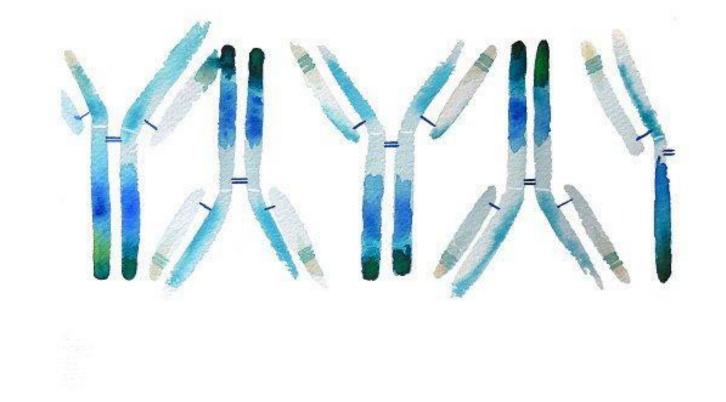
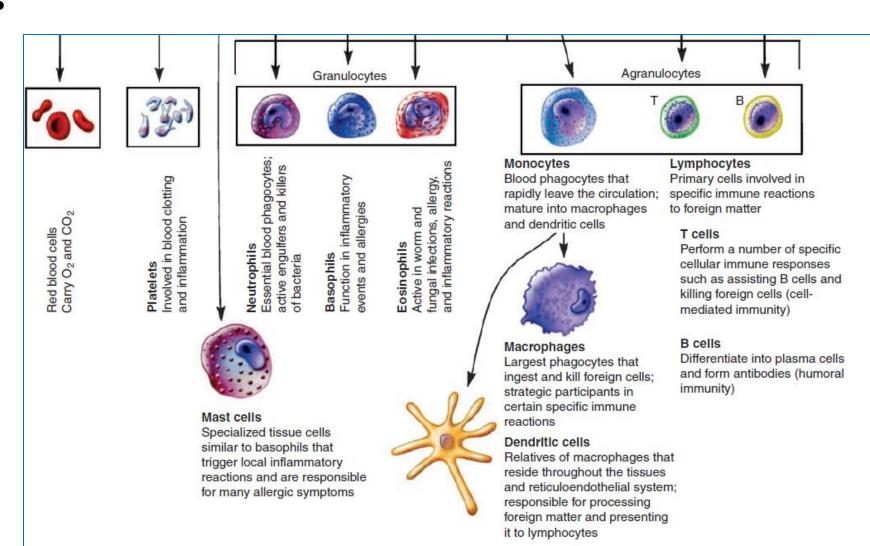
# Medical Immunology



Anas Abu-Humaidan M.D. Ph.D.

#### Cells of the immune system

- Phagocytes
- Mast Cells, Basophils, Eosinophils
- Antigen-Presenting Cells
- Lymphocytes

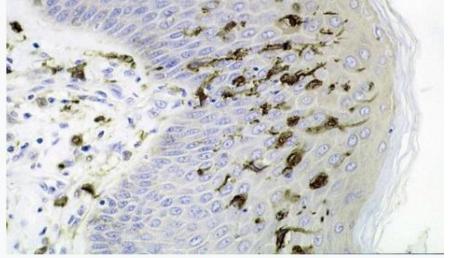


Cells of the immune system / Antigen presenting cells

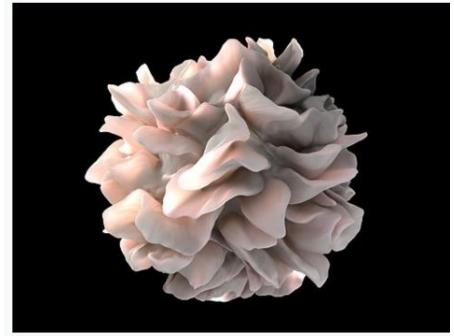
- Antigen-presenting cells (APCs) are cell populations that are specialized to capture
  microbial and other antigens, display them to lymphocytes, and provide signals that
  stimulate the proliferation and differentiation of the lymphocytes.
- The major type of APC that is involved in initiating T cell responses is the **dendritic cell**.
- Macrophages and B cells present antigens to T lymphocytes in different types of immune responses.
- APCs link responses of the innate immune system to responses of the adaptive immune system, and therefore they may be considered components of both systems.

# Cells of the immune system / Antigen presenting cells/ Dendritic Cells

- Dendritic cells are the most important APCs for activating naive T cells, and they play major roles in innate responses to infections and in linking innate and adaptive immune responses.
- They have long membranous projections and phagocytic capabilities and are widely distributed in lymphoid tissues, mucosal epithelium, and organ parenchyma.
- In response to activation by microbes, conventional dendritic cells in skin, mucosa, and organ parenchyma become mobile, migrate to lymph nodes, and display microbial antigens to T lymphocytes.



