic leucocytes

- basophils
- mast cells

Complement proteins



Complement



Specific Immune System

Third Line of Defense

HUMORAL (Ab-MEDIATED) IMMUNE SYSTEM

- antibodies
- B-cells (lymphocytes)

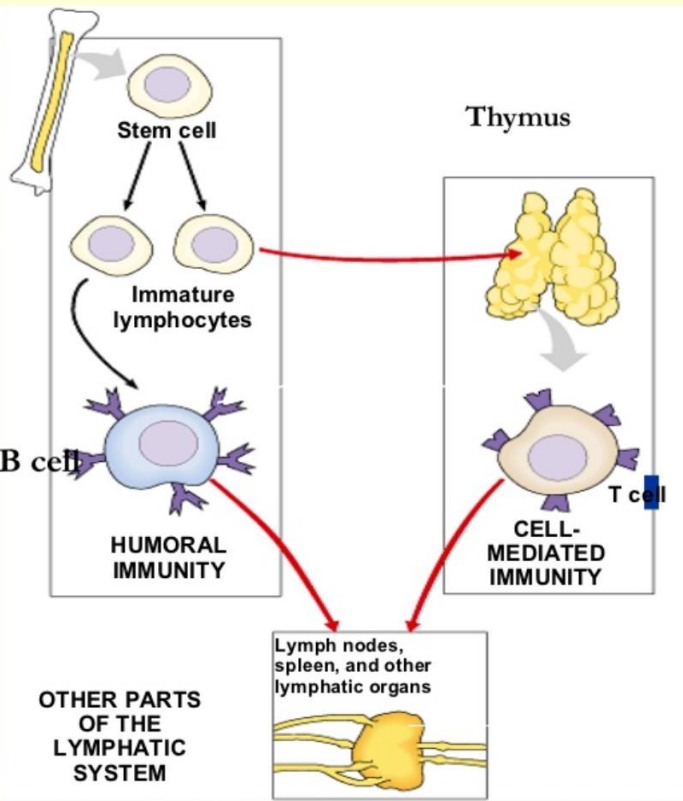
CELL-MEDIATED SYSTEM





- T-cells (lymphocytes)
- helper T lymphocytes (T_H)
 - produce and secrete chemicals that promote large numbers of effector and memory cells
- cytotoxic T lymphocytes (TC)
 - eliminate infected body cells and tumor cells

PHAGOCYTOTIC COMPONENTS

- macrophages, monocytes, neutrophils
- engulf foreign objects and inform T lymphocytes that a specific antigen is present

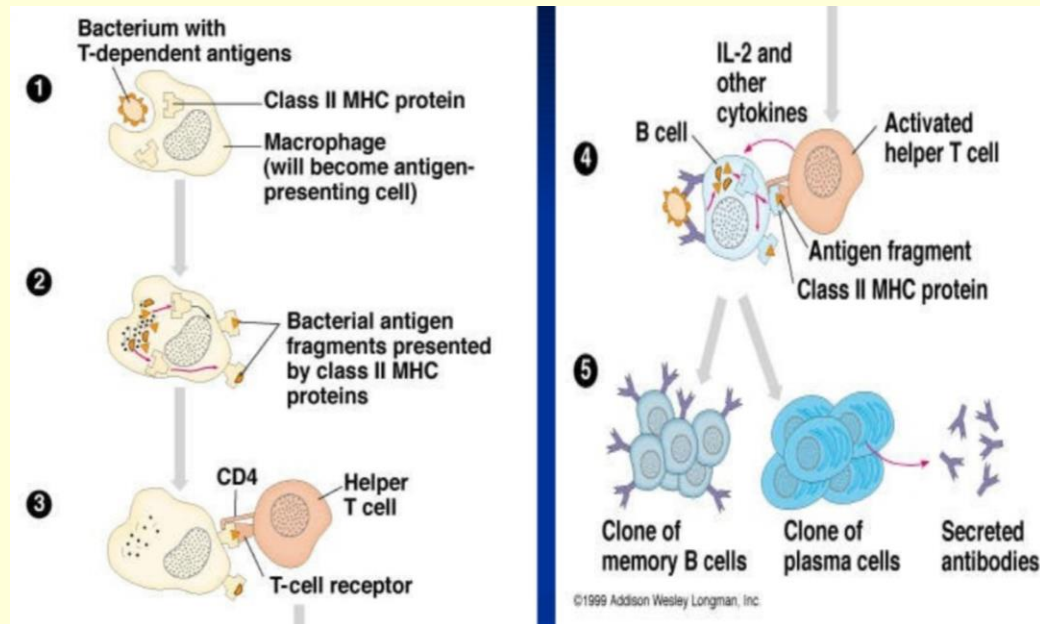
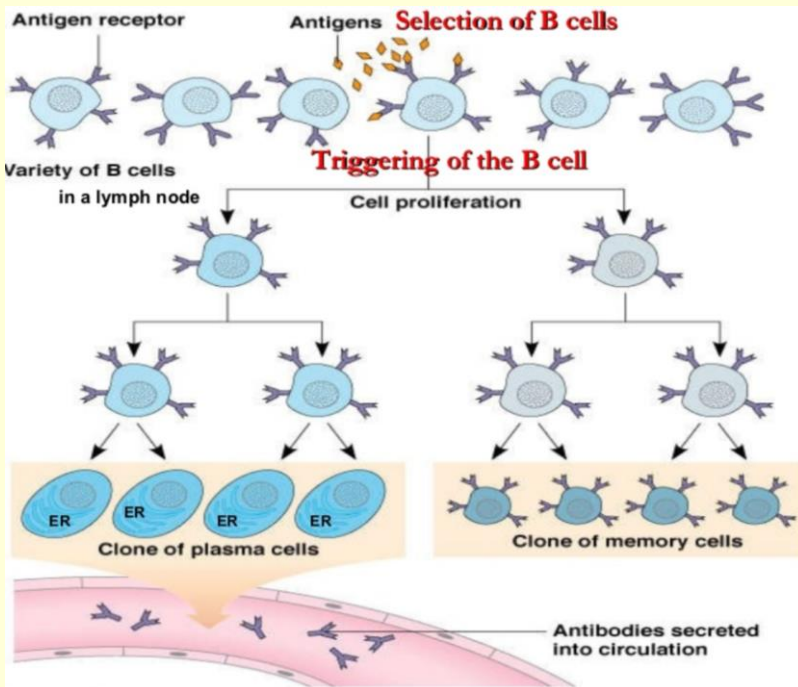
Lymphocytes



LYMPHOCYTES		
	B cells	Differentiate to form antibody-producing cells and memory cells
	Plasma cells	Secrete antibodies
	T cells	Kill virus-infected cells; regulate activities of other white blood cells
	Natural killer cells	Attack and lyse virus-infected or cancerous body cells

Humoral (Antibody-Mediated) Immunity



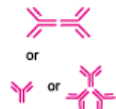


- production of **antibodies** by **B-cells**
- B-cells migrate to the lymphoid organs (spleen, lymph nodes) after maturation



