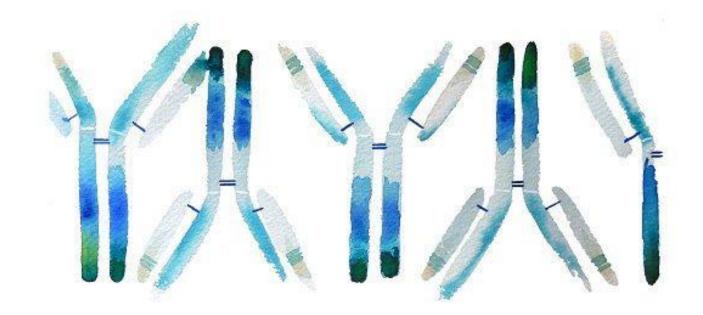
Medical Immunology



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

Lecture 5

What is the process by which phagocytes digest engulfed pathogens or particles within the phagosome using enzymes and reactive oxygen species?

- a) Apoptosis
- b) Exocytosis
- c) Phagolysosome formation
- d) Autophagy

What is the main function of cytotoxic T cells in the immune system?

- a) Phagocytosis of pathogens
- b) Antibody production
- c) Killing infected host cells
- d) Activating B cells

Which of the following cells is part of innate immunity and is of lymphocytic origin?

- a) Dendritic cells
- b) Monocytes
- c) Natural killer cells
- d) B cells
- e) Helper T cells.

Which drug can be used in a patient with a slight fever and the following CBC results?

| Test Name | Results | Reference Range |
|-------------------------|---------|----------------------|
| CBC (INCLUDES DIFF/PLT) | | |
| WHITE BLOOD CELL COUNT | 2800 | 3.8-10.8 Thousand/uL |
| ABSOLUTE NEUTROPHILS | 950 | 1500-7800 cells/uL |
| ABSOLUTE LYMPHOCYTES | 1050 | 850-3900 cells/uL |
| ABSOLUTE MONOCYTES | 519 | 200-950 cells/uL |
| ABSOLUTE EOSINOPHILS | 142 | 15-500 cells/uL |
| ABSOLUTE BASOPHILS | 24 | 0-200 cells/uL |
| NEUTROPHILS | 42% | ę |
| LYMPHOCYTES | 45% | 8 |
| MONOCYTES | 8.8 | 8 |
| EOSINOPHILS | 2.4 | elo |
| BASOPHILS | 0.4 | ę |

- a) The results are normal and no treatment is needed.
- b) Intravenous antibodies
- c) IL-12
- d) Granulocyte colony stimulating factor

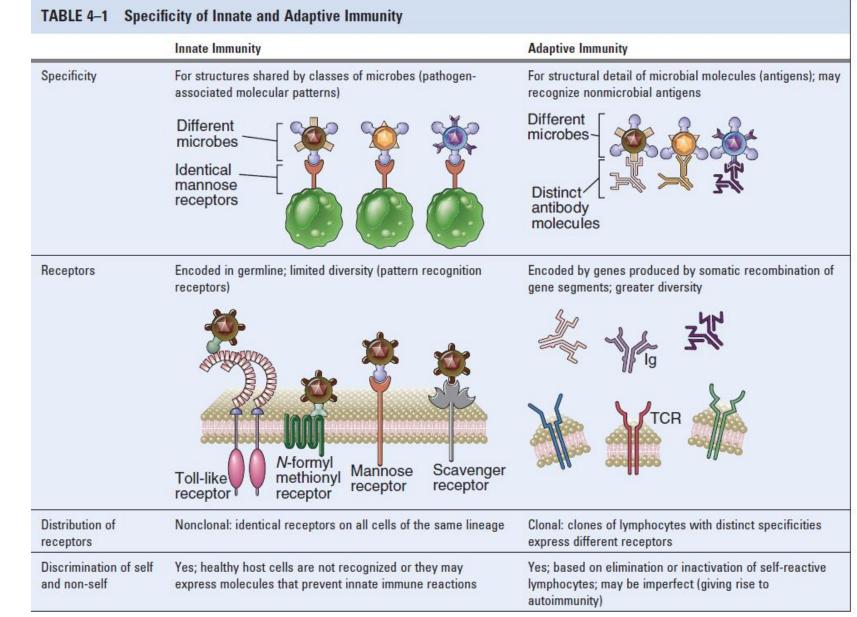
Which drug can be used in a patient with a slight fever and the following CBC results?

| Test Name | Results | Reference Range |
|-------------------------|---------|----------------------|
| CBC (INCLUDES DIFF/PLT) | | |
| WHITE BLOOD CELL COUNT | 5.9 | 3.8-10.8 Thousand/uL |
| ABSOLUTE NEUTROPHILS | 4165 | 1500-7800 cells/uL |
| ABSOLUTE LYMPHOCYTES | 1050 | 850-3900 cells/uL |
| ABSOLUTE MONOCYTES | 519 | 200-950 cells/uL |
| ABSOLUTE EOSINOPHILS | 142 | 15-500 cells/uL |
| ABSOLUTE BASOPHILS | 24 | 0-200 cells/uL |
| NEUTROPHILS | 70.6 | ę. |
| LYMPHOCYTES | 17.8 | 8 |
| MONOCYTES | 8.8 | es. |
| EOSINOPHILS | 2.4 | de la |
| BASOPHILS | 0.4 | ę. |

