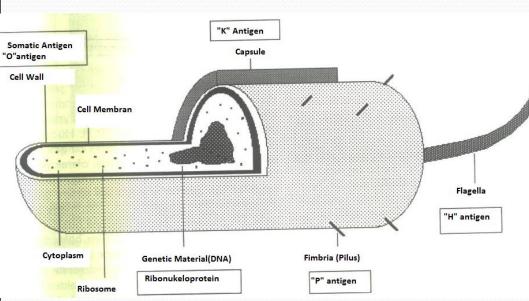
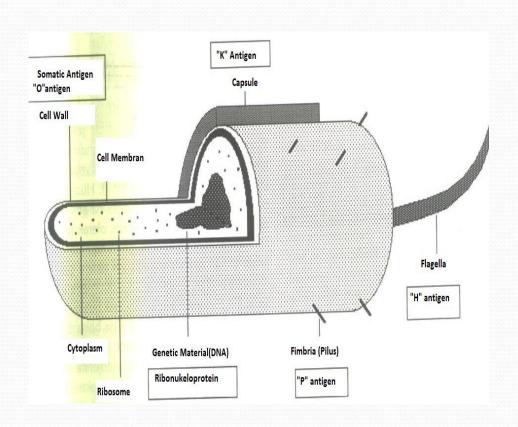


- Cell wall Somatic antigen (O antigen):
- -Gr (+) peptidoglycan teicoic acid and lipoteicoic acid
- -Gr (-) lipopolysaccharide and porine proteins
- Morphological structure, substance exchange, antigenic structure, adhesive molecule content, protection from environmental and host defense factors
- Antibodies to cell wall antigens;
 Opsonization, agglutination,
 neutralization



- Capsule (K antigen):
- -polysaccharide (B.anthracis protein)
- protection from environmental factors and host immune system
- -antifagocytic property
- Anti-capsular immune response; the opsonization

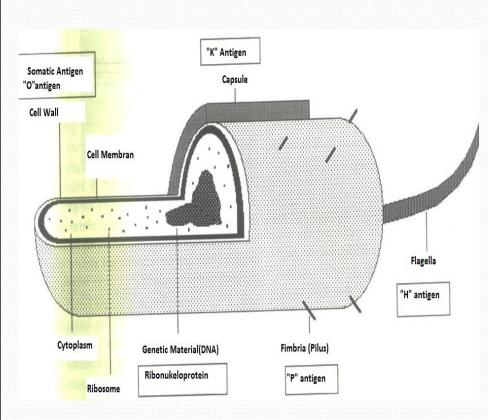


Flagella (H antigen)

- -flagellin (protein)
- movement organelle
- -anti-flagellar immune response; agglutination

Fimbria-Pilus (Pantigen)

- -piline (protein)
- -adhesion organelle
- -anti-fimbrial immune response; neutralization



exotoxins;

- -protein
- lysis of tissue cells (neurotoxin, leukotoxin, hepatotoxin)
- -antitoxin immune response; neutralization

Internal structures; intrastoplasmic organelles Heat shock proteins (stress proteins)

Pathogenesis of bacterial infections

Exotoxigenic mechanism;

Exotoxins break cells, disrupt cell functions, stimulate excessive amounts of harmful cytokines.

Endotoxogenic mechanism;

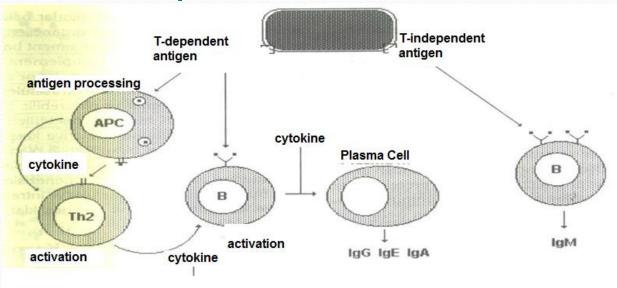
- initiates cytokine synthesis by stimulating inflammation cells (neutrophils, macrophages, endothelial cells), causing inflammation-shock-depression-fever

Invasive mechanism;

- They emit strong enzymes (hyaluronidase, collagenase, coagulase, etc.) that spread in tissues and disrupt the structure and function of tissues or live in cells (intracellular bacteria).