Dendritic cells in skin



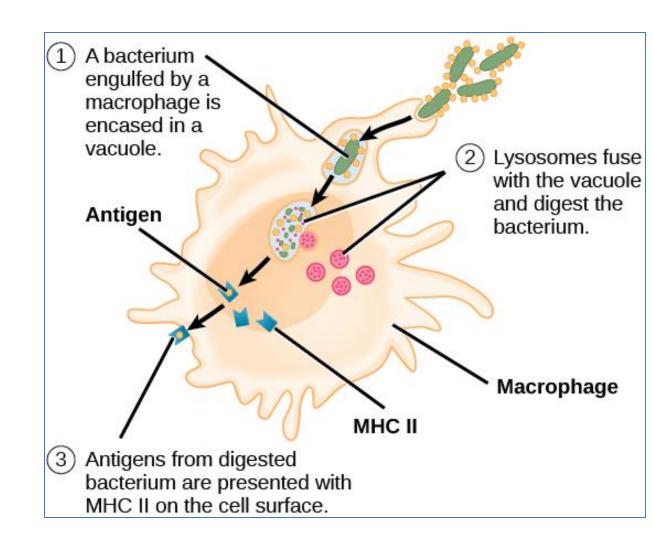
Artistic rendering of the surface of a human dendritic cell illustrating sheet-like processes that fold back onto the membrane surface.

Cells of the immune system / Antigen-Presenting Cells/ Follicular Dendritic Cells

- Follicular dendritic cells (FDCs) are cells with membranous projections that are
  found intermingled in specialized collections of activated B cells, called germinal
  centers, in the lymphoid follicles of the lymph nodes, spleen, and mucosal lymphoid
  tissues.
- FDCs are not derived from precursors in the bone marrow and are of mesenchymal origin, they are non migratory.
- FDCs trap antigens complexed to antibodies or complement products and display these antigens on their surfaces for recognition by B lymphocytes.

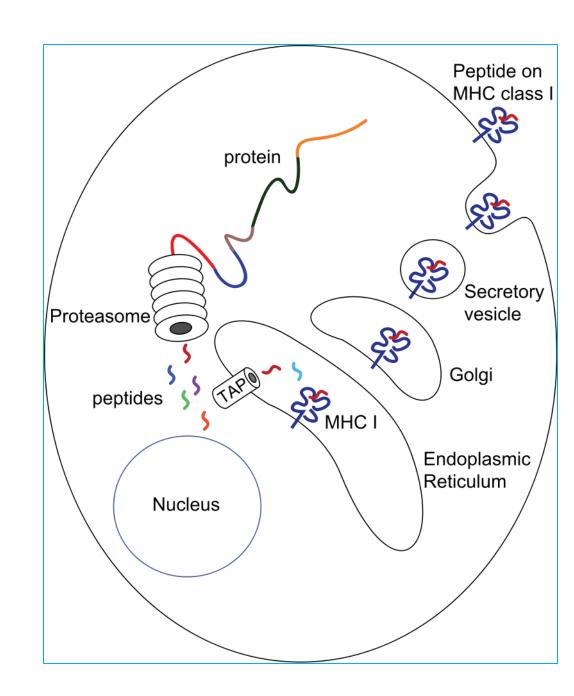
 Macrophages present antigen to helper T lymphocytes at the sites of infection, which leads to helper T cell activation and production of molecules that further activate the macrophages. This process is important for the eradication of microbes that are ingested by the phagocytes but resist killing.

• **B cells present** antigens to helper T cells in lymph nodes and spleen, which is a key step in the cooperation of helper T cells with B cells in humoral immune responses to protein antigens.



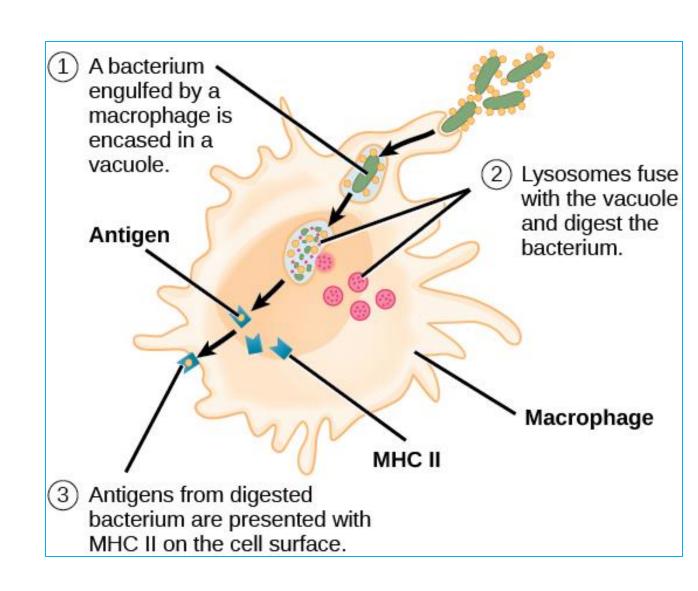
# Major histocompitability complex I

- Their function is to display peptide fragments of proteins from within the cell to cytotoxic T cells; this will trigger an immediate response from the immune system against a particular non-self antigen displayed with the help of an MHC class I protein.
- The proteasome is a macromolecule that consists of 28 subunits, of which half affect proteolytic activity.
- Cells constantly break down proteins and present them on MHC I.



