

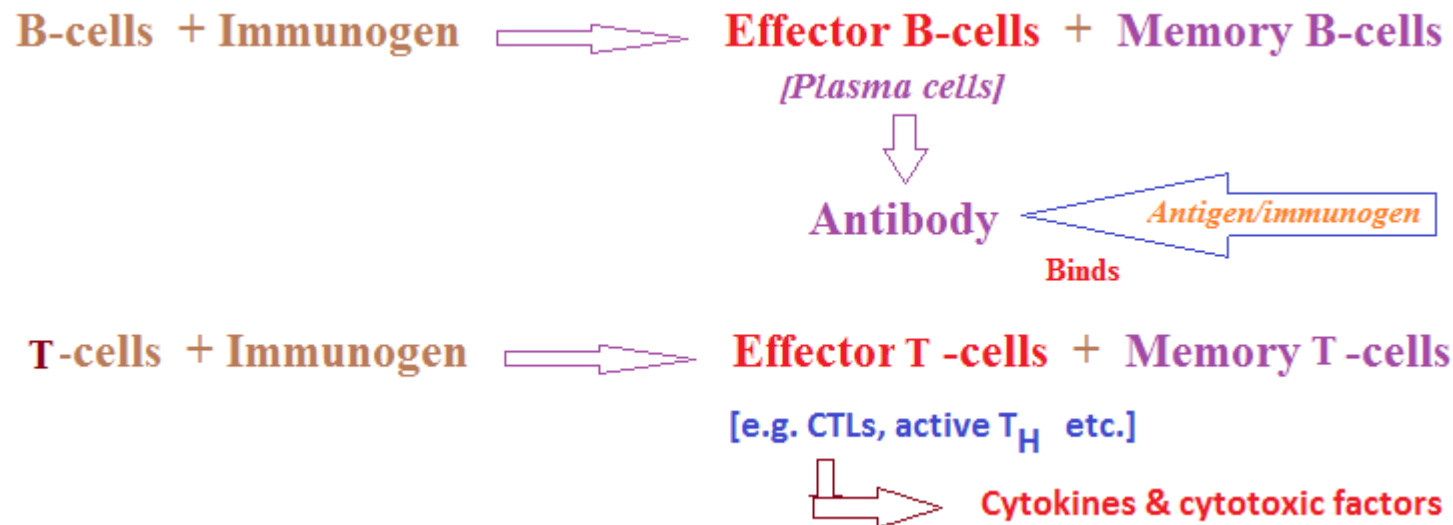
# Antigens

To 'B' or not to 'B'- that is the question

An immunogen refers to a molecule that is capable of eliciting an immune response (humoral and/or cell-mediated) by an organism's immune system, whereas an antigen refers to a molecule that is capable of binding to the product of that immune response.

So, an immunogen is necessarily an antigen, but an antigen may not necessarily be an immunogen.

Hence , all immunogens are antigen but all antigens are not immunogen.



**Haptens are low-molecular-weight compounds that may be bound by antibodies or T cell receptors, but cannot elicit an immune response by themselves.**

**Haptens themselves are nonimmunogenic and they cannot evoke an immune response until they are conjugated with a larger carrier immunogenic molecule such as BSA. The hapten-carrier complex, unlike free hapten, can act as an immunogen and can induce an immune response.**