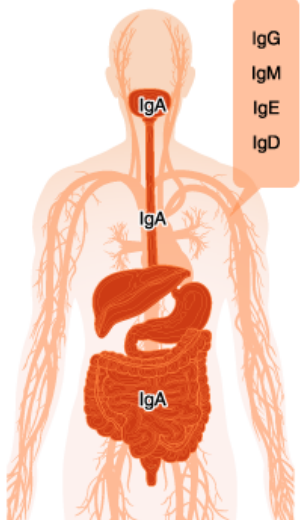
Antibodies

- synthesized by B-cells in soluble or cell-bound form
- each antibody recognizes one specific antigen
- **immunoglobulins** (5 classes) – glycoproteins

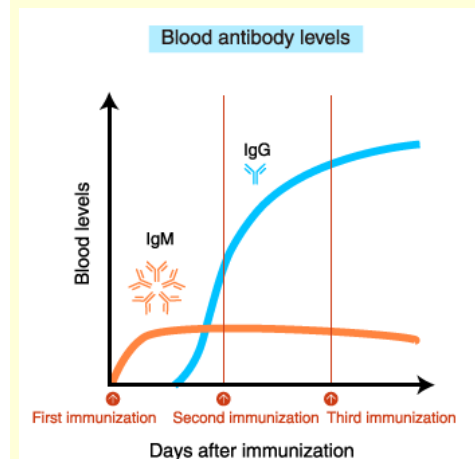
- **Ig G**
- **Ig M**
- **Ig A**
- **Ig D**
- **Ig E**

Types and characteristics of antibodies		
IgG		<ul style="list-style-type: none"> • Highest opsonization and neutralization activities. • Classified into four subclasses (IgG1, IgG2, IgG3, and IgG4).
IgM		<ul style="list-style-type: none"> • Produced first upon antigen invasion. Increases transiently.
IgA		<ul style="list-style-type: none"> • Expressed in mucosal tissues. Forms dimers after secretion.
IgD		<ul style="list-style-type: none"> • Unknown function.
IgE		<ul style="list-style-type: none"> • Involved in allergy.

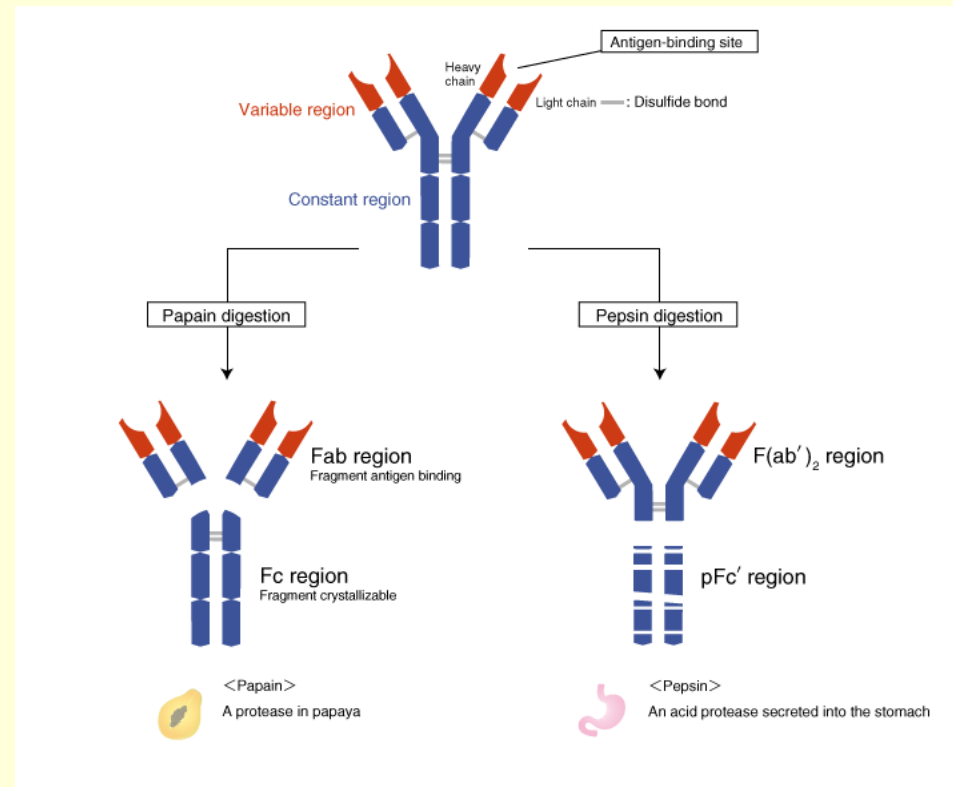
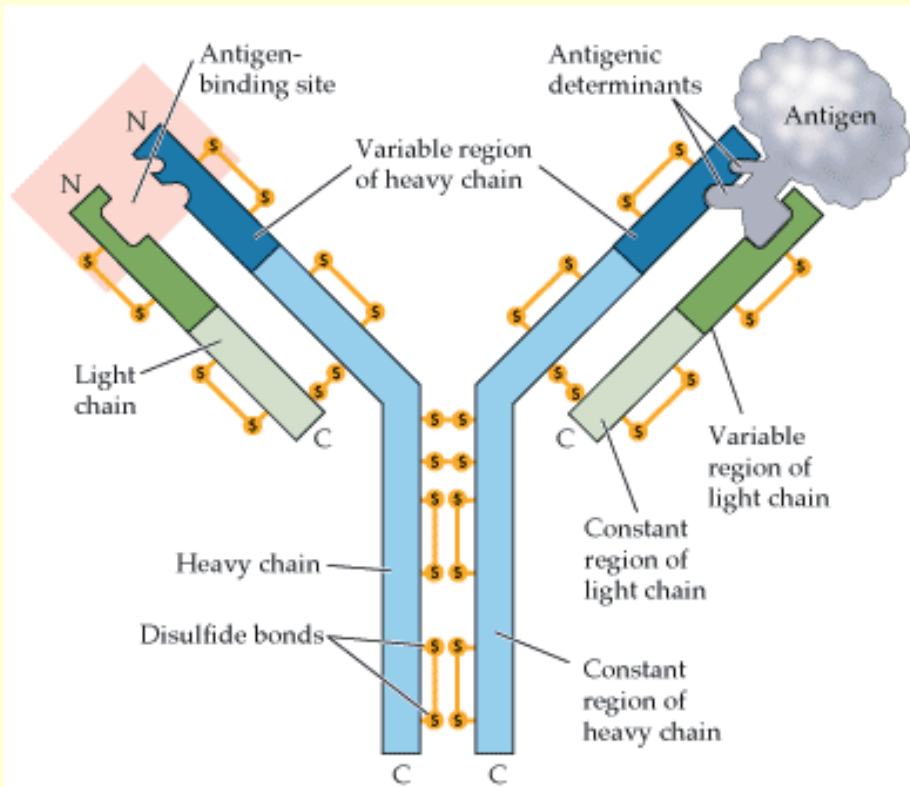
Distribution in the body