# Major histocompitability complex II

- Normally found only on professional antigenpresenting cells such as dendritic cells, Macrophages, B cells and some endothelial cells.
- The antigens presented by class II peptides are derived from extracellular proteins (not cytosolic as in MHC class I).



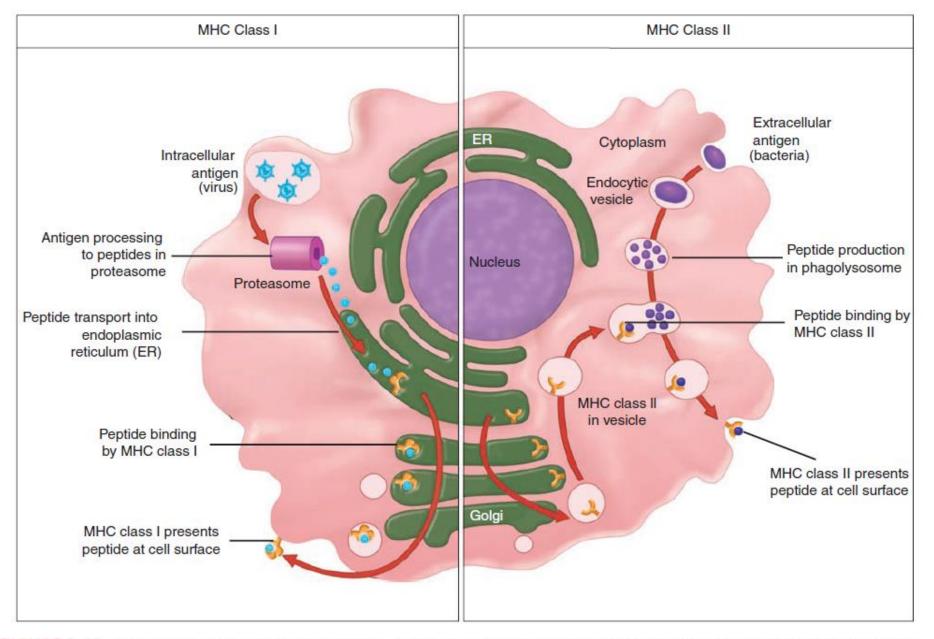
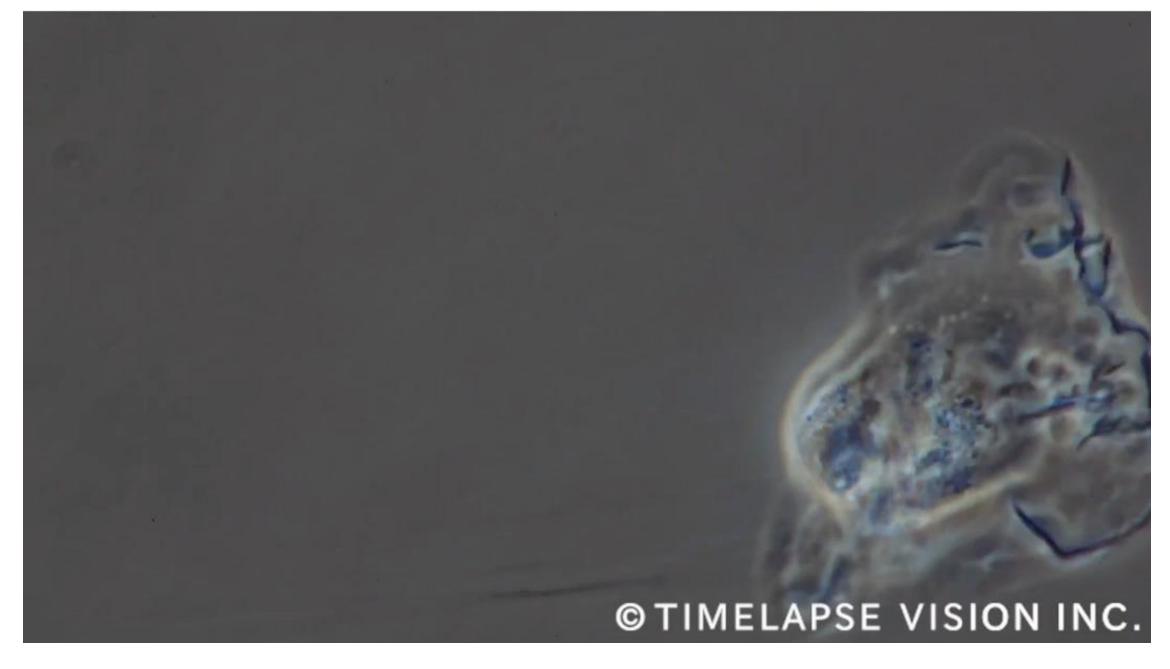