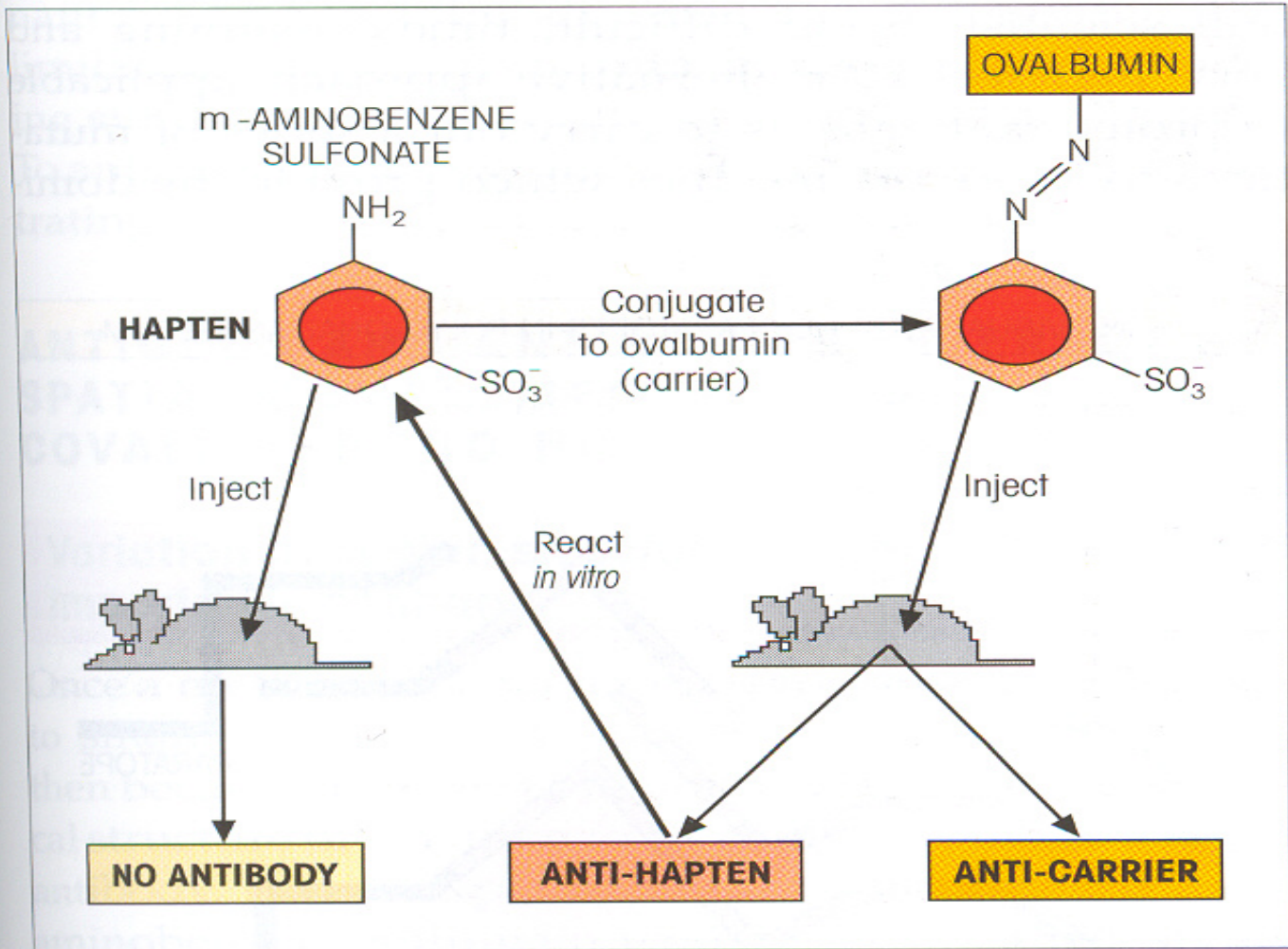
