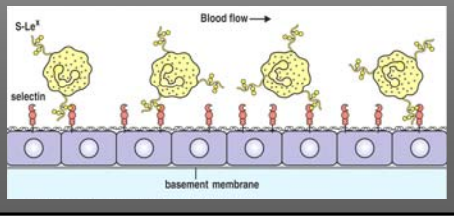
A Day in the Life of a Phagocytic Leukocyte

Selectin-mediated Adhesion is Weak and Promotes "Rolling" of Leukocyte Along Endothelia

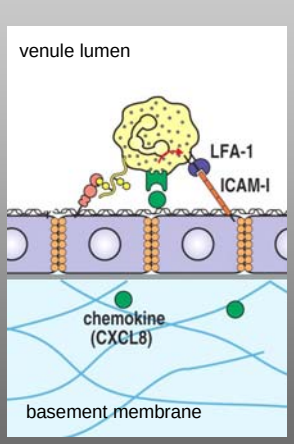
QuickTime™ and a
Sorenson Video decompressor
are needed to see this picture.



Movie, courtesy T. Springer

Firm Adhesion is Triggered by Chemokine Activation of Leukocyte Integrins

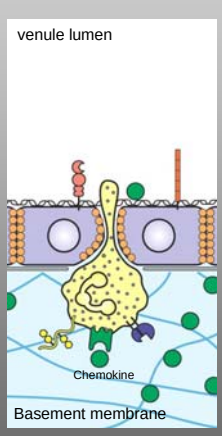
QuickTime™ and a
Sorenson Video decompressor
are needed to see this picture.



Movie, courtesy T. Springer

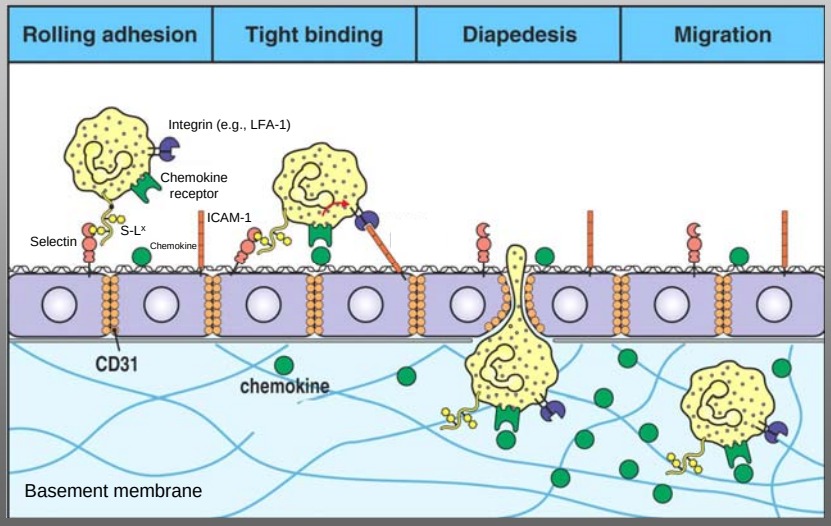
Diapedesis: Crawling Through Endothelial Junctions and Into the Tissue

QuickTime™ and a
Sorenson Video decompressor
are needed to see this picture.



Movie, courtesy T. Springer

Leukocyte Migration, Start to Finish



Modified from: Parham, *The Immune System, 2nd ed.* (Garland: New York), 2005

The Innate Immune Response to Bacterial and Fungal Infections

Relative Risk of Death Associated With Death of a Biological Parent Before the Age of 50

Cause of Death	Relative Risk
All causes	1.7
"Natural causes"	2.0
Infectious	5.8
Cardiovascular	4.5
Cancer	1.2