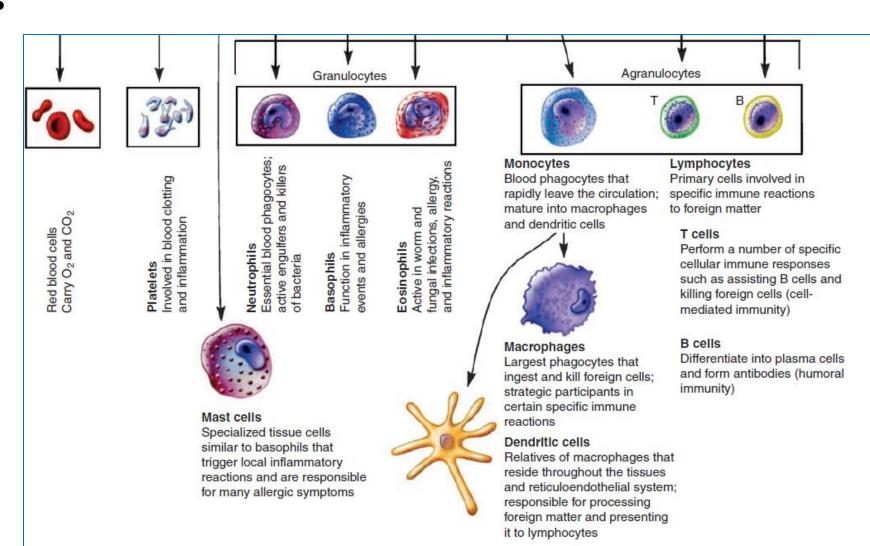
FIGURE 2–II. Antigen processing and presentation. A. Antigens originating in the cytoplasm are digested by the proteasome to peptides. The peptides are bound to the MHC class I molecules in the endoplasmic reticulum (ER) and transported to the surface for presentation. B. Antigens originating outside the cell are endocytosed and digested in the phagolysosome. The digested peptides are bound to MHC class II molecules in the ER and transported to the surface for presentation. MHC, major histocompatibility complex.



Dendritic cell presents antigens to lymphocytes

#### Cells of the immune system

- Phagocytes
- Mast Cells, Basophils, Eosinophils
- Antigen-Presenting Cells
- Lymphocytes



- Lymphocytes consist of distinct subsets that are different in their functions and protein products, but are morphologically similar.
- B (Bursa of Fabricius) lymphocytes originate in the bone marrow and early maturation occurs there.
   Also, T (Thymus) lymphocytes originate in the bone marrow, but mature in the thymus.
- Membrane proteins are used as phenotypic markers to distinguish distinct populations of lymphocytes

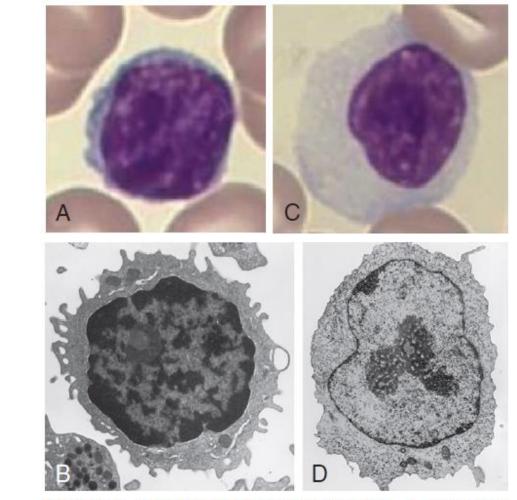
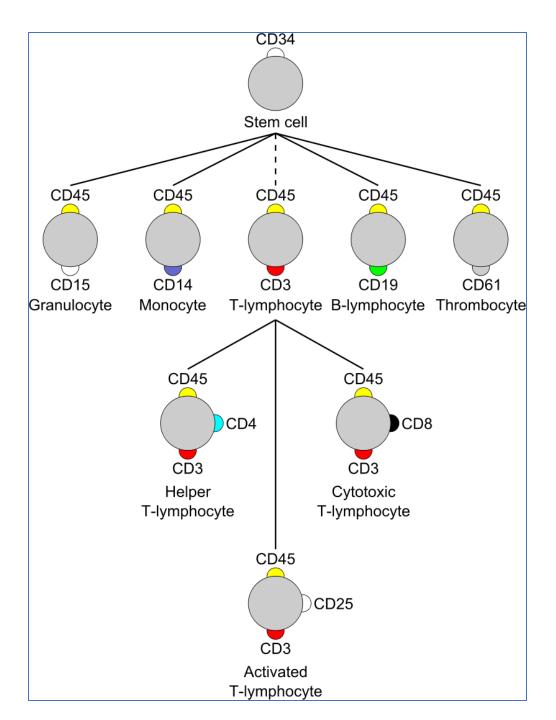