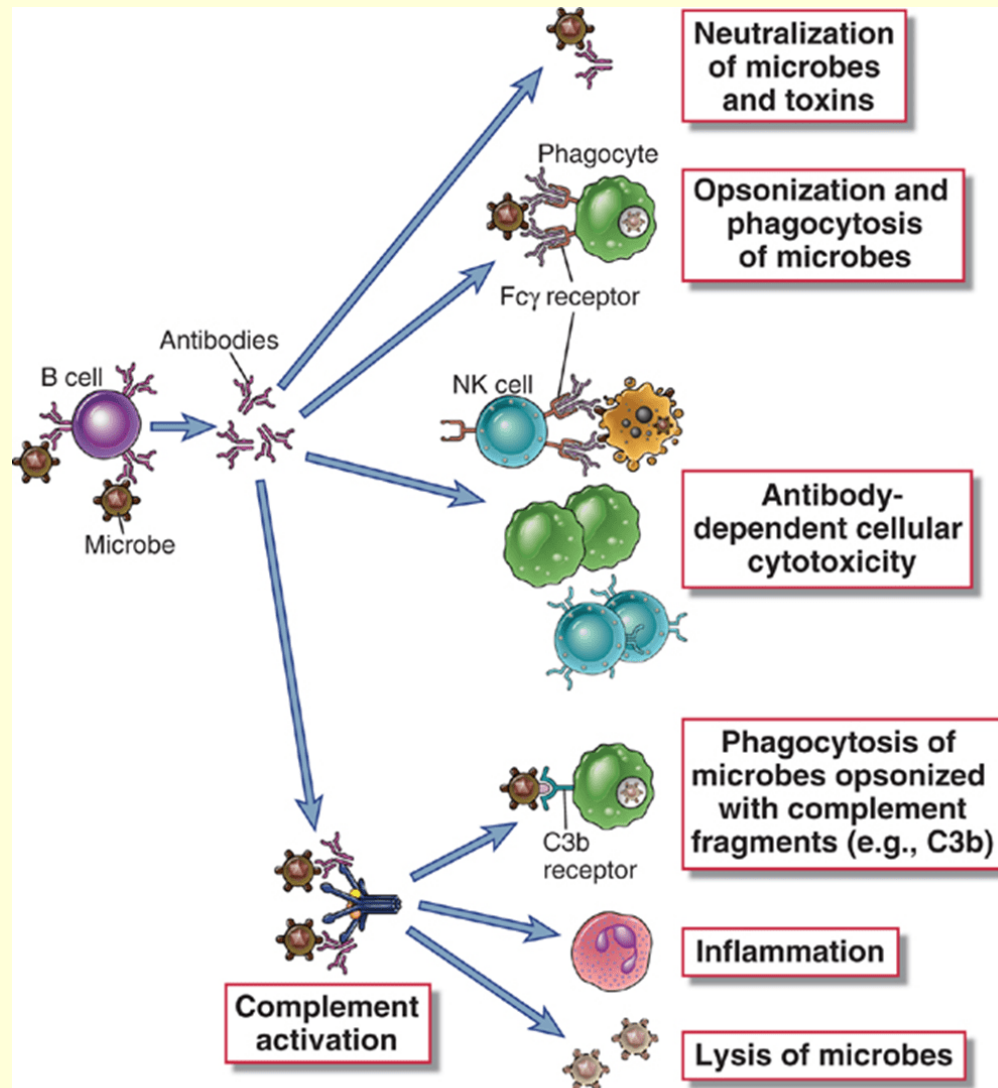
IgG
IgM
IgE
IgD



Antibodies Structure

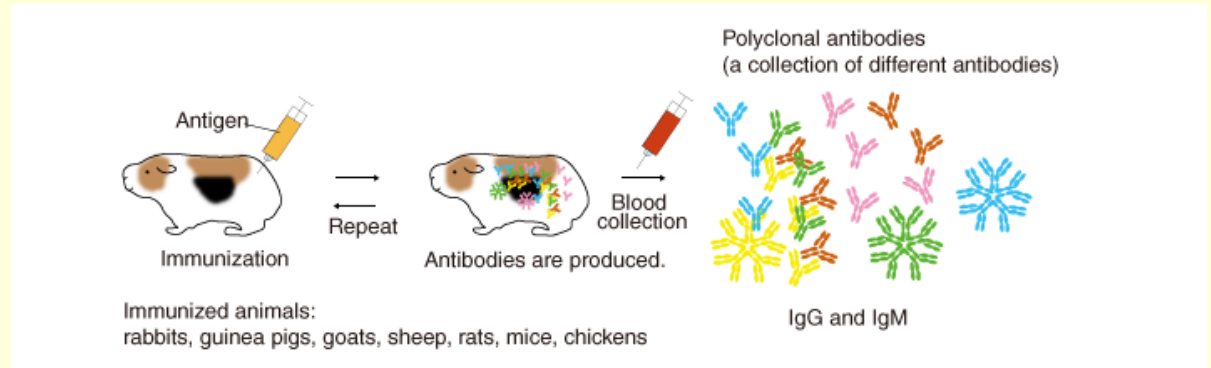


Antibodies Function

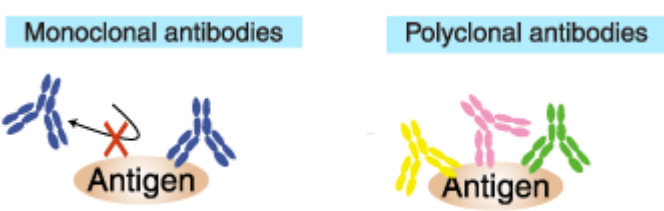
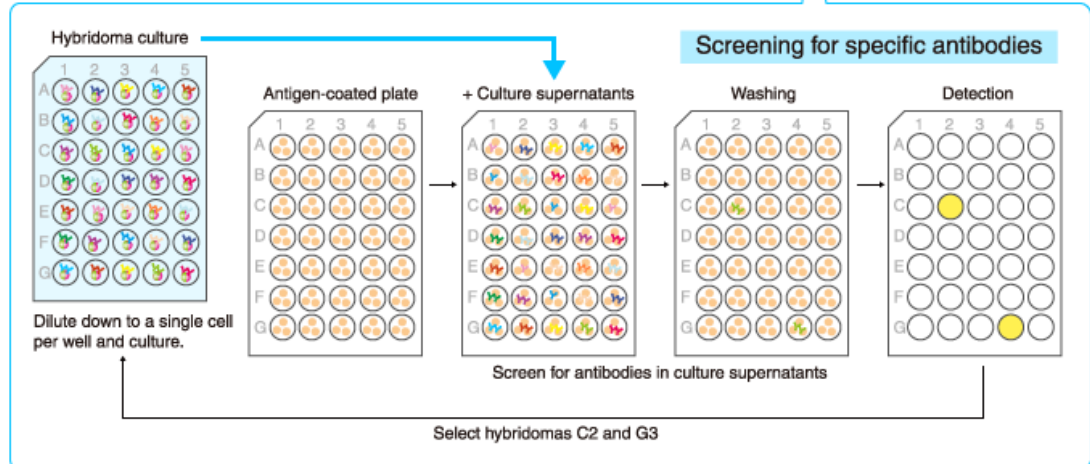
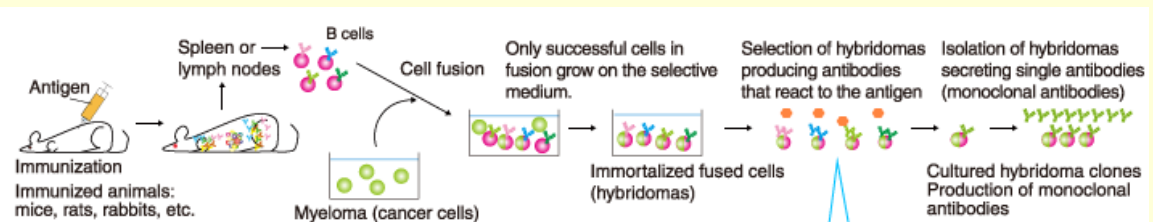