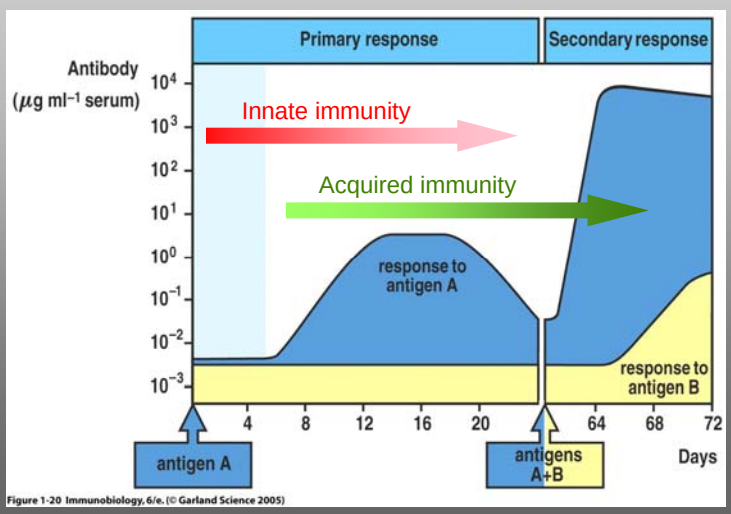
Conclusion: Genes that determine responses to infectious agents have a disproportionate effect on mortality

Source: Sorensen et al., *New Engl. J. Med.*, 318:727, 1988

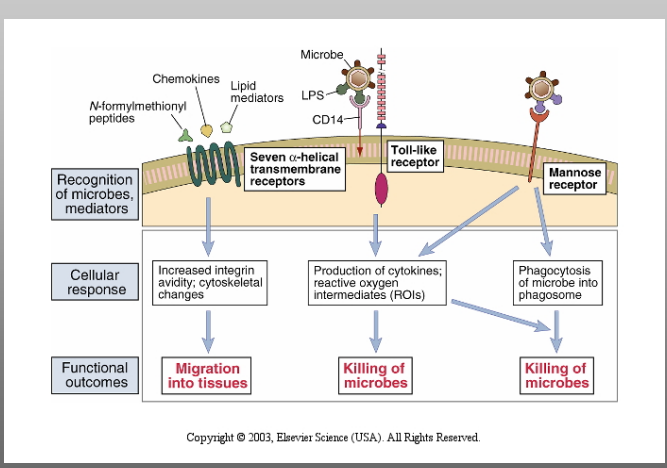
Distinctions Between Innate and Adaptive Immunity

	Innate immune system	Adaptive immune system
Receptors	Germline-encoded	Somatically engineered
Distribution	Non-clonal	Clonal
Kinetics	Rapid	Slow (requires clonal expansion)
Specificity	Recognizes non-self "pattern recognition"	Recognizes "altered self" Primary structure (TCR) Higher order structure (Immunoglobulin; BCR)
Effector Cells	All	Primarily lymphocytes, DCs, Mϕ

What Really Happens During the Lag Period Before the Acquired Immune Response?

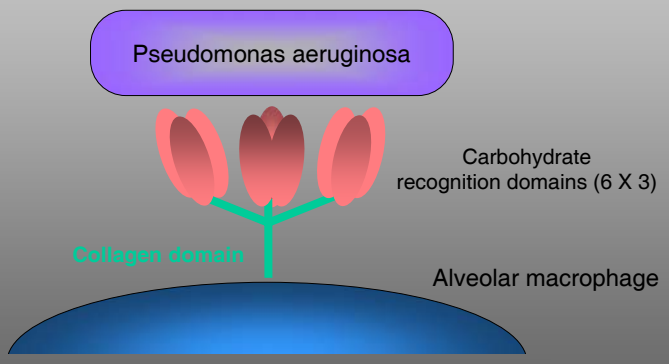


Receptors Important in Innate Immunity



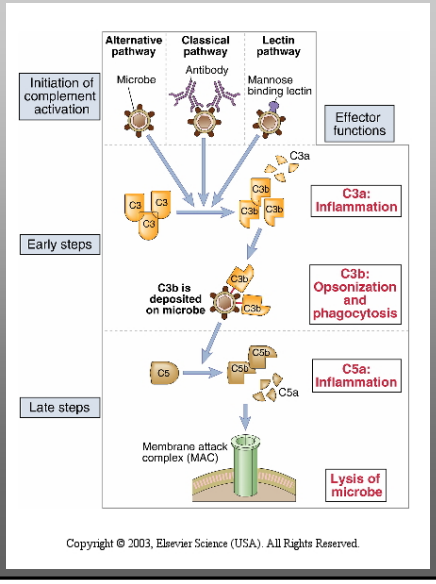
Soluble "Defense Collagens" Participate in Innate Immunity