- a) The results are normal and no treatment is needed.
- b) Intravenous antibodies
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- d) Granulocyte colony stimulating factor

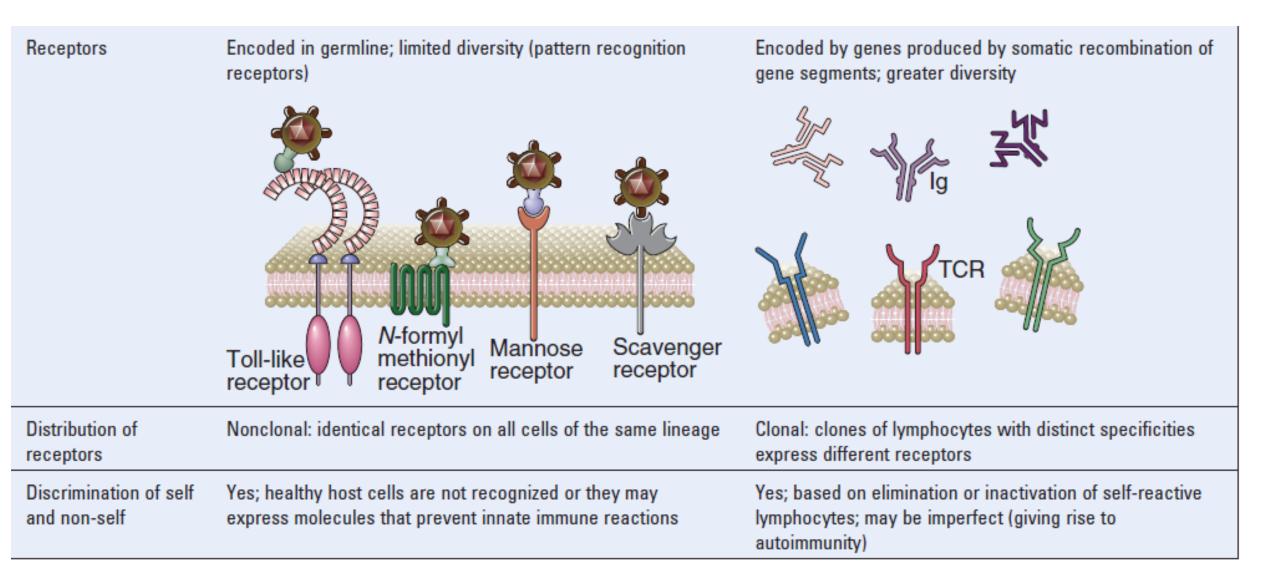
- Much of the interactions between cells of the immune system, and between the immune system and foreign introducers depend on the action of cell bound and secreted molecules.
- In this lecture we will discuss some of those molecules.
- Main topics:
 DAMPs and PAMPs
 TLRs.
- NLRs and the inflammasome.
- RLRs.
- Major inflammatory cytokines (TNF, IL-1, IL-6).

- The cells and soluble molecules of innate immunity either exist in a fully functional state before encounter with microbes or are rapidly activated by microbes
- The innate immune system recognizes molecular structures that are **characteristic of microbial pathogens but not mammalian cells.**
- The innate immune system recognizes microbial products that are often **essential for survival of the microbes.**
- The microbial substances that stimulate innate immunity are called **pathogen-associated molecular patterns (PAMPs).**
- Different classes of microbes (e.g., viruses, gram-negative bacteria, gram positive bacteria, fungi) express different PAMPs.



It is estimated that the innate immune system can recognize about 10³ molecular patterns. In contrast, the adaptive immune system is capable of recognizing 10⁷ or more distinct antigens.

| TABLE 4–1 | Specificity of Innate and Adaptive Immunity | |
|-------------|---|--|
| | Innate Immunity | Adaptive Immunity |
| Specificity | For structures shared by classes of microbes (pathogen- associated molecular patterns) | For structural detail of microbial molecules (antigens); may recognize nonmicrobial antigens |
| | Different microbes | Different microbes- Distinct antibody molecules |



• Characteristics of antigens recognized:

Nucleic acids that are unique to microbes, such as double-stranded RNA found in replicating viruses and unmethylated CpG DNA sequences found in bacteria

Proteins that are found in microbes, such as initiation by N-formylmethionine, which is typical of bacterial proteins.