Antibodies Production

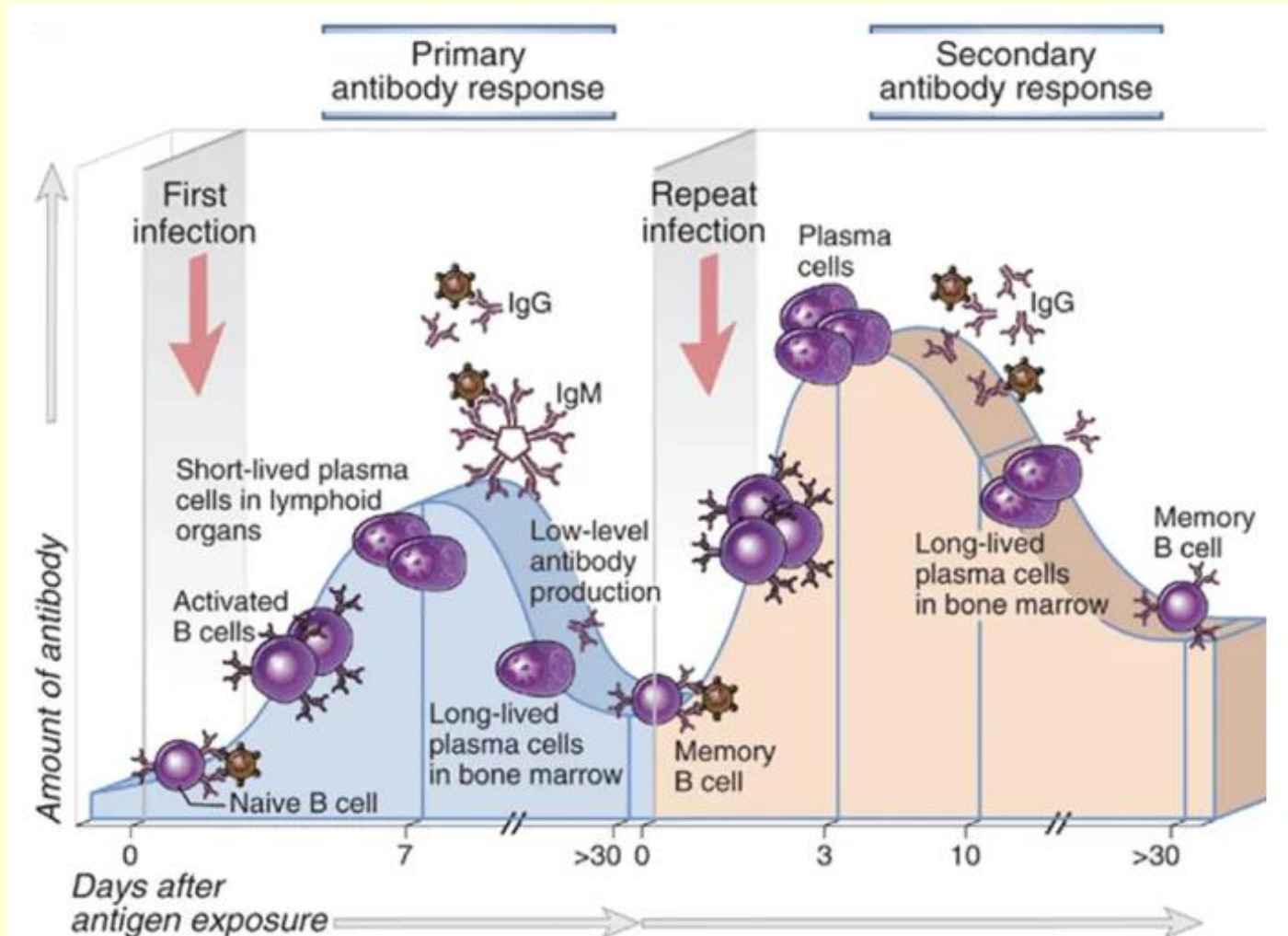
- polyclonal



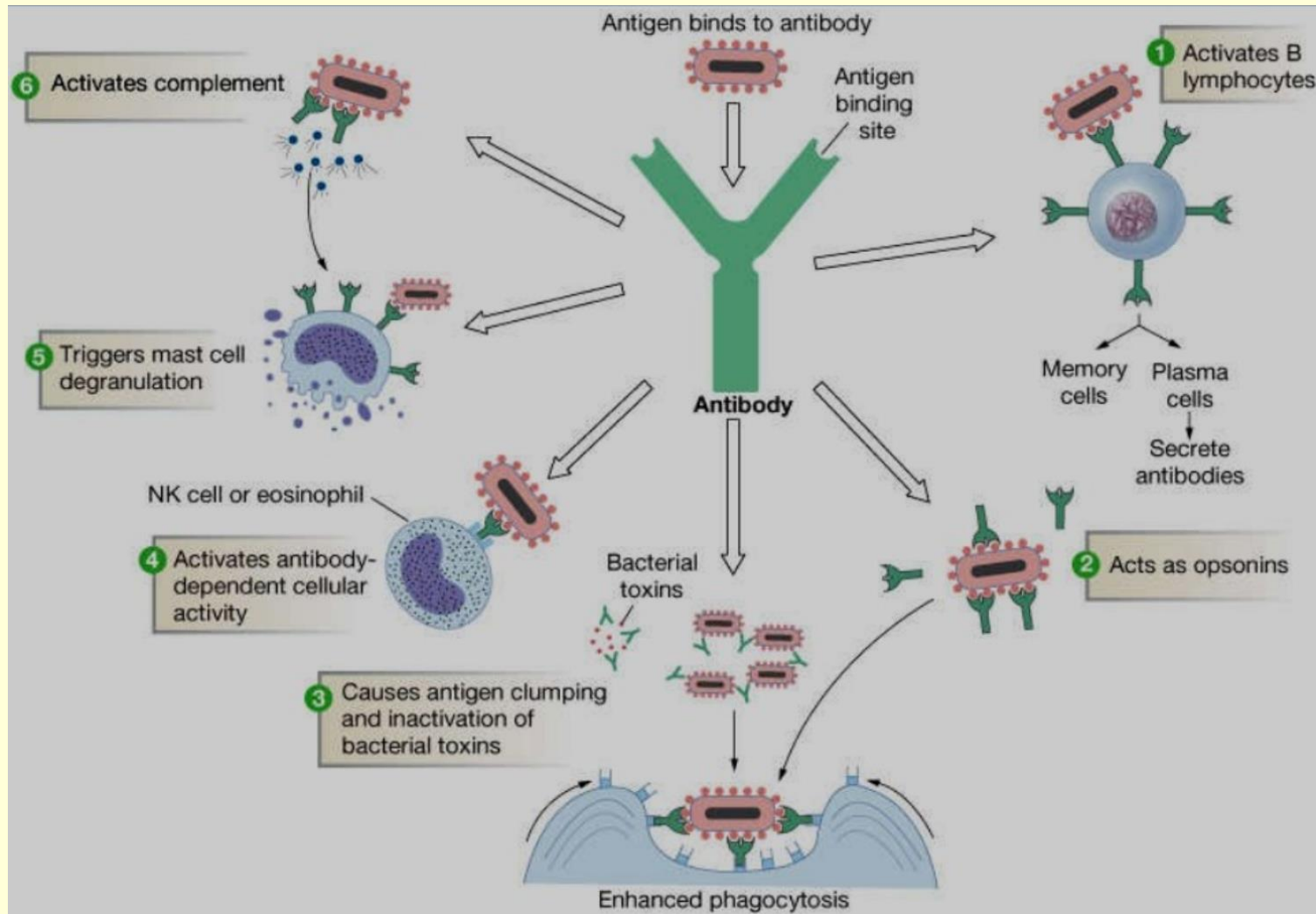
- monoclonal



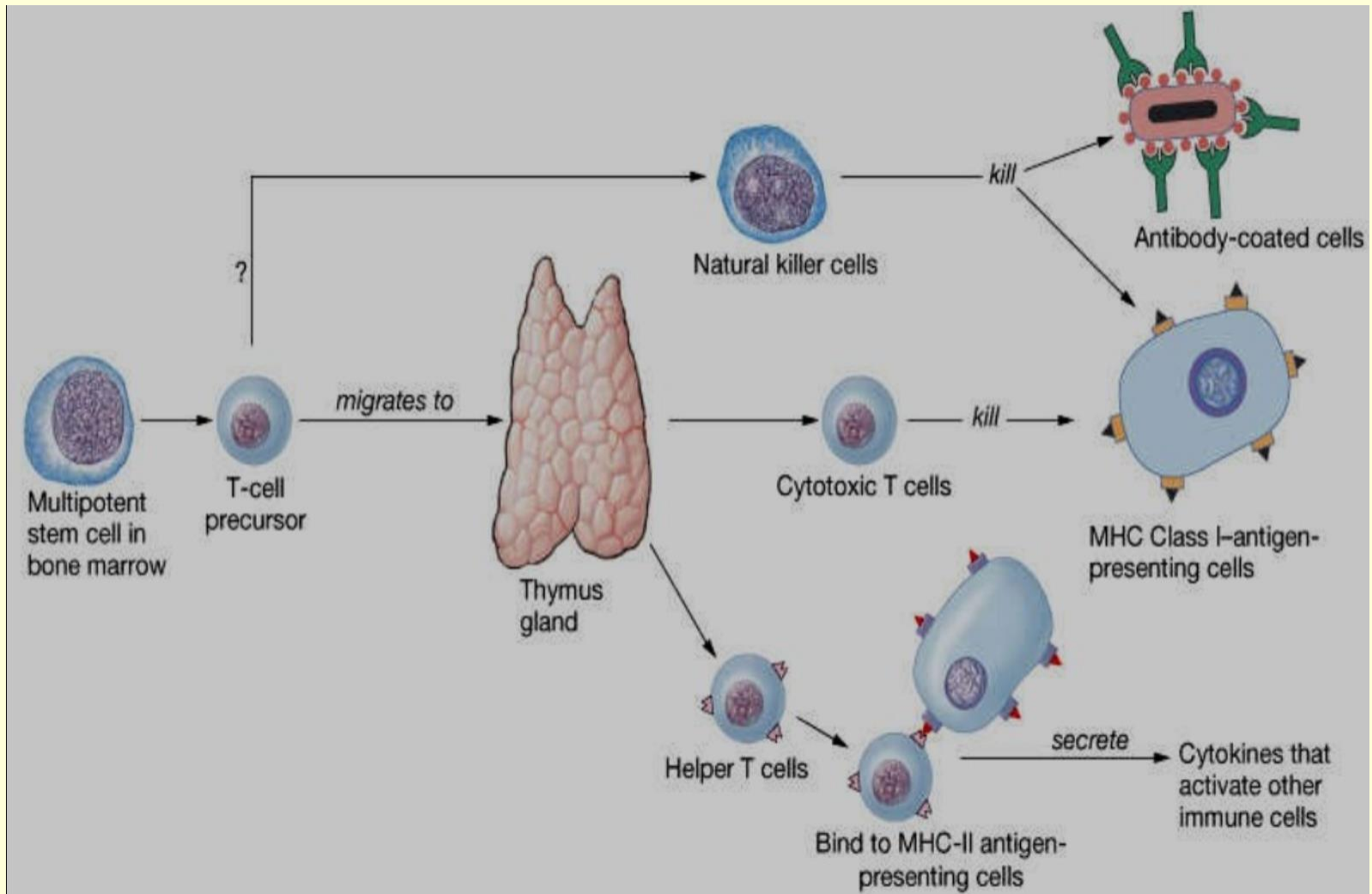
Primary and Secondary Antibody Response



Antigen-Antibody Response