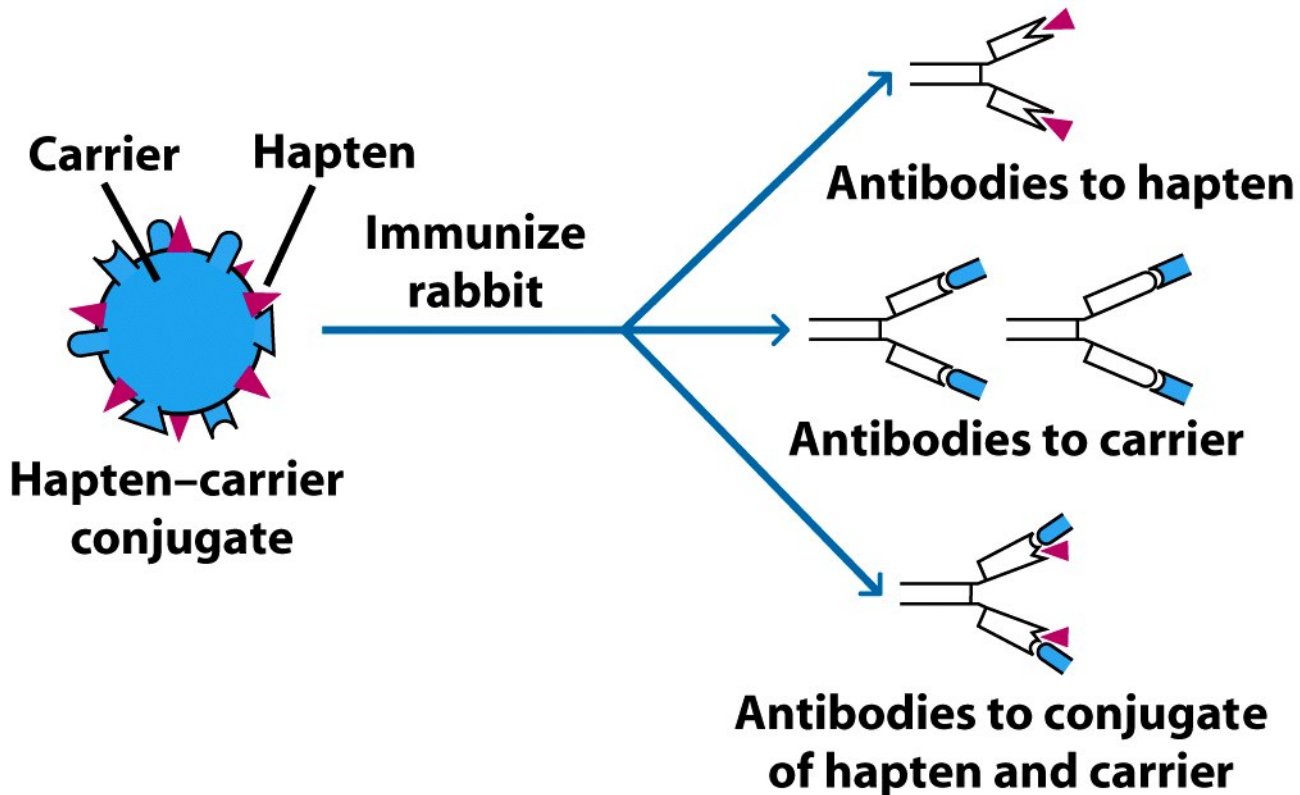