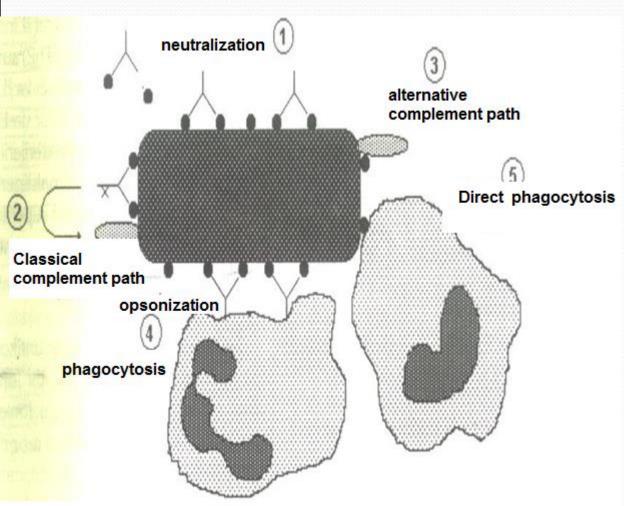
NATURAL DEFENSE MECHANISMS

- Genetic factors
- Hormones (low-dose steroid and estrogen: immunostimulatory effect - high-dose steroid, testesterone and progesterone: immunosuppressive effect)
- Nutrition
- lysozyme
- Free fatty acids (oleic acid)
- Antibacterial peptides (beta lysine, defensin, spermine)
- Iron binding proteins (lactoferrin, transferrin, ferritin)



- Humoral immune response is effective.
- It is part of the bacterial cell wall and the capsule-polysaccharidestructured antigen and a T-independent immune response occurs.
- Bacterial cell wall (peptidoglycan, porine, glycoprotein, etc.), flagella and fimbria - protein-antigen and T-dependent immune response occurs.

Functions of Antibodies to Extracellular Bacteria



Complement activation (IgG and IgM)

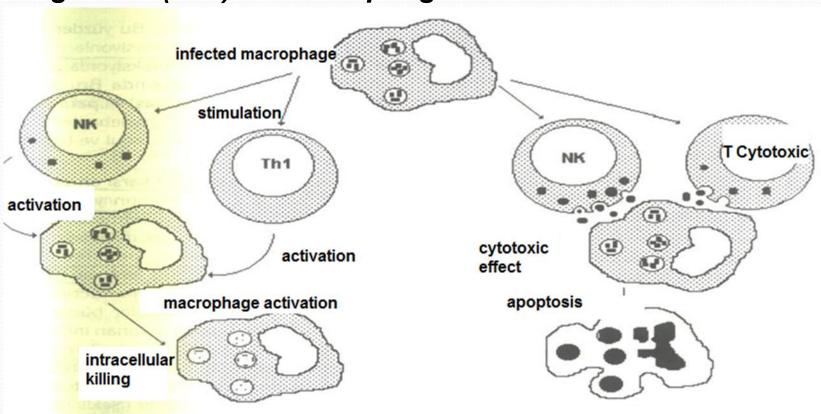
Opsonization (IgG and IgM)

Neutralization (IgA, IgG and IgM)

Inflammation Stimulation

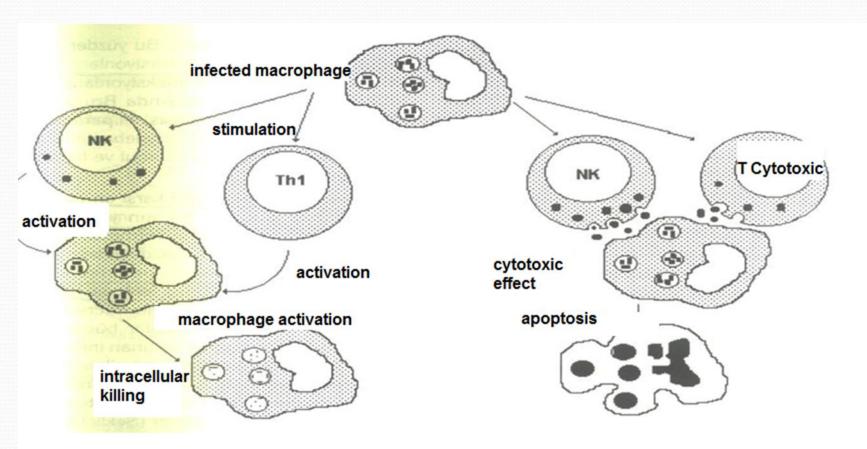
- These bacteria are more resistant than others,
- They can survive for a long time in the cells to which they are phagocytized,
- They cause chronic infections,
- Cellular immunity is effective,
- Three basic mechanisms are effective in cellular immunity