FIGURE 2-7 Morphology of lymphocytes. A, Light micrograph of a lymphocyte in a peripheral blood smear. (Courtesy of Jean Shafer, Department of Pathology, University of California, San Diego. Copyright 1995-2008, Carden Jennings Publishing Co., Ltd.) B, Electron micrograph of a small lymphocyte. (Courtesy of Dr. Noel Weidner, Department of Pathology, University of California, San Diego.) C, Light micrograph of a large lymphocyte (lymphoblast). (Courtesy of Jean Shafer, Department of Pathology, University of California, San Diego. Copyright 1995-2008, Carden Jennings Publishing Co., Ltd.) D, Electron micrograph of a large lymphocyte (lymphoblast). (From Fawcett DW. Bloom and Fawcett: A Textbook of Histology, 12th ed. Chapman & Hall, New York, 1994. With kind permission of Springer Science and Business Media.)

### Cells of the immune system / Cluster of differentiation

- The cluster of differentiation (CD) is a protocol used for the identification and investigation of cell surface molecules providing targets for immunophenotyping of cells.
- In terms of physiology, CD molecules can act in numerous ways, often acting as receptors or ligands important to the cell and some function as adhesion molecules.



Humoral immunity

- B-lymphocytes are the only cells capable of producing antibodies.
- They recognize extracellular (including cell surface) antigens and differentiate into antibody-secreting plasma cells, thus functioning as the mediators of humoral immunity.

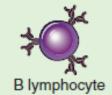


Microbe



Extracellular microbes

Responding lymphocytes



Effector mechanism



Transferred by

Serum (antibodies)

**Functions** 

Block infections and eliminate extracellular microbes

Cells of the immune system / Lymphocytes/ T-lymphocytes

 T lymphocytes, the cells of cell-mediated immunity, recognize the antigens of intracellular microbes and either help phagocytes to destroy these microbes or directly kill the infected cells.

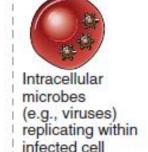
- T cells do not produce antibody molecules.
- Their antigen receptors are membrane molecules distinct from but structurally related to antibodies.

#### Cell-mediated immunity

Microbe



Phagocytosed microbes in macrophage



Responding lymphocytes



Cytotoxic T lymphocyte

Effector mechanism





Transferred by

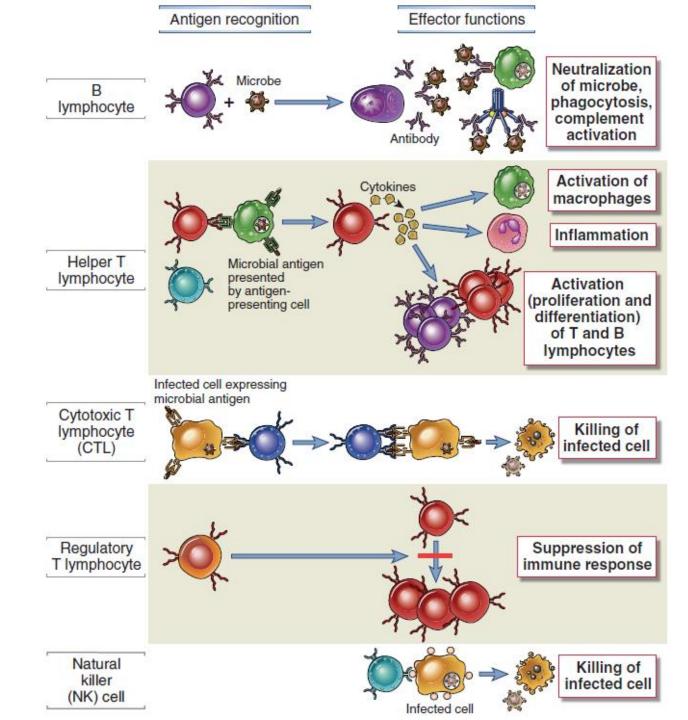
Functions

Cells (T lymphocytes)

Activate macrophages to kill phagocytosed microbes Cells (T lymphocytes)

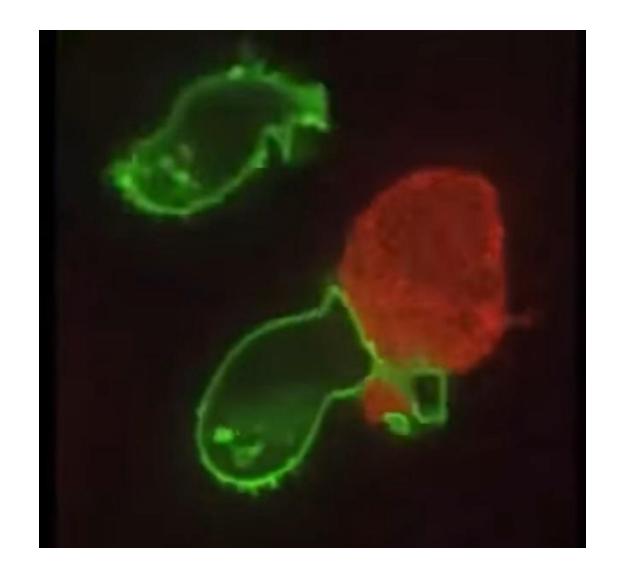
Kill infected cells and eliminate reservoirs of infection