Complex lipids and carbohydrates that are synthesized by microbes but not by mammalian cells, such as lipopolysaccharide (LPS) in gram-negative bacteria, **lipoteichoic acid** or peptidoglycan (PGN) in gram positive bacteria, and mannose-rich **oligosaccharides**.

 limited number of fundamental differences between microbial molecules and the molecules that higher organisms produce. Thus, the innate immune system has evolved to recognize only a limited number of molecules.

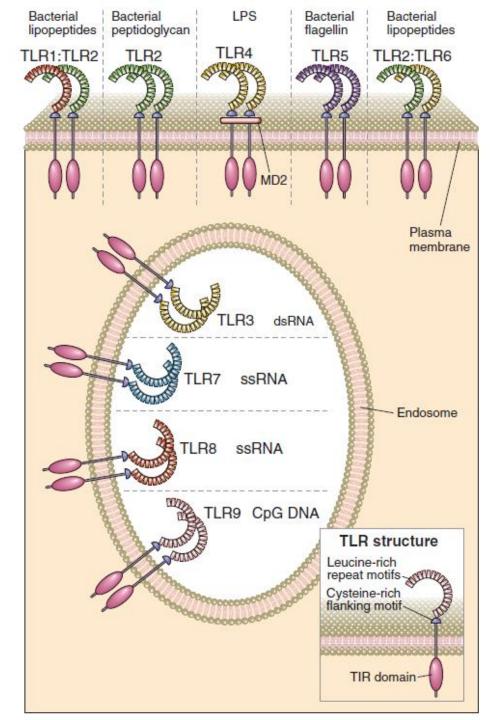
- The innate immune system also recognizes endogenous molecules that are produced by or released from damaged and dying cells. These substances are called damage associated molecular patterns (DAMPs).
- DAMPs may be produced as a result of cell damage caused by infections, but they may also indicate sterile injury to cells caused by any of myriad reasons, such as chemical toxins, burns, trauma, or decreased blood supply.
- DAMPs are generally not released from cells dying by apoptosis. In some cases, healthy
 cells of the immune system are stimulated to produce and release DAMPs, which enhances
 an innate immune response to infections.

| TABLE 4–2 Examples of PAMPs and DAMPs | | | |
|---|--------------------------|--|--|
| Pathogen-Associated Molecular Patterns Microbe Type | | | |
| Nucleic acids | ssRNA dsRNA CpG | Virus Virus Virus, bacteria | |
| Proteins | Pilin Flagellin | Bacteria Bacteria | |
| Cell wall lipids | LPS Lipoteichoic acid | Gram-negative bacteria Gram-positive bacteria | |
| Carbohydrates | Mannan Dectin glucans | Fungi, bacteria Fungi | |
| Damage-Associated Molecular Patterns | | | |
| Stress-induced proteins | HSPs | | |
| Crystals | Monosodium urate | | |
| Nuclear proteins | HMGB1 | | |
| CpG, cytidine-guanine dinucleotide; dsRNA, double-stranded RNA; HMGB1, high-mobility group box 1; HSPs, heat shock proteins; LPS, lipopolysaccharide; ssRNA, single-stranded RNA. | | | |

- **Pattern recognition receptors (PRRs)** play a crucial role in the proper function of the innate immune system. PRRs are germline-encoded host sensors, which detect molecules typical for the pathogens.
- They are proteins expressed, mainly, by cells of the innate immune system, such as dendritic cells, macrophages, monocytes, neutrophils and epithelial cells, to identify two classes of molecules: **PAMPs** and **DAMPs**.
- PRR can be **cell bound** or **soluble**.
- Cell bound PRR can be found on **different compartments of the cell**. (membrane, cytosol)

Molecules of the immune system / cell bound PRR/ TLRs

- Toll-like receptors (TLR), are proteins that respond to the presence of pathogenic microbes by activating antimicrobial defense mechanisms in the cells in which they are expressed.
- TLR are found in every life form in the evolutionary tree from insects up to mammals.
- TLRs are also involved in response to endogenous molecules whose expression or location indicates cell damage (DAMP).
- Ligand binding to the leucine-rich domains causes physical interactions between TLR molecules and the formation of TLR dimers.

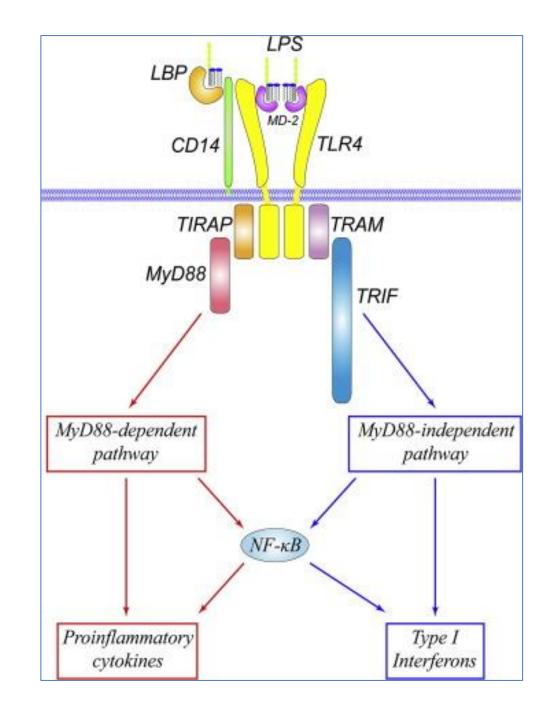


Toll-like receptors (TLRs) play a crucial role in innate immunity by recognizing:

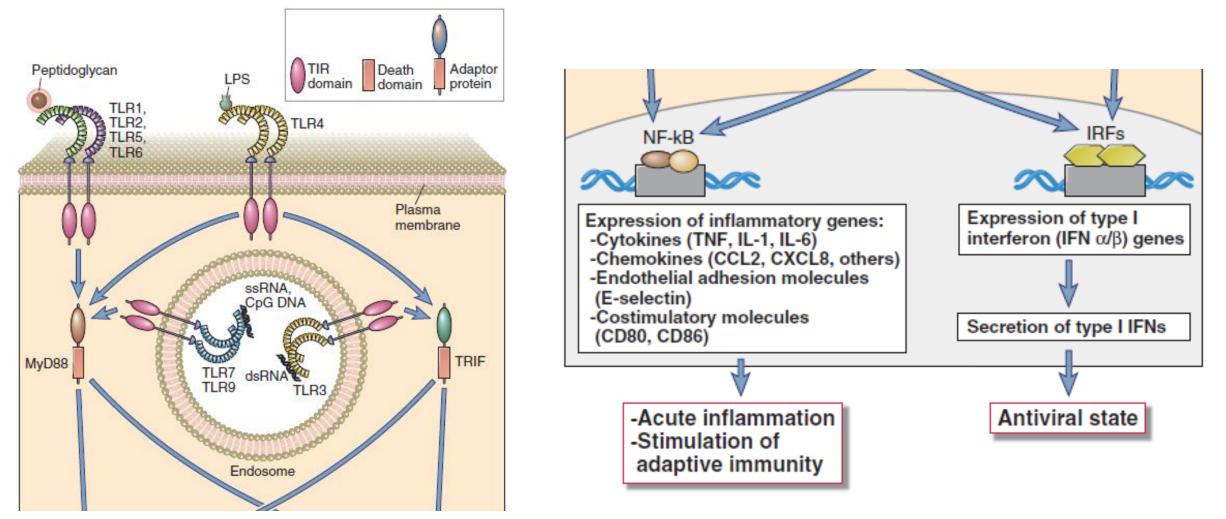
- A. Self-antigens
- B. Pattern recognition molecules
- C. T-cell receptors
- D. MHC class II molecules
- E. pathogen-associated molecular patterns (PAMPs).

Molecules of the immune system / cell bound PRR/ TLRs

- Adapter and accessory molecules can be needed for proper signalling.
- An extracellular protein called **MD2** (myeloid differentiation protein 2) binds the lipid A component of **LPS**, forming a complex that then interacts with **TLR4** and initiates signaling.
- Another protein called CD14 is also required for efficient LPS-induced signaling.
- Both CD14 and MD2 can also associate with other TLRs.



Molecules of the immune system / cell bound PRR/ TLRs



- adaptor proteins (MyD88, TRIF) facilitate the recruitment and activation of various protein kinases, leading to the activation of different transcription factors.
- All TLRs except TLR3 signal through MyD88 and are therefore capable of activating NF-κB and inducing an inflammatory response. TLR3 signals through TRIF and therefore activates IRF3 and induces expression of type I interferons.

How many different Toll-like receptors (TLRs) have been identified in humans?

- a) 2
- b) 5
- c) 10
- d) 20

Molecules of the immune system / cell bound PRR/ Other receptors

- Receptors for Carbohydrates recognize carbohydrates on the surface of microbes, they
 facilitate the phagocytosis of the microbes and stimulate subsequent adaptive immune
 responses. These receptors belong to the C-type lectin family, so called because they bind
 carbohydrates (hence, lectins) in a Ca++-dependent manner (hence, C-type). Some of these
 are soluble proteins found in the blood and extracellular fluids; others are integral membrane
 proteins found on the surfaces of macrophages, dendritic cells, and some tissue cells.
 (examples, mannose and dectin receptors).
- Scavenger receptors comprise a structurally and functionally diverse collection of cell surface proteins found mainly on macrophages.
- N-Formyl met-leu-phe receptors, expressed by neutrophils and macrophages, recognize bacterial peptides containing N-formylmethionyl residues and stimulate directed movement of the cells. (i.e those residues are chemoattractans that help phagocytic cells trace the bacteria producing it)

Molecules of the immune system / cytoplasmic PRR

- In addition to the membrane-bound TLRs, which sense pathogens outside cells or in endosomes, the innate immune system has evolved to equip cells with pattern recognition receptors that detect infection or cell damage in the cytoplasm.
- The two major classes of these cytoplasmic receptors are **NOD-like receptors** and **RIG-like receptors**. These cytoplasmic receptors, like TLRs, are linked to signal transduction pathways that promote inflammation or type I interferon production.
- The normal life cycles of some microbes, such as viral gene translation and viral particle assembly, **take place in the cytoplasm**.
- Some microbes can produce toxins that create pores in host cell plasma membranes, including endosomal membranes, through which microbial molecules can enter the cytoplasm.