 Macrophage activation: IL12 (infected macrophage) and IFN gamma (NK cell) → Th1 stimulation → IL2 and IFN gamma (Th1) → macrophage activation

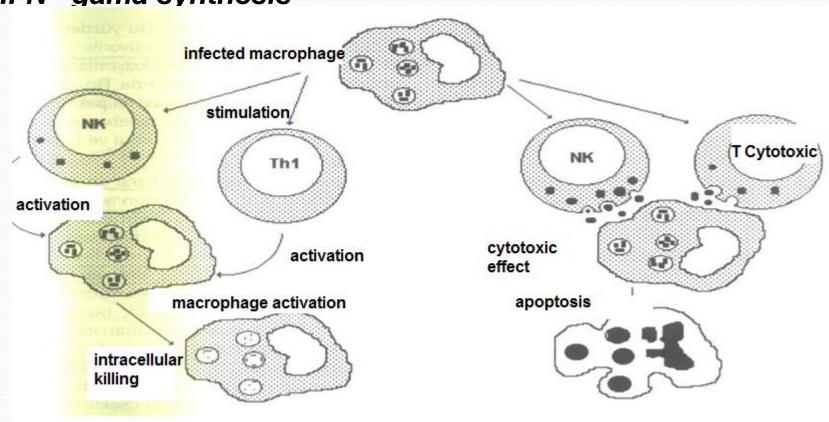


Live Bacteria **Dead Bacteria** Macrophage Phagocytosis Macrophage Invasion IL-12 Th2 Cell Th1-Cell IL4-IL5 IFN-gamma **Antibody Production Macrophage Activation**

Cytotoxic T-lymphocyte: Intrastoplasmic protein antigens
 → presentation with MHC class I → cytotoxic T-lymphocyte stimulation → apoptosis



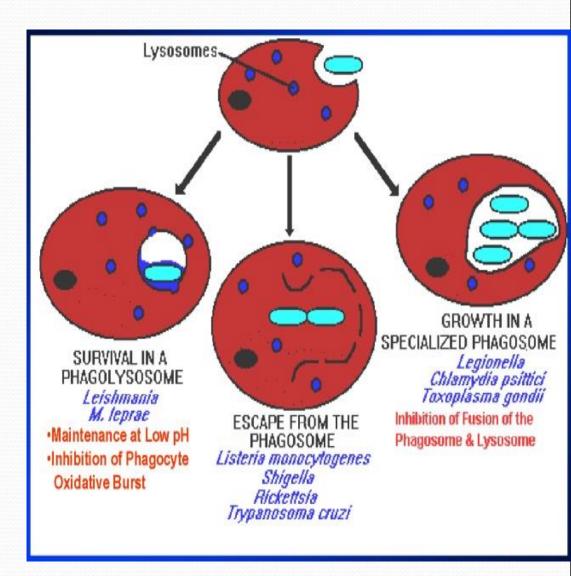
 NK cells: IL12 (infected macrophage) → NK cell stimulation → macrophage activation by apoptosis and IFN –gama synthesis



Ways of Bacteria to Avoid Immune Response

Resistance to phagocytosis

- Resistance to ingestion (capsule)
- Live on the phagolysosome stay
- Survival of cell stoplasm by escape from phagosome
- Phagolysosome formation blocking and phagosome development



Ways of Bacteria to Avoid Immune Response

- Resistance to complement effect (capsule structure)
- Destruction of immune system molecules (IgA and IL-2 degrading enzymes)
- Antigenic variation