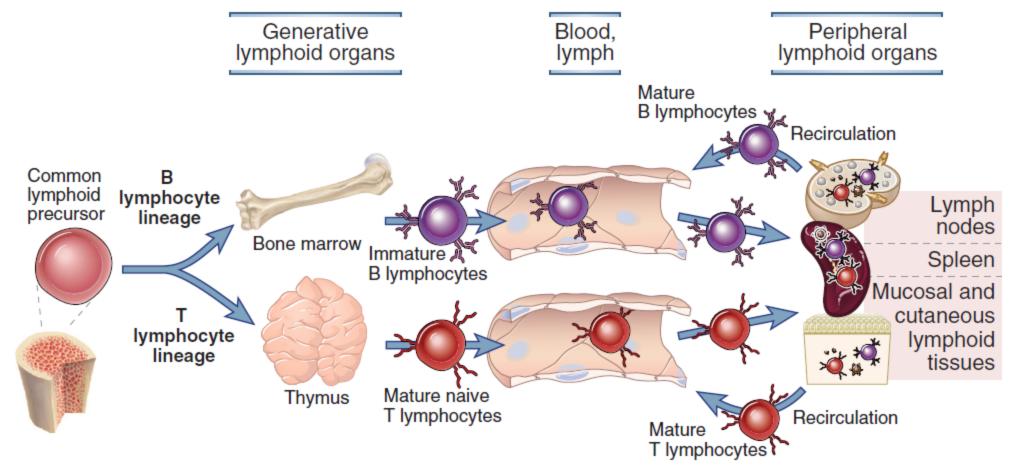
- Humoral immunity is mediated by molecules in the blood and mucosal secretions, called antibodies, which are produced by cells called B lymphocytes.
- Cell-mediated immunity is mediated by T lymphocytes. viruses and some bacteria, survive and proliferate inside phagocytes and other host cells, where they are inaccessible to circulating antibodies. Defense against such infections is a function of cell mediated immunity, which promotes the destruction of microbes residing in phagocytes or the killing of infected cells to eliminate reservoirs of infection.



Cells of the immune system / Lymphocytes/ T-lymphocytes

Through the action of **perforin**, **granzymes** enter the cytoplasm of the target cell and their serine protease function triggers the **caspase** cascade, which eventually lead to **apoptosis** (programmed cell death).