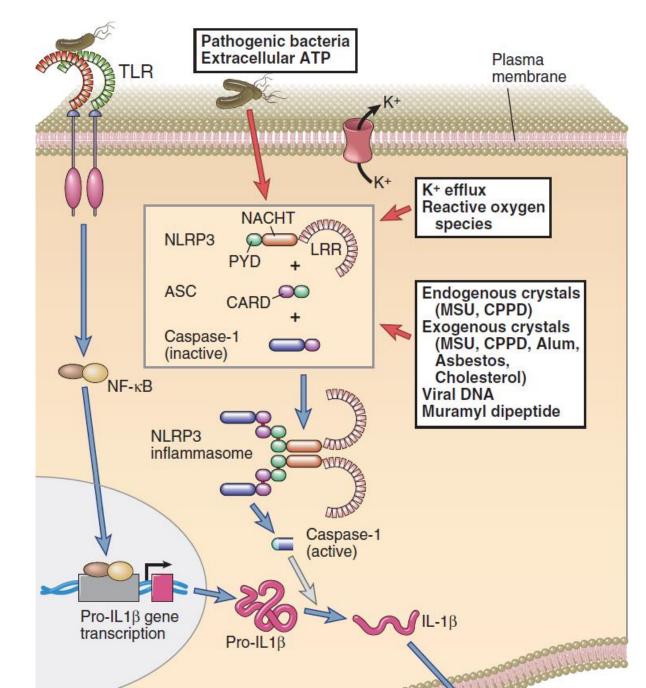
Molecules of the immune system / cytoplasmic PRR/ NOD-like receptors (NLRs)

- NOD-like receptors (NLRs) are a family of more than 20 different cytosolic proteins, some of which sense cytoplasmic PAMPs and DAMPs and recruit other proteins to form signaling complexes that promote inflammation.
- **NOD1** and **NOD2**, are expressed in the cytoplasm of several cell types including mucosal epithelial cells and phagocytes, and they respond to bacterial cell wall peptidoglycans.
- The NLRP* subfamily of NLRs respond to cytoplasmic PAMPs and DAMPs by forming signaling complexes called inflammasomes, which generate active forms of the inflammatory cytokine IL-1.

*(NLR family, pyrin-domain-containing proteins)

Molecules of the immune system / cytoplasmic PRR/ NOD-like receptors (NLRs)

 The inflammasome is a multiprotein intracellular complex that detects pathogenic microorganisms and sterile stressors, and that activates the highly pro-inflammatory cytokines interleukin-1b (IL-1b) and IL-18. Dysregulation of inflammasomes is associated with a number of autoimmune diseases.



Molecules of the immune system / cytoplasmic PRR/ NOD-like receptors (NLRs)



NEWS • 10 MAY 2019 • CLARIFICATION 24 MAY 2019

NLRP3 inhibitors stoke anti-inflammatory ambitions

Inhibitors of the innate immune system's NLRP3 inflammasome promise potential in Parkinson disease, Alzheimer disease, non-alcoholic steatohepatitis, gout and much more, catching the eye of Novartis, Genentech and others.

Molecules of the immune system / cytoplasmic PRR/ RIG-like receptors (RLRs)

- **RIG-like receptors*** RLRs are **cytosolic sensors of viral RNA** that respond to viral nucleic acids by inducing the production of the antiviral type I interferons.
- RLRs can recognize double-stranded and single-stranded RNA, which includes the genomes of RNA viruses and RNA transcripts of RNA and DNA viruses
- RLRs also can discriminate viral single-stranded RNA from normal cellular single-stranded RNA transcripts.
- RLRs are expressed in a wide variety of cell types, including bone marrow–derived leukocytes and various tissue cells.

*RIG (retinoic acid–inducible gene)

Which NOD-like receptor (NLR) is associated with inflammasome formation and the release of pro-inflammatory cytokines like IL-1 β ?