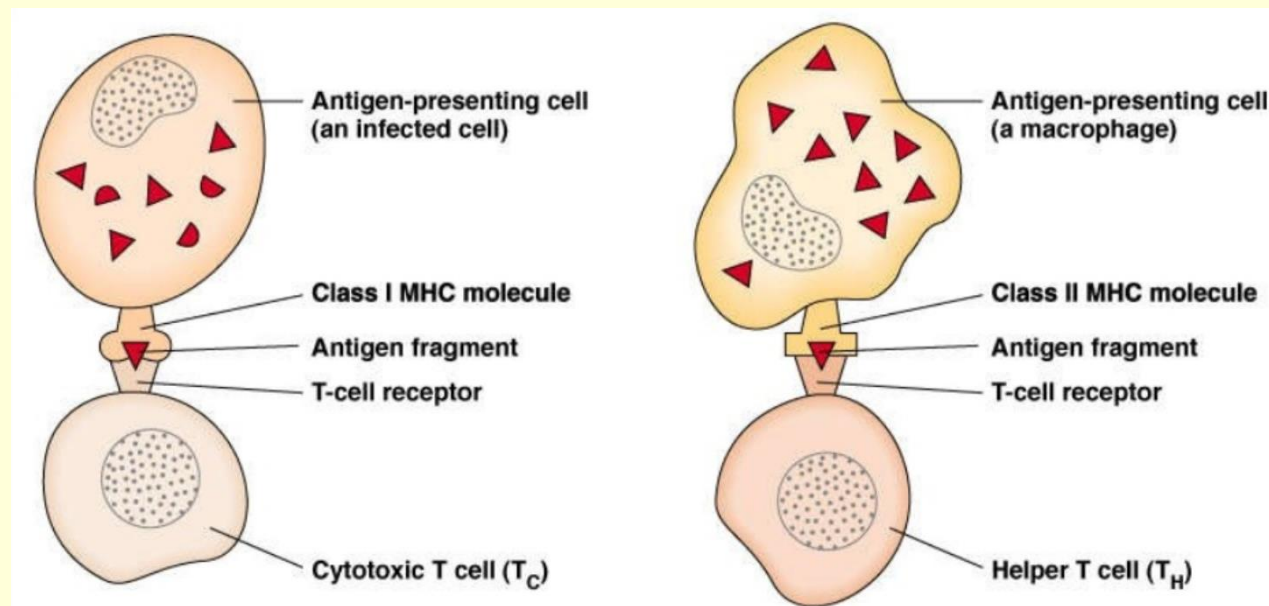


Cell-Mediated Immunity T-cells



Cell-Mediated Immunity T-cells

- T-cells recognize antigen associated with **MHC (major histocompatibility complex)** molecules in the cells
 - **MHC class I**
 - **MHC class II**