Domain Structure of Surfactant Protein A (SP-A), a Lung Soluble Defense Collagen (Collectin)

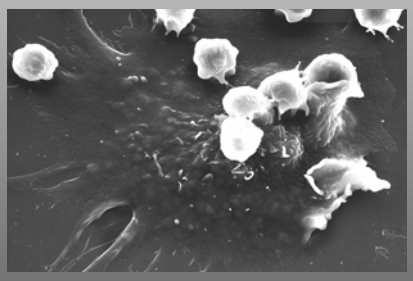


The Complement System is Critical for Innate Immunity and is Triggered by Multiple Ligands

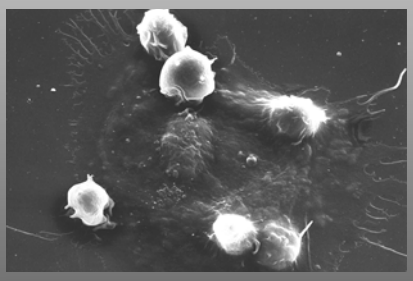
Complement is a Group of Proteins that Deposit On Microbial Pathogens and Help Kill Them



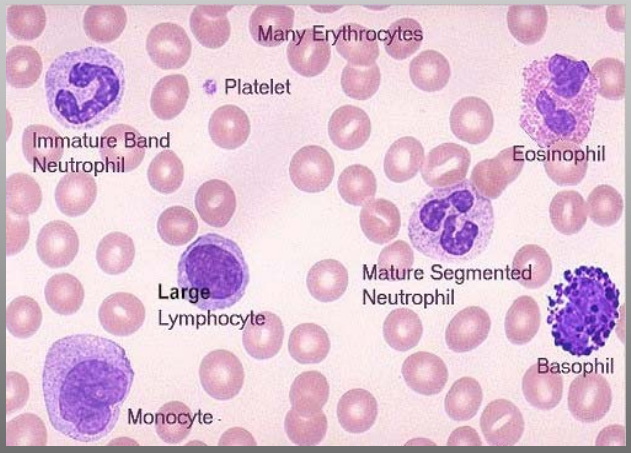
Phagocytosis of IgG-coated Targets by Macrophages



3 min

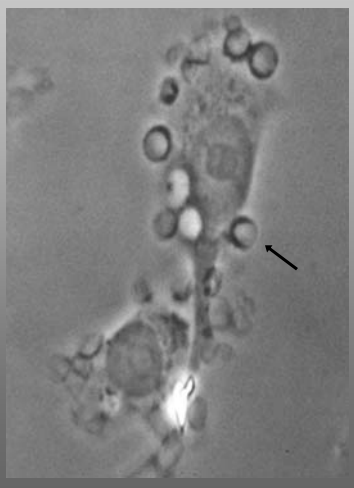


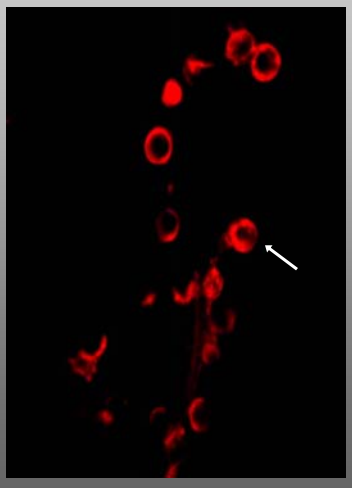
10 min



Mast Cells Can Phagocytose Too!