a) NOD1

b) NOD2

c) NLRP3

d) NLRP6

Molecules of the immune system / cell bound PRR

| Cell-Associated Pattern Recognition Receptors | Location | Specific Examples | PAMP/DAMP Ligands |
|--|---|--|--|
| Toll-like receptors (TLRs) | Plasma membrane and endosomal membranes of dendritic cells, phagocytes, B cells endothelial cells, and many other cell types | TLRs 1-9 | Various microbial molecules including bacterial LPS and peptidoglycans, viral nucleic acids |
| NOD-like receptors (NLRs) | Cytoplasm of phagocytes epithelial cells, and other cells | NOD1/2 NALP family (inflammasomes) | Bacterial cell wall peptidoglycans Flagellin, muramyl dipeptide, LPS; urate crystals; products of damaged cells |
| RIG-like receptors (RLRs) | Cytoplasm of phagocytes and other cells | RIG-1, MDA-5 | Viral RNA |
| C-type lectin–like receptors | Plasma membranes of phagocytes | Mannose receptor | Microbial surface carbohydrates with terminal mannose and fructose |
| 8-8 | | Dectin | Glucans present in fungal cell walls |
| Scavenger receptors | Plasma membranes of phagocytes | CD36 | Microbial diacylglycerides |
| N-Formyl met-leu-phe receptors | Plasma membranes of phagocytes | FPR and FPRL1 | Peptides containing <i>N</i> -formylmethionyl residues |

Molecules of the immune system / The Major Proinflammatory Cytokines

- One of the earliest responses of the innate immune system to infection and tissue damage is the secretion of cytokines by tissue cells, which is critical for the acute inflammatory response.
- Three of the most important proinflammatory cytokines of the innate immune system are **TNF**, **IL-1**, and **IL-6**.
- Tissue **macrophages** and **mast cells** are the major source of these cytokines, although other cell types, including **endothelial** and **epithelial cells**, can also produce IL-1 and IL-6.

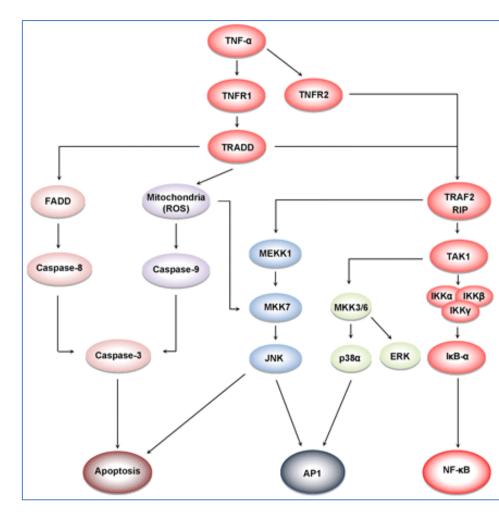
Molecules of the immune system / Important cytokines

| | TABLE 2-2 | Some Cytokines Acting in Infection | |
|--|--|---|---|
| | | CELL SOURCE | FUNCTIONS |
| Cytokines are a | Interleukins (IL |) | |
| broad and loose category of small | IL-1 | Macrophages, endothelium, fibroblasts, epithelial | Differentiation and function of immune effectors, PMN response ($T_{_{\rm H}}$ 17) |
| proteins that are | IL-2 | T cells $(T_H I)$ | T-cell proliferation, cytolytic activity of natural killer (NK) cells |
| important in cell signalling. | IL-4 | T cells (T _H 2), macrophages, B cells | Differentiation of naïve T cells to helper T cells, prolif- eration of B cells |
| | IL-5 | T cells (T _H 2) | Eosinophil activation |
| Cytokines include chemokines , | IL-8 | Macrophages, endothelial,T cells, keratinocytes, PMNs | Chemoattractant for PMNs and T cells, PMN degranu- lation, migration of PMNs |
| interferons, | IL-17 | T cells (T _H I7) | Inflammation, PMN response |
| interleukins, | IL-22 | T cells (T _H 17) | Antimicrobial peptides |
| lymphokines, and | Interferons (IFN | ۷) | |
| tumour necrosis | IFN-α/β | T cells, B cells, macrophages, fibroblasts | Antiviral activity, stimulates macrophages, MHC class I expression |
| factors | IFN-γ | T cells (T _H I, CTLs), NK cells | T-cell activation, macrophage activation, PMNs, NK cells, antiviral, MHC class I and II expression |
| Cytokines are | Tumor Necrosis | s Factor (TNF) | |
| produced by a broad range of cells, | TNF-α | T cells, macrophages, NK cells | Expression of multiple cytokines, (growth and tran- scription factors), stimulates inflammatory response, cytotoxic for tumor cells |
| including immune | TNF-β | T cells, B cells | Same as TNF- $lpha$ |
| and non-immune | MUC Mainta history | | |
| cells | MHC, Major histocompatibility complex; PMN, Polymorphonuclear neutrophil | | |

cells

Molecules of the immune system / The Major Proinflammatory Cytokines/ TNF

- Tumor necrosis factor (TNF) is a mediator of the acute inflammatory response to bacteria and other infectious microbes.
- TNF production by macrophages is stimulated by PAMPs and DAMPs. TLRs, NLRs, and RLRs can all induce TNF gene expression, in part by activation of the NF-κB transcription factor.
- TNF can also mediate cell proliferation and in some cases cell death.
- TNF superfamily plays highly diversified roles in the body.