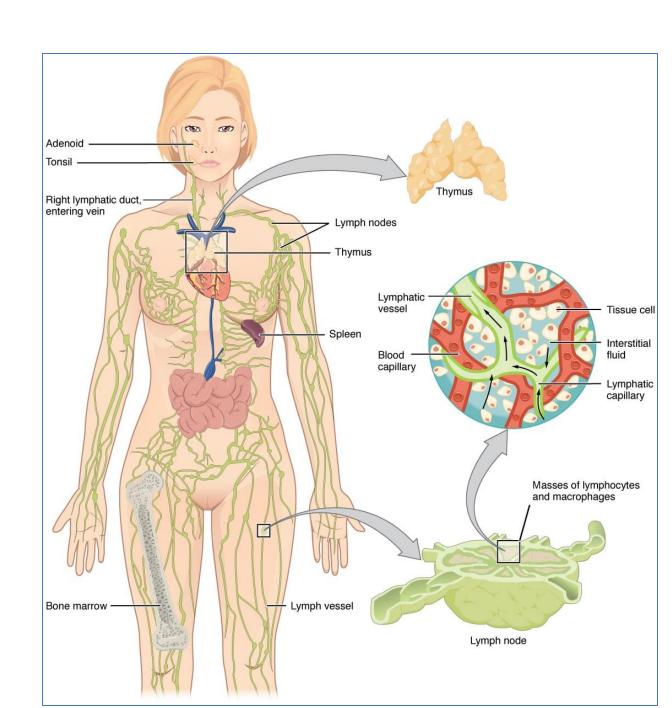




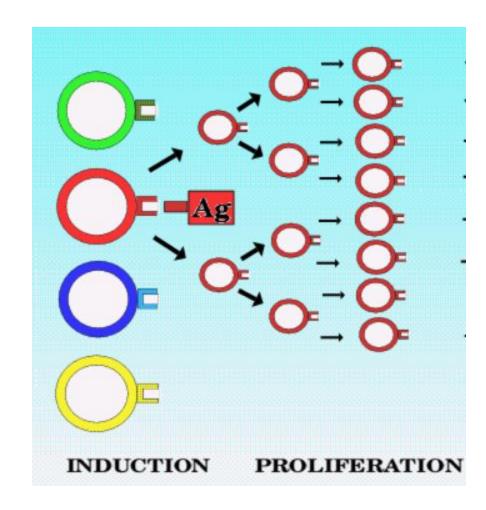
**FIGURE 2–5 Maturation of lymphocytes.** Lymphocytes develop from bone marrow stem cells and mature in the generative lymphoid organs (bone marrow and thymus for B and T cells, respectively) and then circulate through the blood to secondary lymphoid organs (lymph nodes, spleen, regional lymphoid tissues such as mucosa-associated lymphoid tissues). Fully mature T cells leave the thymus, but immature B cells leave the bone marrow and complete their maturation in secondary lymphoid organs. Naive lymphocytes may respond to foreign antigens in these secondary lymphoid tissues or return by lymphatic drainage to the blood and recirculate through other secondary lymphoid organs.

The total number of lymphocytes in a healthy adults about  $5 \times 10^{11}$ . Of these:

- $\sim$ 2% are in the blood,
- $\sim$ 10% in the bone marrow,
- ~15% in the mucosal lymphoid tissues of the gastrointestinal and respiratory tracts, and
- $\sim$ 65% in lymphoid organs (mainly the lymph nodes and spleen)



- Lymphocytes, the unique cells of adaptive immunity, are the only cells in the body that express clonally distributed antigen receptors, each with a fine specificity for a different antigenic determinant.
- Each clone مستنسخ of lymphocytes consists of the **progeny of one cell** and expresses antigen receptors with a **single specificity**.
- There are millions of lymphocyte clones in the body, enabling the organism to recognize and respond to millions of foreign antigens.



 How the enormously diverse repertoire of antigen receptors (millions), and therefore specificities, is generated from a small number of genes for these receptors in the germline\*?

- Genes encoding the **antigen receptors** of lymphocytes are formed by **recombination of DNA segments during the maturation of these cells**. There is a random aspect to these somatic recombination events that results in the generation of millions of different receptor genes and a highly diverse repertoire of antigen specificities among different clones of lymphocytes.
- The antigen receptors are basically antibodies bound to the cell surface.

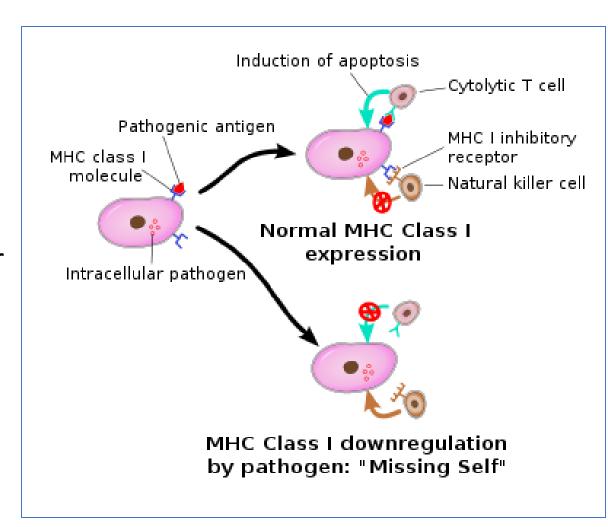
\*Germline DNA: The DNA in germ cells (egg and sperm cells that join to form an embryo). Germline DNA is the source of DNA for all other cells in the body. Also called constitutional DNA.

- In contrast with most organs, such as the heart, which does the same job throughout life, the immune system needs to adapt to an environment that is always changing. This problem is solved by investing in strategies that exploit the power of random change itself.
- Using randomness in this way creates waste, but preserves responsiveness.

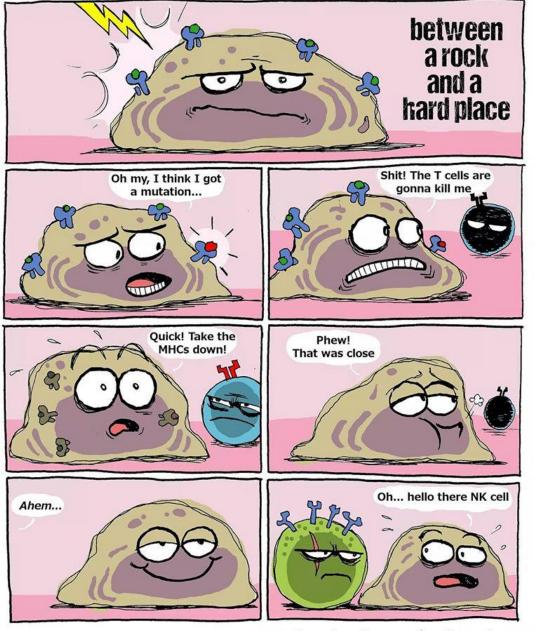
Tonegawa's Nobel Prize work elucidated the genetic mechanism of the adaptive immune system, which had been the central question of immunology for over 100 years. Prior to Tonegawa's discovery, one early idea to explain the adaptive immune system suggested that each gene produces one protein; however, there are under 19,000 genes in the human body which nonetheless can produce millions of antibodies.