Extension of an F-actin-rich "Phagocytic Cup" Around Phagocytic Targets

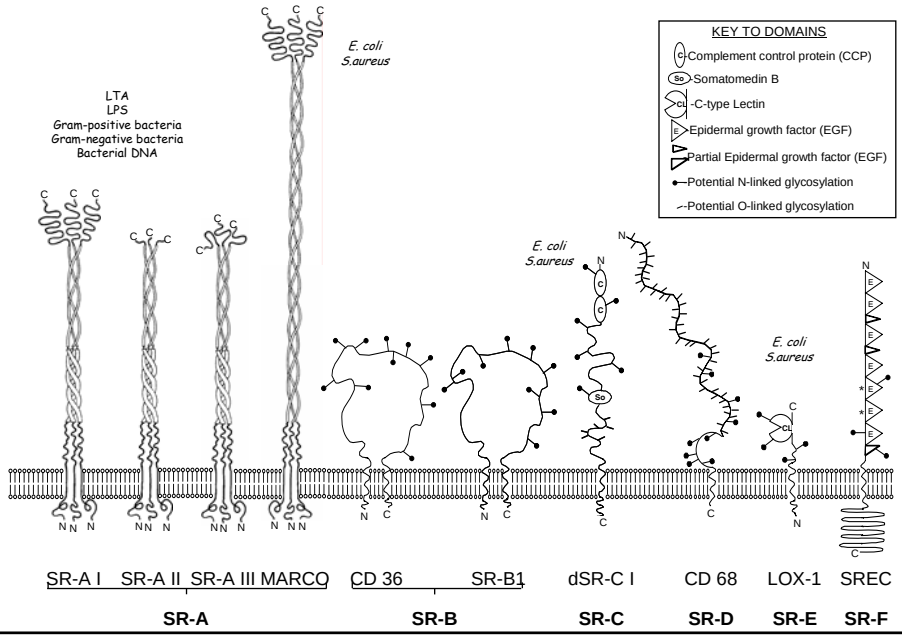




Examples of "Pattern Recognition Receptors" that Participate in Phagocytosis

Receptor	Expression	Target	Ligand
Integrins			
CR3 (CD11b/CD18; $\alpha_M\beta_2$)	PMN, Mo, Mϕ	Yeast	β-glucan
			C3bi, fibrinogen, LPS, ICAM
β_1 Integrins	Leuk	*Yersinia*	Invasin
Scavenger Receptors			
SR-AI/SR-AII	Mϕ	Gram-positive bacteria	Leipoteichoic acid
		Gram-negative bacteria	?
MARCO	Mϕ	*E. coli, S. aureus*	?
Lectins			
Dectin-1	Mϕ, DC	Yeast	β-glucan
CR3 (CD11b/CD18; $\alpha_M\beta_2$)	PMN, Mo, Mϕ	Yeast	β-glucan

The Scavenger Receptor Superfamily



Elie Metchnikoff, 1845-1916

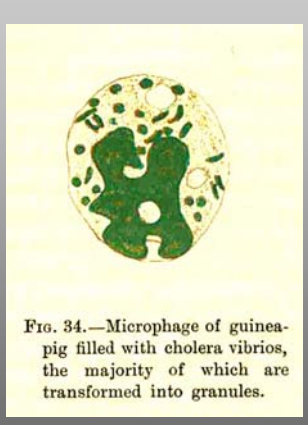
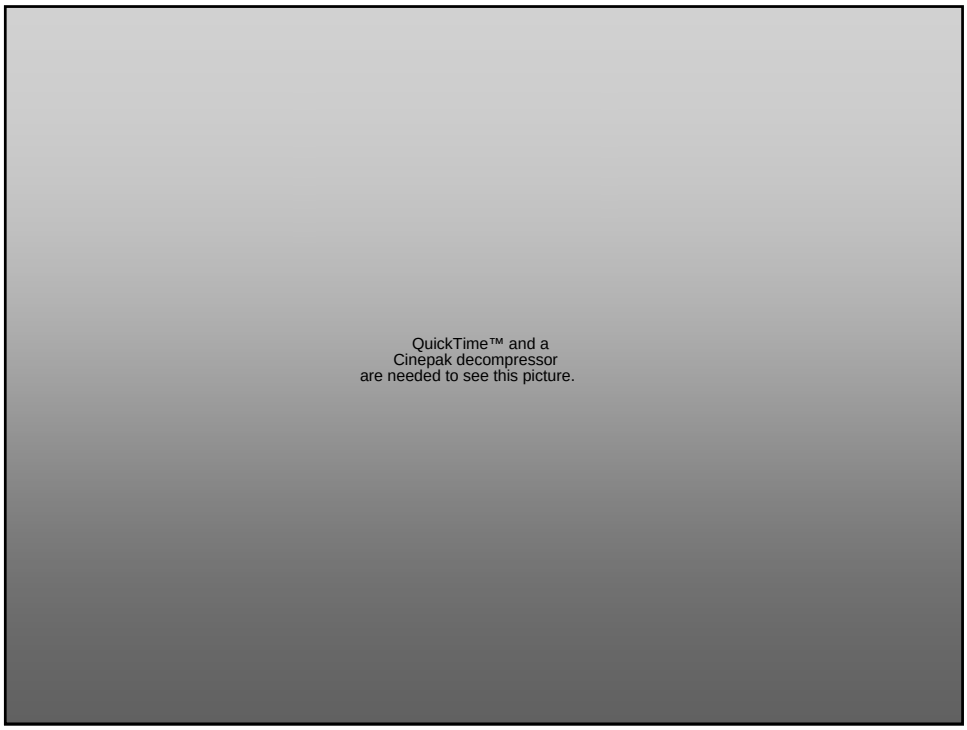