Molecules of the immune system / The Major Proinflammatory Cytokines/ IL-1

- Interleukin-1 (IL-1) is also a mediator of the acute inflammatory response and has many similar actions as TNF.
- Unlike TNF, IL-1 is also produced by many cell types other than macrophages, such as neutrophils, epithelial cells (e.g., keratinocytes), and endothelial cells.
- There are two forms of IL-1, called IL-1α and IL-1β, The main biologically active secreted form is IL-1β.
- IL-1β gene transcription is induced by TLR and NOD signaling pathways that activate NF-κB, whereas pro–IL- 1β cleavage is mediated by the NLRP3 inflammasome.
- IL-1 mediates its biologic effects through a membrane receptor called the type I IL-1 receptor

Molecules of the immune system / The Major Proinflammatory Cytokines/ IL-6

 IL-6 is another important cytokine in acute inflammatory responses that has both local and systemic effects, including the induction of liver synthesis of a variety of other inflammatory mediators, the stimulation of neutrophil production in the bone marrow, and the differentiation of IL-17–producing helper T cells

Molecules of the immune system / The Major Proinflammatory Cytokines

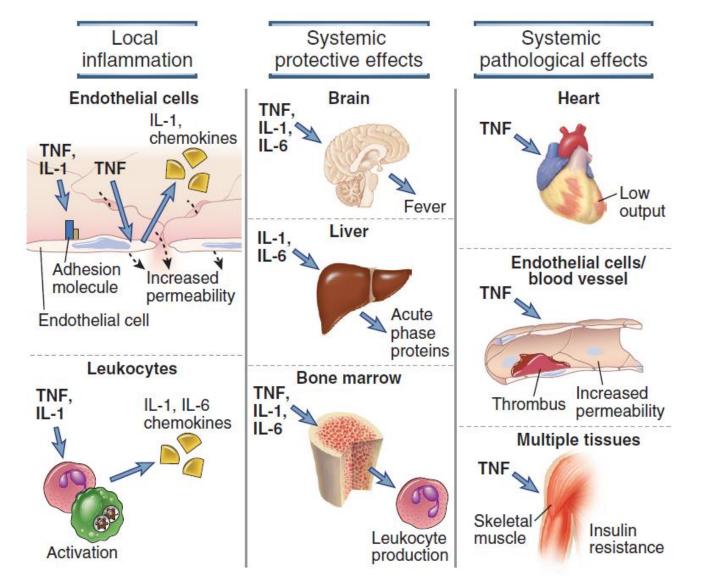
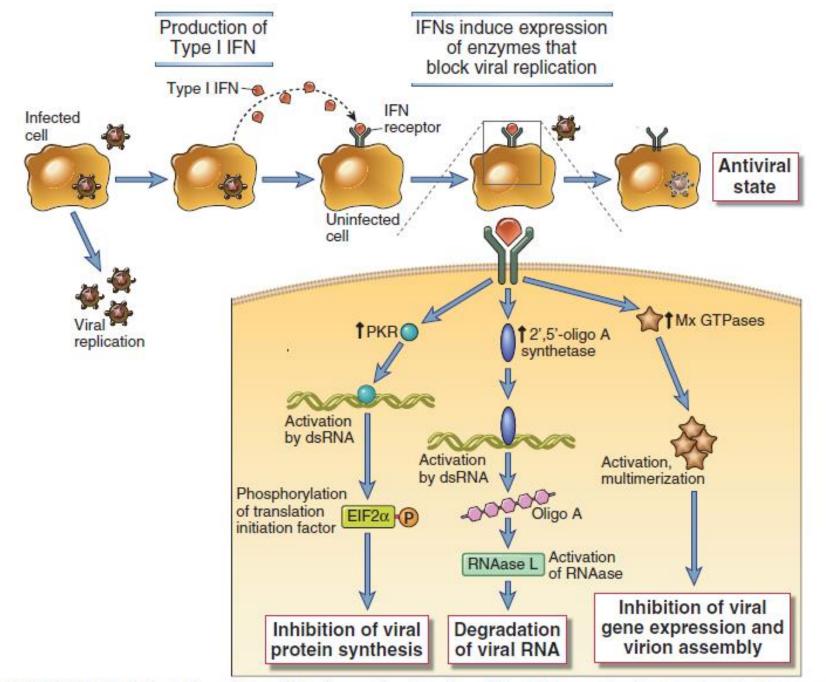