- Natural killer (NK) cells are lymphocytes
  distinct from T and B cells that play important
  roles in innate immune responses mainly
  against intracellular viruses and bacteria.
- The term natural killer derives from the fact that these cells are capable of performing their killing function without a need for clonal expansion and differentiation.
- Most NK cells express inhibitory receptors that recognize class I major histocompatibility complex (MHC) molecules, which are cell surface proteins normally expressed on almost all healthy cells in the body



# Cells of the immune system / Lymphocytes / Natural killer (NK) cells



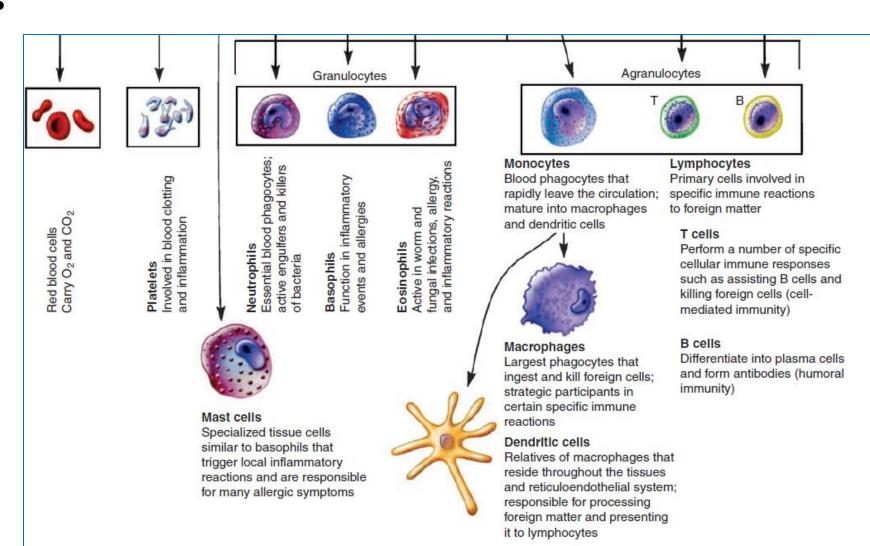
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TABLE 2–2 Lymphocyte Classes

Class	Functions	Antigen Receptor and Specificity	Selected Phenotype Markers	Percentage of Total Lymphocytes (Human)		
				Blood	Lymph Node	Spleer
αβ T lymphocytes						
CD4 <sup>+</sup> helper T lymphocytes	B cell differentiation (humoral immunity) Macrophage activation (cell-mediated immunity) Stimulation of inflammation	αβ heterodimers Diverse specificities for peptide–class II MHC complexes	CD3 <sup>+</sup> , CD4 <sup>+</sup> , CD8 <sup>-</sup>	50-60*	50-60	50-60
CD8 <sup>+</sup> cytotoxic T lymphocytes	Killing of cells infected with viruses or intracellular bacteria; rejection of allografts	αβ heterodimers Diverse specificities for peptide–class I MHC complexes	CD3 <sup>+</sup> , CD4 <sup>+</sup> , CD8 <sup>-</sup>	20-25	15-20	10-15
Regulatory T cells	Suppress function of other T cells (regulation of immune responses, maintenance of self-tolerance)	αβ heterodimers Unresolved	CD3 <sup>+</sup> , CD4 <sup>+</sup> , CD25 <sup>+</sup> (most common, but other phenotypes as well)	Rare	10	10
γδ T lymphocytes	Helper and cytotoxic functions (innate immunity)	γδ heterodimers Limited specificities for peptide and nonpeptide antigens	CD3 <sup>+</sup> , CD4 <sup>+</sup> , and CD8 variable			
B lymphocytes	Antibody production (humoral immunity)	Surface antibody Diverse specificities for all types of molecules	Fc receptors; class II MHC; CD19, CD21	10-15	20-25	40-45
Natural killer cells	Cytotoxic killing of virus- infected or damaged cells (innate immunity)	Various activating and inhibitory receptors Limited specificities for MHC or MHC-like molecules	CD16 (Fc receptor for IgG)	10	Rare	10
NKT cells	Suppress or activate innate and adaptive immune responses	αβ heterodimers Limited specificity for glycolipid-CD1 complexes	CD16 (Fc receptor for IgG); CD3	10	Rare	10

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- Lymphocytes



# **Further reading:**

• Cellular and Molecular Immunology. 7th Edition.. Chapter 2. Cells and tissues of the immune system