FIG. 34.—Microphage of guinea-pig filled with cholera vibrios, the majority of which are transformed into granules.

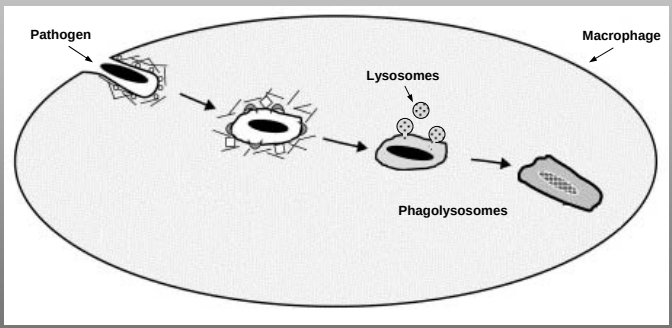
Phagosome-Lysosome Fusion?

QuickTime™ and a
Cinepak decompressor
are needed to see this picture.

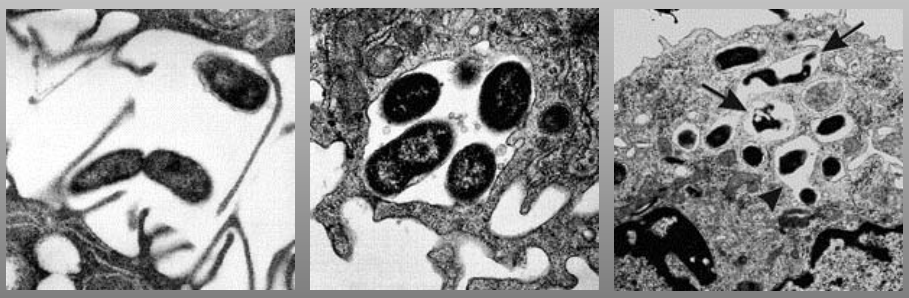

QuickTime™ and a
Cinepak decompressor
are needed to see this picture.


QuickTime™ and a
Cinepak decompressor
are needed to see this picture.

Post-phagocytic Events: Phagosome-Lysosome Fusion



Phagocytosis of Bacteria is Followed by Phagosome-Lysosome Fusion

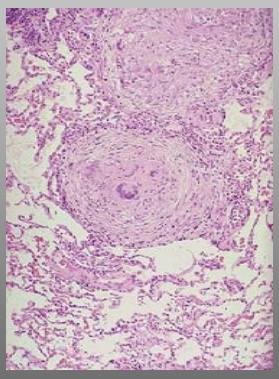


0-3 min | 1-5 min | 30 min-hrs

From: Allen et al., *J. Exp. Med.* 191:115, 2000

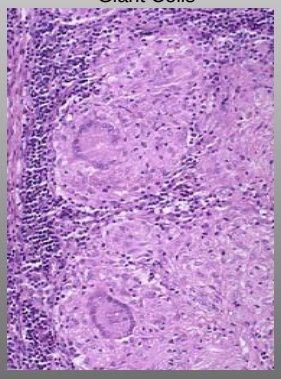
The Granuloma: a Delayed Response to Indigestible Pathogens and Particles in Macrophages

Granulomas



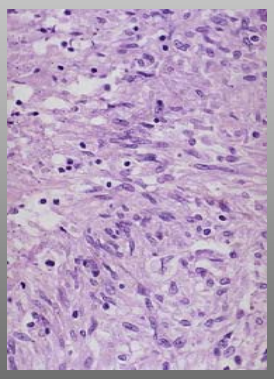
Granulomatous inflammation consists of epithelioid macrophages, giant cells, lymphocytes, plasma cells, and fibroblasts.