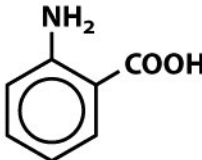
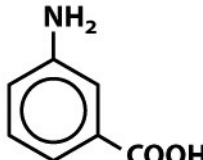
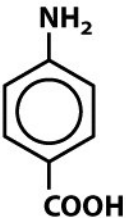
Figure 5.1. A hapten on its own will not induce antibodies. However, it will react *in vitro* with antibodies formed to a conjugate with an immunogenic carrier.



| <b>Injection with:</b>                    | <b>Antibodies formed:</b>   |
|---|---|
| <b>Hapten (DNP)</b>                       | <b>None</b>   |
| <b>Protein carrier (BSA)</b>              | <b>Anti-BSA</b>   |
| <b>Hapten-carrier conjugate (DNP-BSA)</b> | <b>Anti-DNP (major)</b><br><b>Anti-BSA (minor)</b><br><b>Anti-DNP/BSA (minor)</b> |

**Figure 4-1**  
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**TABLE 4-1**
**Reactivity of antisera with various haptens**

| Antiserum against           | REACTIVITY WITH   |  |   |   |
|-----------------------------|---|--|---|---|
|                             |  |  |  |  |
|                             | Aminobenzene (aniline)  | <i>o</i> -Aminobenzoic acid  | <i>m</i> -Aminobenzoic acid   | <i>p</i> -Aminobenzoic acid   |
| Aminobenzene                | +   | 0  | 0   | 0   |
| <i>o</i> -Aminobenzoic acid | 0   | +  | 0   | 0   |
| <i>m</i> -Aminobenzoic acid | 0   | 0  | +   | 0   |
| <i>p</i> -Aminobenzoic acid | 0   | 0  | 0   | +   |

KEY: 0 = no reactivity; + = strong reactivity

SOURCE: Based on K. Landsteiner, 1962, *The Specificity of Serologic Reactions*, Dover Press. Modified by J. Klein, 1982, *Immunology: The Science of Self-Nonself Discrimination*, Wiley.

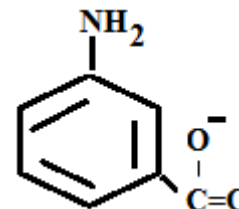
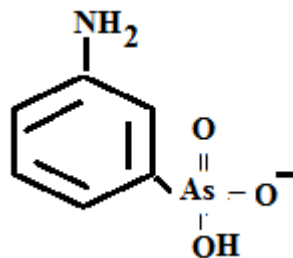
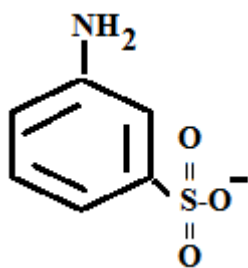
**Table 4-1**
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**Cross reactivity between antisera against m-amino benzene sulfonate and *o*-, *m*- and *p*- amino benzene sulfonate/arsonate /carbonate.**

|                         | Ortho- | Meta- | Para- |
|-------------------------|--------|-------|-------|
| Amino benzene sulfonate | +      | +++   | -     |
| Amino benzene arsonate  | -      | +     | -     |
| Amino benzene carbonate | -      | -     | -     |

**Explain these results.**



•**Epitope: the portion of an antigen that is recognized and bound by an Ab or TCR/MHC complex (antigenic determinant).**

•**Paratope: “The site in the variable (V) domain of an antibody or T-cell receptor that binds to an epitope on an antigen**

# Epitopes

- Antigenic determinants recognized by B cells and T cells
  - B cell epitopes tend to be on the outside of the antigen
    - For example, the hydrophilic amino acids on a protein's surface
  - T cell epitopes from proteins derived from enzymatic digestion of peptide and then association with MHC