(a)

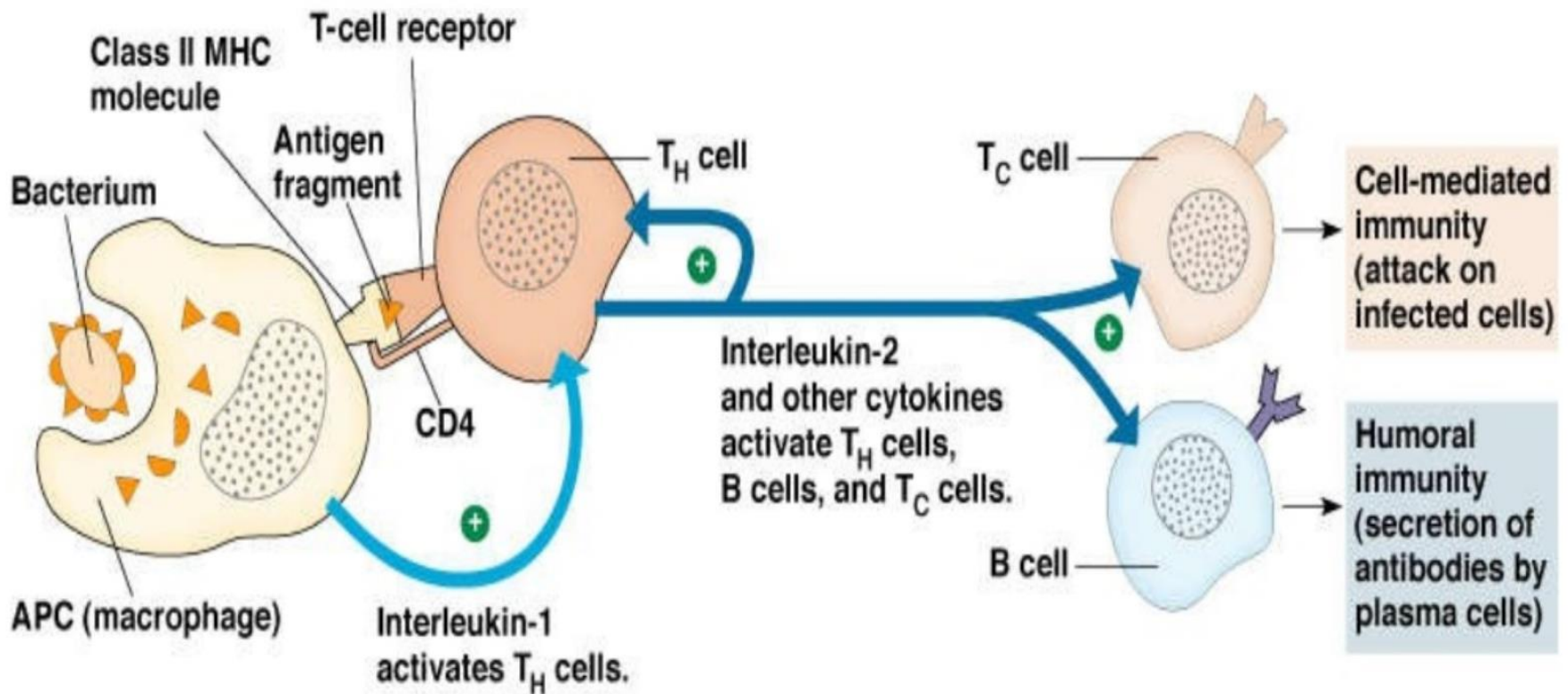
(b)

Cell-Mediated Immunity T-cells

T_H helper cells

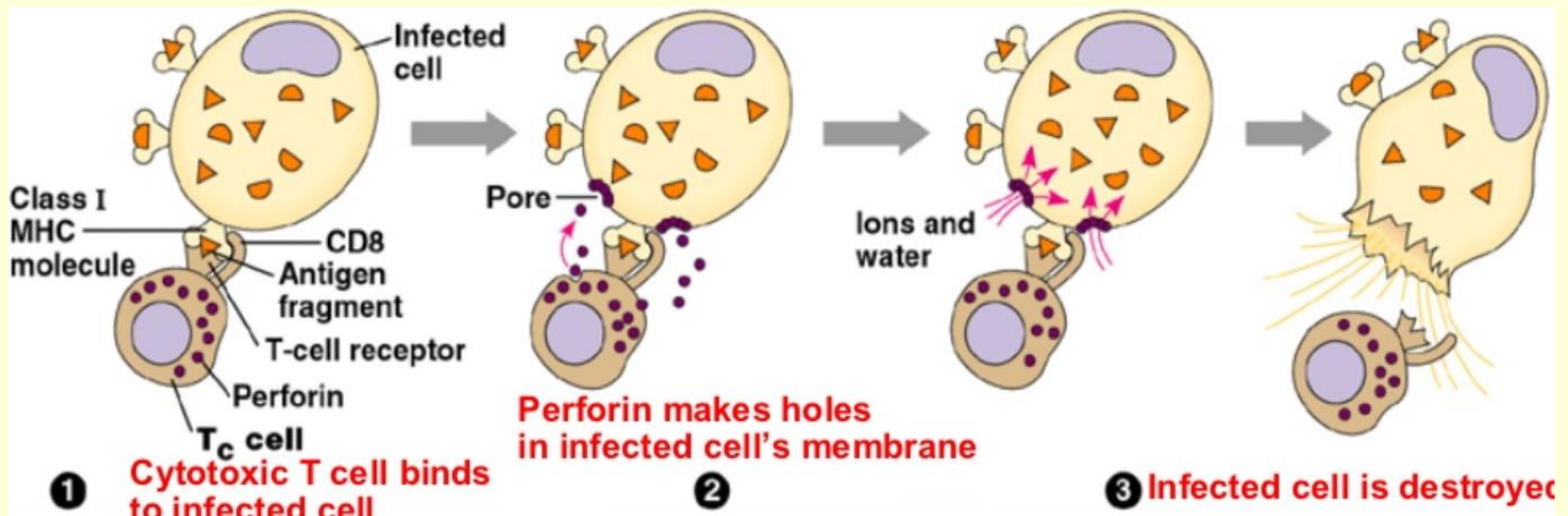
- T_H cells receptors recognize antigen on the surface of APC (macrophages)
- release of cytokines and activation of T_C and B-cells
- T_C cells directly attack and destroy infected cells \Rightarrow **CELL IMMUNE RESPONSE**
- B-cells (plasma cells) produce antibodies \Rightarrow **HUMORAL IMMUNE RESPONSE**