Langhans-type Giant Cells



Langhans-type giant cells represent fused macrophages. The nuclei are lined up around the peripheryof the cell.

Epithelioid Cells

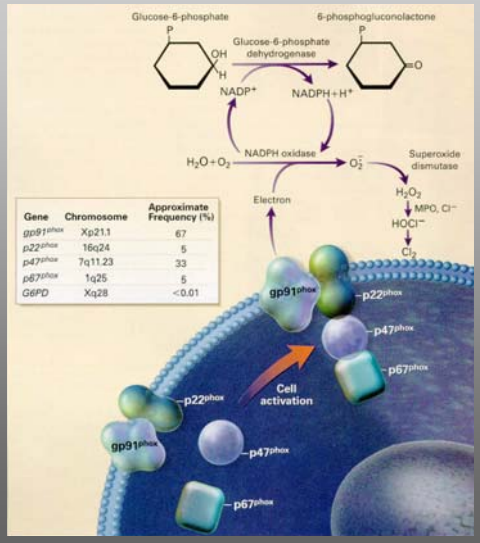


Epithelioid cells accumulate around the center of a granuloma. They get their name from the fact that they have pink cytoplasm similar to squamous epithelia.

Oxidant-dependent Killing of Bacteria and Fungi

QuickTime™ and a
Cinepak decompressor
are needed to see this picture.

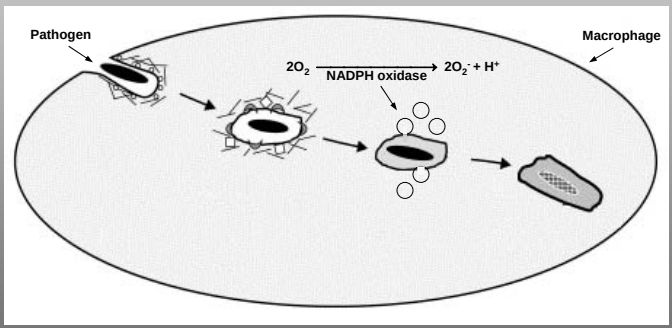
Chronic Granulomatis Disease, an Inherited Defect of the NADPH Oxidase Complex



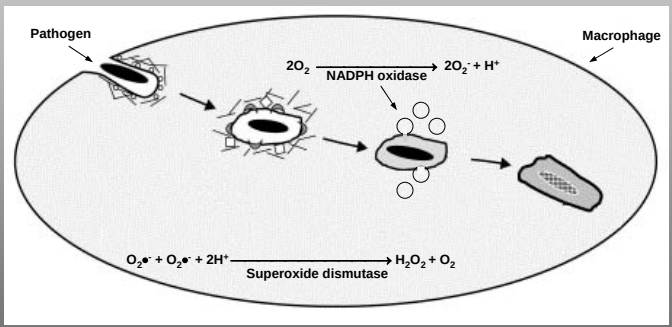
Gene	Chromosome	Approximate Frequency (%)
gp91phox	Xp21.1	67
p22phox	16q24	5
p47phox	7q11.23	33
p67phox	1q25	5
G6PD	Xq28	<0.01

From: Lekstrom-Himes and Gallin, *N Engl J Med*, 343:1703, 2000

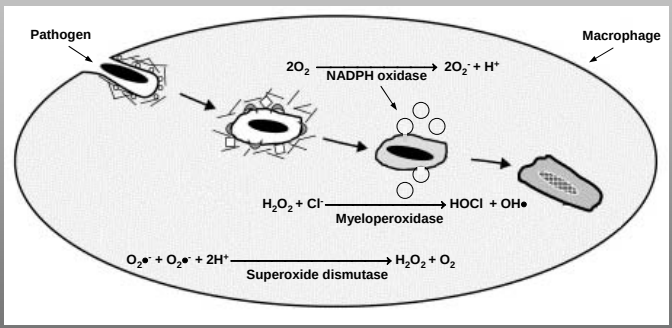
Post-phagocytic Events: "Phagosome-Oxidase Fusion"