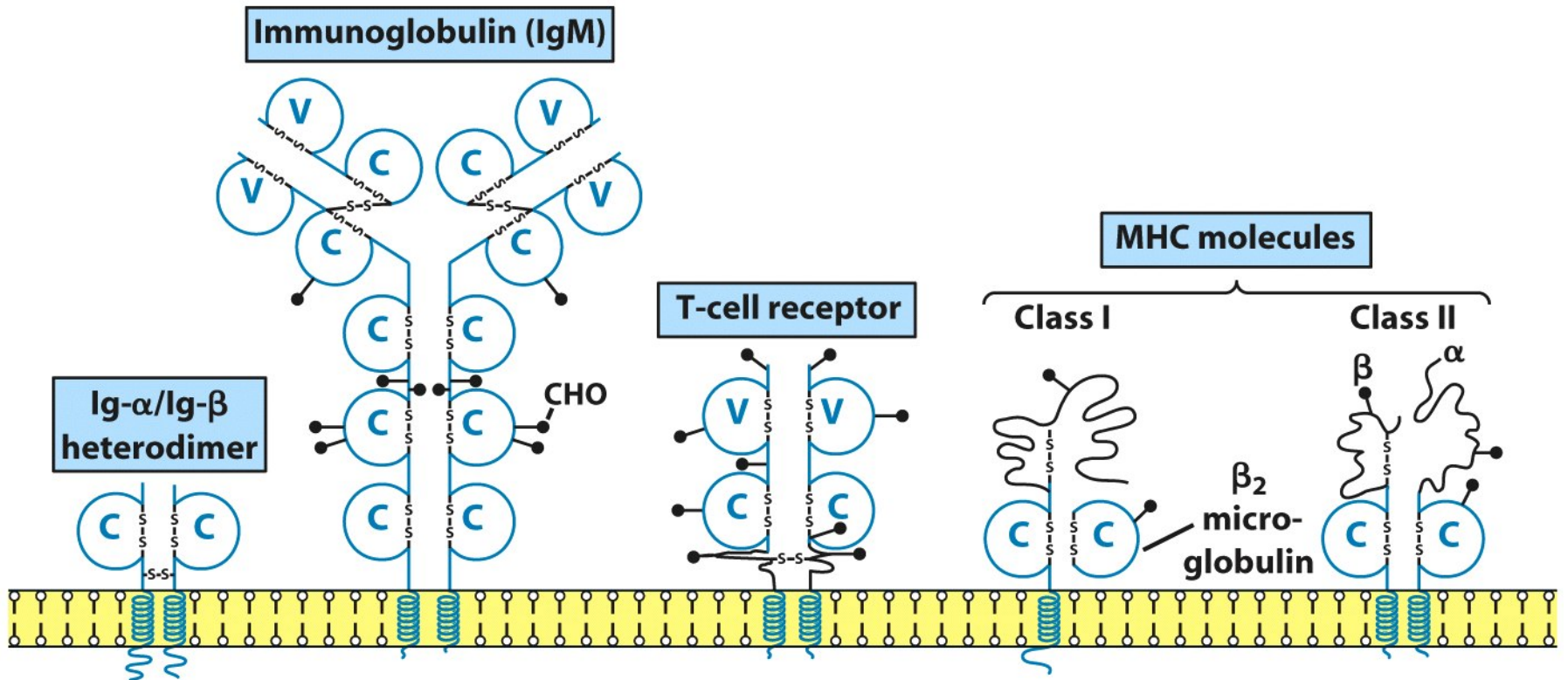


# Receptor-Ligand Interactions

- Antigen receptors of the adaptive immune system are transmembrane proteins
  - B cells – the B cell receptor
  - T cells – the T cell receptor
- Multiple noncovalent bonds
  - Hydrogen bonds
  - Ionic bonds
  - Van der Waals
  - Hydrophobic interactions

# Immunoglobulin Superfamily

- All have similar structures
- Examples:
  - Antibodies
  - T-cell receptors
  - Class I and II MHC molecules
  - Part of B cell receptor
- Most members of immunoglobulin superfamily cannot bind antigen



**Figure 4-24 part 1**  
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## Characteristics of B-cell epitopes

- B cell epitopes on native proteins generally are composed of hydrophilic amino acids on the protein surface that are topographically accessible to membrane bound or free antibody.
  - When talking about proteins, the epitopes can be **sequential or nonsequential** (referring to amino acid sequence) depending on protein folding

**TABLE 4-2** Comparison of antigen recognition by T cells and B cells

| <b>Characteristic</b>               | <b>B cells</b>  | <b>T cells</b>  |
|-------------------------------------|---|---|
| <b>Interaction with antigen</b>     | Involves binary complex of membrane Ig and Ag   | Involves ternary complex of T-cell receptor, Ag, and MHC molecule                     |
| <b>Binding of soluble antigen</b>   | Yes   | No  |
| <b>Involvement of MHC molecules</b> | None required   | Required to display processed antigen   |
| <b>Chemical nature of antigens</b>  | Protein, polysaccharide, lipid  | Mostly proteins, but some lipids and glycolipids presented on MHC-like molecules      |
| <b>Epitope properties</b>           | Accessible, hydrophilic, mobile peptides containing sequential or nonsequential amino acids | Internal linear peptides produced by processing of antigen and bound to MHC molecules |

Table 4-2  
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**Naïve T cells: cells never meet antigens before.**

They can only be activated by dendritic cells.