Helper T_H-cells



Cytotoxic T_C-cells

- recognize the antigens on the surface of the cells
- destroy infected cells (bacteria, viruses, fungi, parasites), tumour cells, transplanted tissue



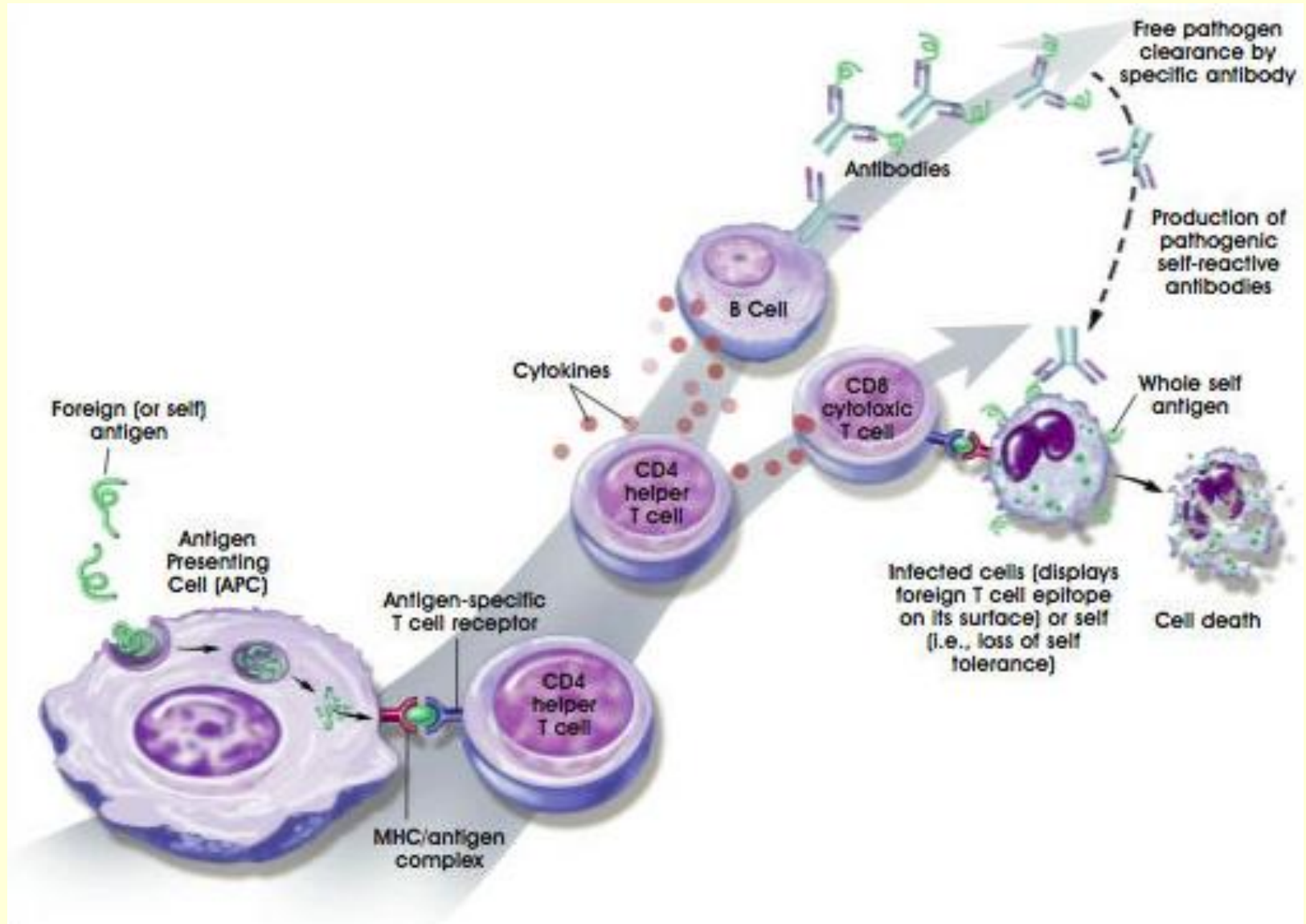
Cytokines

- small proteins involved in cell signaling
- amplify and regulate immune responses
- **interleukins** (IL-1, IL-2...)
- **TNF- α** (tumor necrosis factor alpha)
- **INF- γ** (interferon beta)
- **colony-stimulating factors** (G-CSF)

Complex Immune Reaction

- antigen ingested and presented by **antigen presenting cells (APC)**, like macrophages
- **helper T-cells** react with MHC-antigen complex
- **T-cell activation**, proliferation and cytokine production
- **cytokines** activate other cells (macrophages, NK cells, T_C cells)
- IL-2 stimulates **B-cells** to be developed to **plasma cells** with antibody production