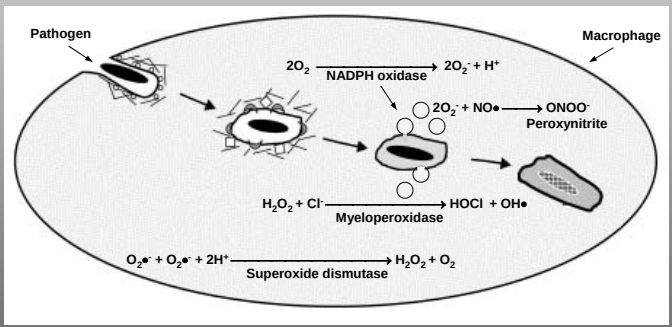
Post-phagocytic Events: Generation of H_2O_2



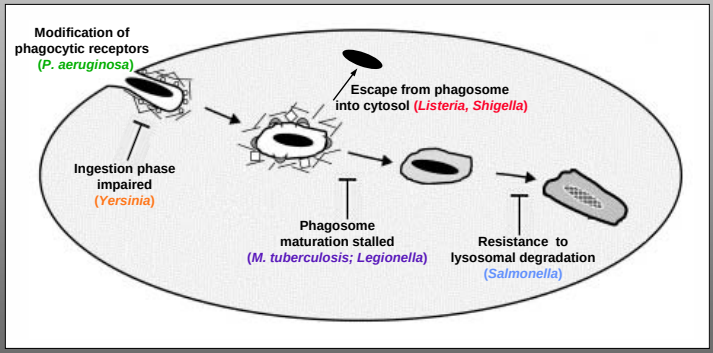
Post-phagocytic Events: Myeloperoxidase Activity



Post-phagocytic Events: Peroxynitrite Production



Bacterial Virulence Factors Subvert Host Defenses



Non-oxidative Killing Mechanisms

QuickTime™ and a
TIFF (Uncompressed) decompressor
are needed to see this picture.

Epithelial Cells Express Defensins, Too

HD-5

HBD-1

HBD-2

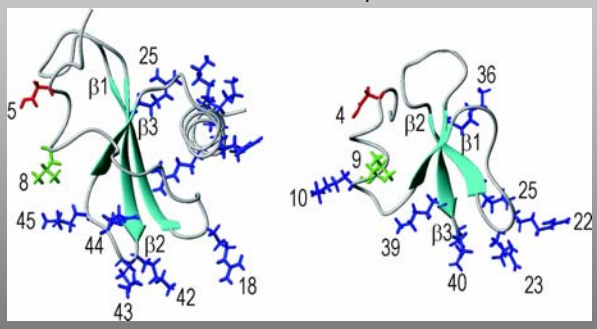
HBD-2 *H. pylori*, untreated

HBD-2 *H. pylori*, treated

From: Wehkamp et al., *J. Clin. Path.*56:352, 2003; Hamanka et al., *Gut* 49:481, 2001

Structural Similarity of a Chemokine and a Defensin

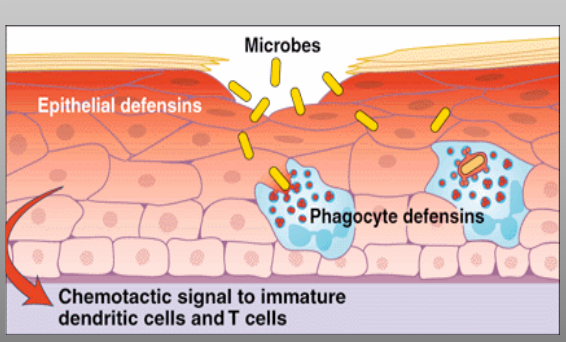
Chemokine CCL20 β-Defensin BD2



β-pleated sheets are represented by ***green arrows***;
arginine and lysine residues are shown in ***blue***