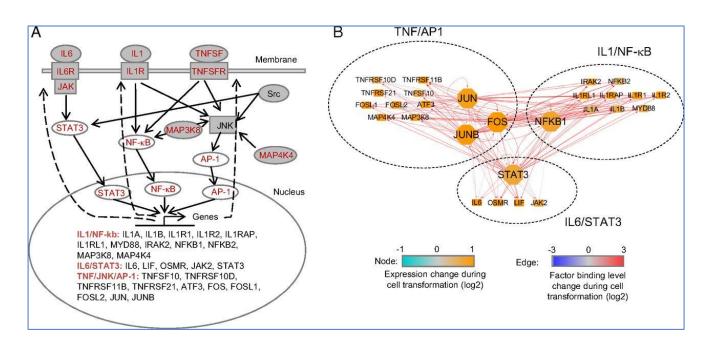
FIGURE 4–14 Local and systemic actions of cytokines in inflammation. TNF, IL-1, and IL-6 have multiple local and systemic inflammatory effects. TNF and IL-1 act on leukocytes and endothelium to induce acute inflammation, and both cytokines induce the expression of IL-6 from leukocytes and other cell types. TNF, IL-1, and IL-6 mediate protective systemic effects of inflammation, including induction of fever, acute-phase protein synthesis by the liver, and increased production of leukocytes by the bone marrow. Systemic TNF can cause the pathologic abnormalities that lead to septic shock, including decreased cardiac function, thrombosis, capillary leak, and metabolic abnormalities due to insulin resistance. Molecules of the immune system / Important cytokines / Interferons

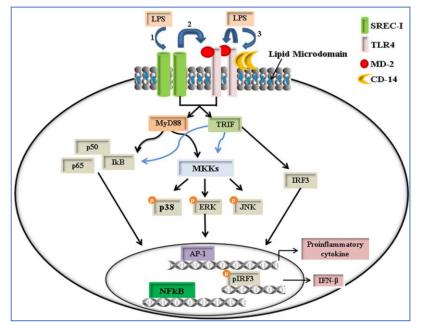
- The major way by which the innate immune system deals with viral infections is to induce the expression of type I interferons. Type I interferons are a large family of structurally related cytokines that mediate the early innate immune response to viral infections.
- Type I interferons, signaling through the type I interferon receptor, activate transcription of several genes that confer on the cells a resistance to viral infection, called an **antiviral state**.
- Type I interferons cause **sequestration of lymphocytes in lymph nodes**, thus maximizing the opportunity for encounter with microbial antigens.
- Type I interferons increase the cytotoxicity of NK cells and CD8+ CTLs
- Upregulate expression of class I MHC molecules and thereby increase the probability that virally infected cells will be recognized and killed by CD8+ CTLs.

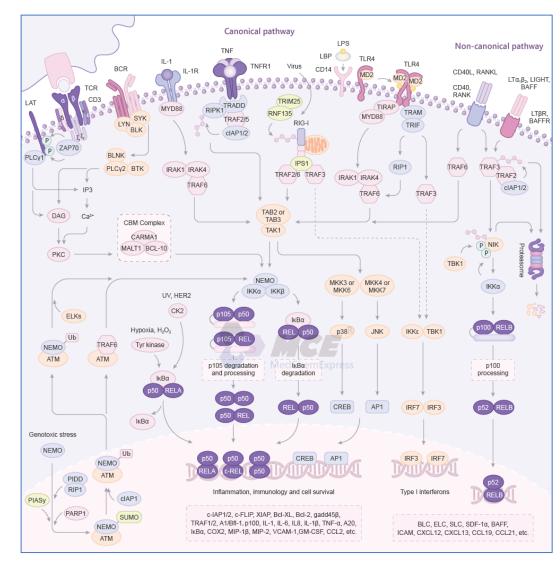
Molecules of the immune system / Important cytokines / Interferons



Molecules of the immune system / The Major Proinflammatory Cytokines







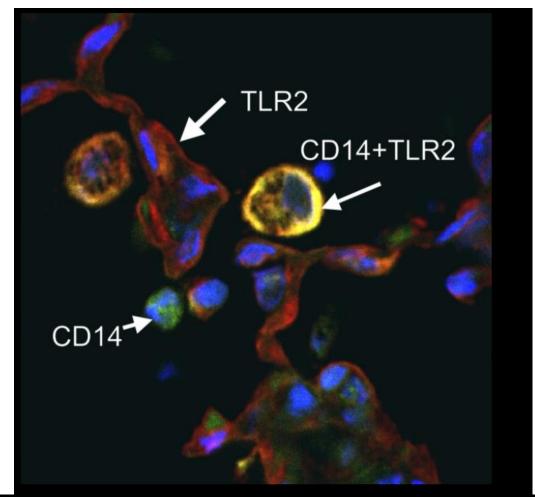


Figure 3. TLR2 and CD14 in the lungs of a rabbit. TLR2 is labeled *red* and CD14 is labeled *green*. Colocalization of TLR2 and CD14 is shown in *yellow*. TLR2 is visible on the alveolar epithelium and on alveolar macrophages in the airspace. CD14 is visible on alveolar macrophages, and neutrophils in the intravascular and alveolar space. The *bright yellow* alveolar macrophage shows high levels of expression of both TLR2 and CD14. Similar results are found when the sections are labeled for TLR4 and CD14.

Further reading:

• Cellular and Molecular Immunology. 7th Edition.. Chapter 4. Innate immunity

CELL-ASSOCIATED PATTERN RECOGNITION RECEPTORS OF INNATE IMMUNITY

SOLUBLE RECOGNITION AND EFFECTOR MOLECULES OF INNATE IMMUNITY