- **Effector cells: short-lived cells with special functions** such as cytokine secretion and B-cell help and cytotoxic killing activity. Effector cells are derived from naïve or memory cells after antigen activation. TH1 and TH2 subsets.
- **Memory cells: long-lived resting cells that are** derived from naïve and effector cells. They respond faster and stronger to a subsequent challenge with the same antigen.

# Properties of Immunogen contribute to Immunogenicity

- Properties
  - **Foreignness**
  - **Molecular size**
  - **Chemical composition and complexity**
  - **Ability to be processed and presented on MHC**

- Foreignness

- Lymphocytes that do not bind to self antigens are allowed to further develop

- Therefore they will later only recognize nonself antigens

- For example:

- Bovine serum albumin (BSA) is not immunogenic when injected into cow but is when injected into chicken

- Some macromolecules are highly conserved throughout evolution and display little immunogenicity

- » Cytochrome c, collagen



- Molecular Size
  - Active (good) immunogens
    - » > 100,000 Daltons
  - Poor immunogens
    - » < 5,000-10,000 Daltons

- Chemical Composition
  - Polymers composed of multiple copies of same amino acid or sugar tend to be poor immunogens
  - Lipids are haptens and need to be conjugated with carrier to produce antibodies
    - Important for assays for detection of some steroids, vitamins

- Susceptibility to antigen processing
  - Large, insoluble macromolecules are more likely to be phagocytized for processing

# The biological system contributes to immunogenicity

- Host Genetic make-up
- Manner in which material is presented
- Use of agents (adjuvants) to enhance immunogenicity

- Genotype of recipient animal
  - Genes of MHC
  - Genes in coding for specific antibodies

# FACTORS GOVERNING IMMUNOGENICITY

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## Properties of the Host

### Genetics of the responding host

#### 1. Major Histocompatibility Complex (MHC)

Immune response genes

IA and IE: Class II in Mouse

ImgenFx2.pox

Called “H2” in Mice; “HLA” in Human

HLA = Human Leucocyte-associated Antigens

TABLE 4.2

# Effect of MHC haplotype on the immune response to the Two different antigens in mice

Ag&MHCd.ppt  
Kuby 2nd Ed (4th Build)

| MHC Haplotype    | Representative mouse strains | Antibody response to Ag 1 | Antibody response to Ag 2 |
|------------------|------------------------------|---------------------------|---------------------------|
| H-2 <sup>b</sup> | C57                          | Low                       | High                      |
| H-2 <sup>b</sup> | C57BL/6                      | Low                       | High                      |
| H-2 <sup>b</sup> | C3H.SW                       | Low                       | High                      |
| H-2 <sup>b</sup> | I29/J                        | Low                       | High                      |
| H-2 <sup>d</sup> | BALB/c                       | Intermediate              | Intermediate              |
| H-2 <sup>d</sup> | B10.D2                       | Intermediate              | Intermediate              |
| H-2 <sup>d</sup> | DBA/2                        | Intermediate              | Intermediate              |
| H-2 <sup>d</sup> | NZB                          | Intermediate              | Intermediate              |
| H-2 <sup>k</sup> | CBA                          | High                      | Low                       |
| H-2 <sup>k</sup> | C3H/HeJ                      | High                      | Low                       |
| H-2 <sup>k</sup> | C58J                         | High                      | Low                       |
| H-2 <sup>k</sup> | B10.BR                       | High                      | Low                       |
| H-2              | B10.S                        | Low                       | Low                       |
| H-2              | SJL                          | Low                       | Low                       |