From: Perez-Canadillas et al., *J. Biol. Chem.* 2001 276:28372

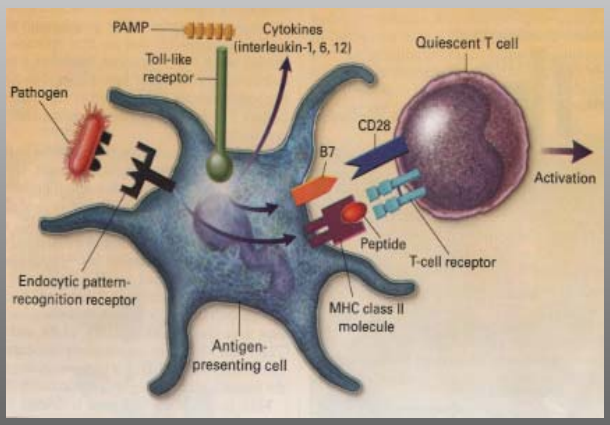
The Role of Defensins in Orchestrating the Immune Response



Inflamed defenders. A model of defensin activity in an infected epithelium. Epithelial cells synthesize antimicrobial defensins (red) both constitutively and in response to infectious and inflammatory stimuli. Other defensins are introduced by the influx of phagocytic cells that use them to kill ingested microbes. Released defensins attract dendritic cells and memory T cells, setting the stage for the adaptive phase of the immune response. From Ganz, *Science* 286:420, 1999.

The Relationship Between the Innate Immune Response and Acquired Immunity

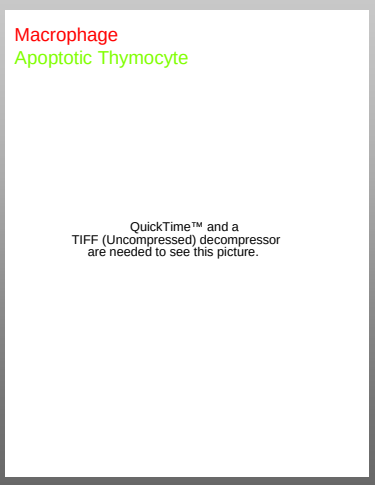
The (Primary) Acquired Immune Response is Initiated by Innate Immune Recognition



Phagocytosis: Not Just for Bugs

Phagocytosis is the Principal Mechanism of Disposal of Apoptotic Corpses

Macrophage
Apoptotic Thymocyte



Implications: Disposal of apoptotic corpses occurs continuously during the lifetime of an individual. In this setting, phagocytosis is not accompanied by inflammation, but rather by an anti-inflammatory signal (the production of TGF-β). As apoptotic corpses contain many potential self antigens, the lack of an appropriate anti-inflammatory signal has the potential to trigger autoimmunity.

From: Jennings et al., *Am. J. Resp. Cell Mol. Biol.* 32:108, 2005

Immunological Consequences of Phagocytosis

Clearance of pathogens

Death of pathogenic microbe
Resolution of infection

Persistence of pathogenic microbe
Failure of resolution of infection

Clearance of apoptotic corpses

Suppression of inflammation
Tolerance

Inappropriate inflammation
Break in tolerance

Summary

1. Innate immunity represents the first-line of host defense. Its receptors are germline-encoded and recognize pathogen-associated “molecular patterns.”

2. Phagocytosis is a component of innate and aquired immunity. It is the principal means of destroying pathogenic bacteria and fungi. Phagocytosis initiates the process of antigen presentation.

3. Many phagocytic receptors recognize a diverse array of microbial pathogens. Some pathogens (e.g., *S. pneumoniae*) require opsonization for their clearance. However, bugs fight back.

4. Phagocytic leukocytes employ oxidative and non-oxidative means of killing. The NADPH oxidase generates reactive oxidants, such as superoxide anion and hypochlorous acid (bleach).

5. Innate immunity ushers in acquired immunity: innate immune activation of APCs results in up-regulation of co-stimulatory molecules and enhances the effectiveness of antigen presentation.

6. Phagocytosis is an essential component of development and tissue remodelling. Ingestion of apoptotic bodies is immunologically “silent” and is normally accompanied by a suppression of inflammation. Failure of this mechanism may result in autoimmunity.