Complex Immune Reaction



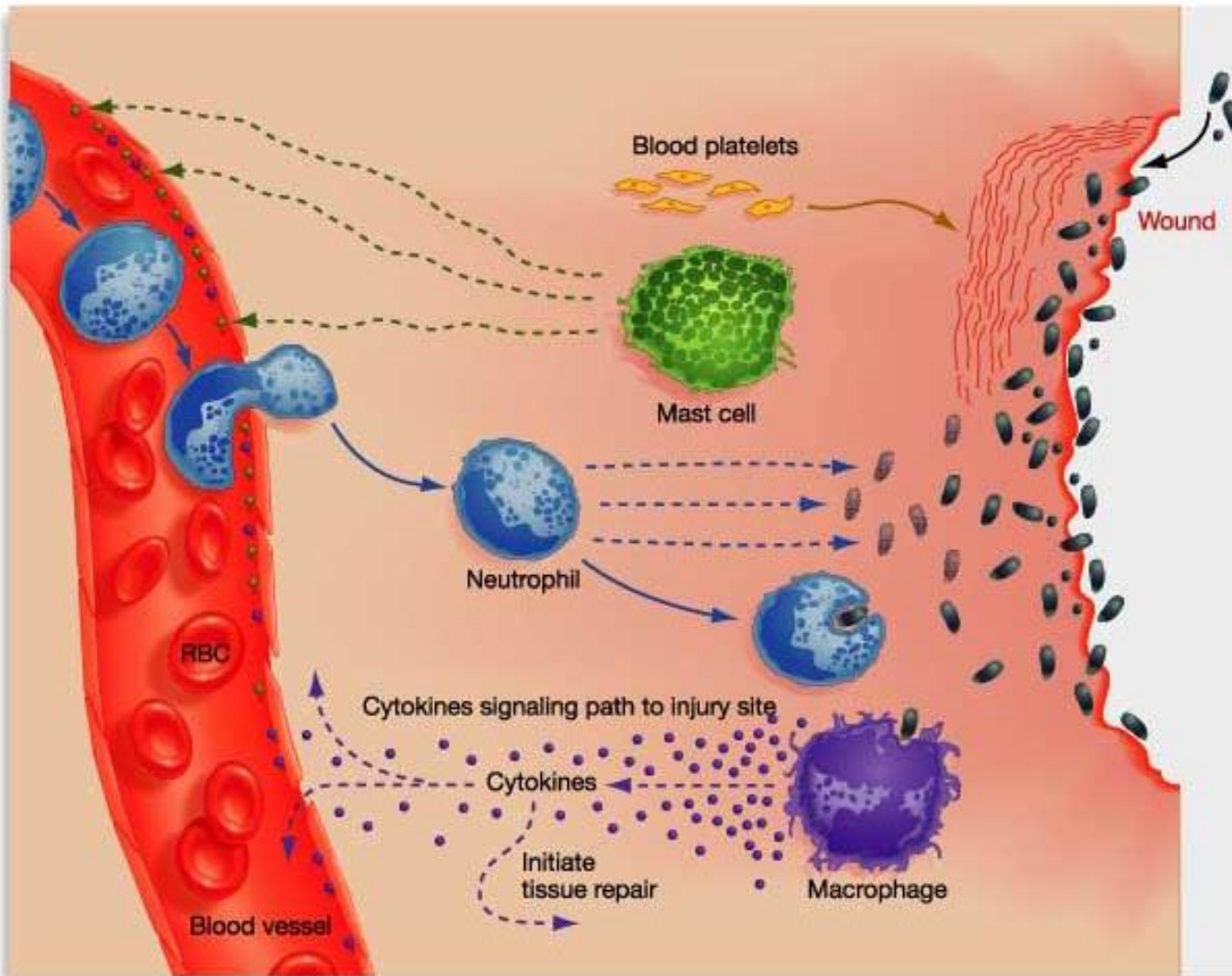
Inflammation

- defense reaction of the organism to injurious stimuli

MAIN AIMS

- to neutralize and destroy invading and harmful agents
- to limit the spread of harmful agents to other tissues
- to prepare any damaged tissue for repair

Inflammation



1. Bacteria and other pathogens enter wound.
2. Platelets from blood release blood-clotting proteins at wound site.
3. Mast cells secrete factors that mediate vasodilation and vascular constriction. Delivery of blood, plasma, and cells to injured area increases.
4. Neutrophils secrete factors that kill and degrade pathogens.
5. Neutrophils and macrophages remove pathogens by phagocytosis.
6. Macrophages secrete hormones called cytokines that attract immune system cells to the site and activate cells involved in tissue repair.
7. Inflammatory response continues until the foreign material is eliminated and the wound is repaired.

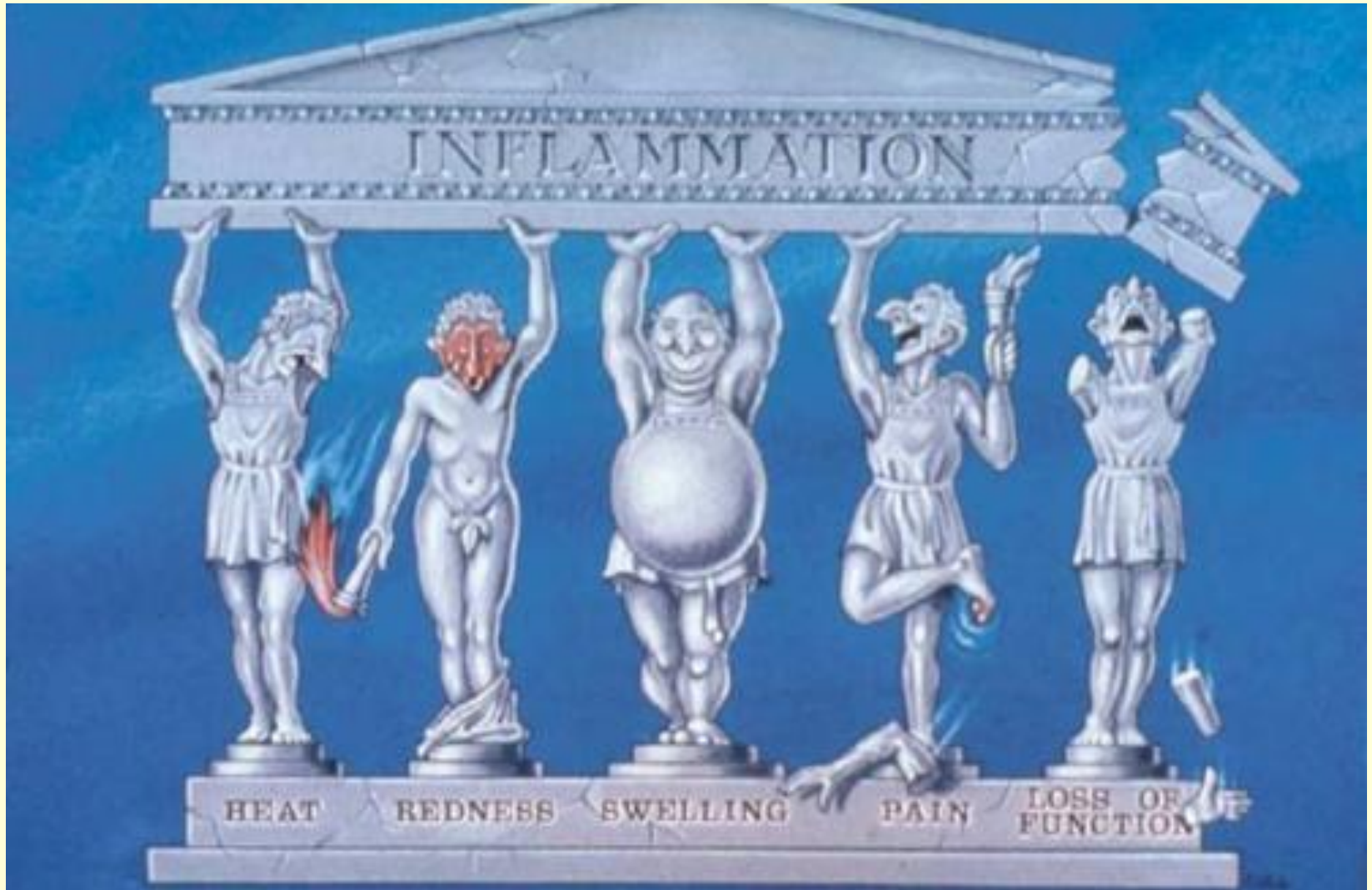
Local Signs of Inflammation

FIVE CARDINAL-LOCAL SIGNS OF INFLAMMATION:

1. RUBOR (REDNESS)
2. TUMOR (SWELLING)
3. CALOR (HEAT)
4. DOLOR (PAIN)
5. FUNCTIO LAESA (LOSS OF FUNCTION)

.... - ITIS

Inflammation



Inflammation

SYSTEMIC MANIFESTATION OF INFLAMMATION:

- fever
- Increased leukocyte counts (leukocytosis)
- lethargy
- muscle catabolism
- increased acute phase proteins (CRP C-reactive protein)
- increased erythrocyte sedimentation rate (ESR/FW)

Inflammation

INFLAMMATION AND INFECTION ARE COMMONLY CONFUSED
BECAUSE THEY OFTEN COEXIST

INFECTION IS ALWAYS ACCOMPANIED
BY INFLAMMATION;

HOWEVER,

NOT ALL INFLAMMATION
INVOLVES AN INFECTIOUS AGENT

Inflammation



Causes of Inflammation

Biological

- bacteria, viruses, fungi, parasites

Physical

- UV, temperature, X-rays

Chemical

- strong acids and alkalies

Endogenous

- autoimmune diseases, disintegrating tumor cells

Surgery, trauma