# FACTORS GOVERNING IMMUNOGENICITY

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## Properties of the Host

### **Genetics of the responding host**

2. Variable region (V-region)  
genes available for Antibody  
and T-cell Receptor generation
3. Genes controlling factors for  
Immune regulation



# Immunogen dosage and route of Administration

- Too low or high of dosage can induce tolerance (WHY?)
- Single dose is often not enough – booster is needed
- Route
  - Intravenous (iv)
  - Intradermal (id)
  - Subcutaneous (sc)
  - Intramuscular (im)
  - Intraperitoneal (ip)
    - » Antigen administered iv would travel to spleen; administered sc would travel to lymph nodes

# **Conformational Properties of Epitopes for Antibodies**

## ADJUVANTS

An **adjuvant** (from **Latin adjuvare – to help**) is any substance, distinct from antigen, which enhances immune response by various mechanisms: by recruiting of professional Antigen-Presenting Cells (APCs) to the site of antigen exposure, by increasing the delivery of antigens by its delayed/slow release (depot generation), immunomodulation by cytokine production (selection of Th1 or Th2 response), by induction of T-cell response (prolonged exposure of peptide-MHC complexes (signal 1) and stimulation of expression of T-cell-activating co-stimulators (signal 2) on the APC's surface).

**Freund's adjuvant** (**toxic for human use, used only for mice , rabbit etc.**)

Freund's complete adjuvant – antigen in aqueous solution, mineral oil, emulsifying agent (mannide monooleate), heat killed cells of Mycobacteria (muramyl di-peptide activates dendritic cells and macrophages)

**Freund's incomplete adjuvant – antigen in aqueous solution, mineral oil, and emulsifying agent.**

Aluminum potassium sulfate (**Alum**) → **Approved for general human use**

- Adjuvants

Enhance immunogenicity by:

- Prolong antigen persistence (Antigen is released very slowly from injection site)
- Enhance co-stimulatory signals (B7 etc.)
- Induce granuloma formation (Chronic inflammatory response attract both phagocytes and lymphocytes formation of dense, macrophage-rich mass)
- Stimulate lymphocyte proliferation non-specifically

# **Activation of Systemic Inflammatory Responses by Potent Inflammatory Signals from Infectious Organisms and Plants**

**Endotoxins**

**Exotoxins**

**Super-antigens**

**Mitogens and Lectins**

# Immunopathology of Bacterial Septic Shock

Caused by Gram Negative Bacterial Endotoxins:

Examples -

E. coli

Pseudomonas aeruginosa

Neisseria meningitidis

Meningococcus

70,000 Deaths per year

Diarrhea, Fever, Blood Clotting, Blood Pressure Drop

Macrophage Activation by Endotoxin:

Generates systemic cytokine production -

IL1 and TNF-alpha (Tumor necrosis factor)

# Immunopathology of Bacterial Toxic Shock

**Exotoxins Secreted or Presented Membrane Bound:**

**Act as "Superantigens"**

**Examples -**

**Staphylococcus aureus:**

**TSST1 (Toxic Shock Syndrome Toxin)**

**Streptococcus pyogenes - Rheumatic fever and shock**

**Staphylococcus enterotoxins: Food poisoning**

**Superantigens cross-link TCR with Antigen-presenting Cell**

**MHC Class II Proteins**

**Generates systemic cytokine production - IL1 and TNF-alpha  
(Tumor necrosis factor)**

Antigen-presenting cell

Generalized  
Pathological  
T-Cell Activation  
By Superantigens

Class II  
MHC  
Protein

Staphylococcal enterotoxins (food poisoning)  
Streptococcal pyrogenes exotoxins (rheumatic  
fever, toxic shock)

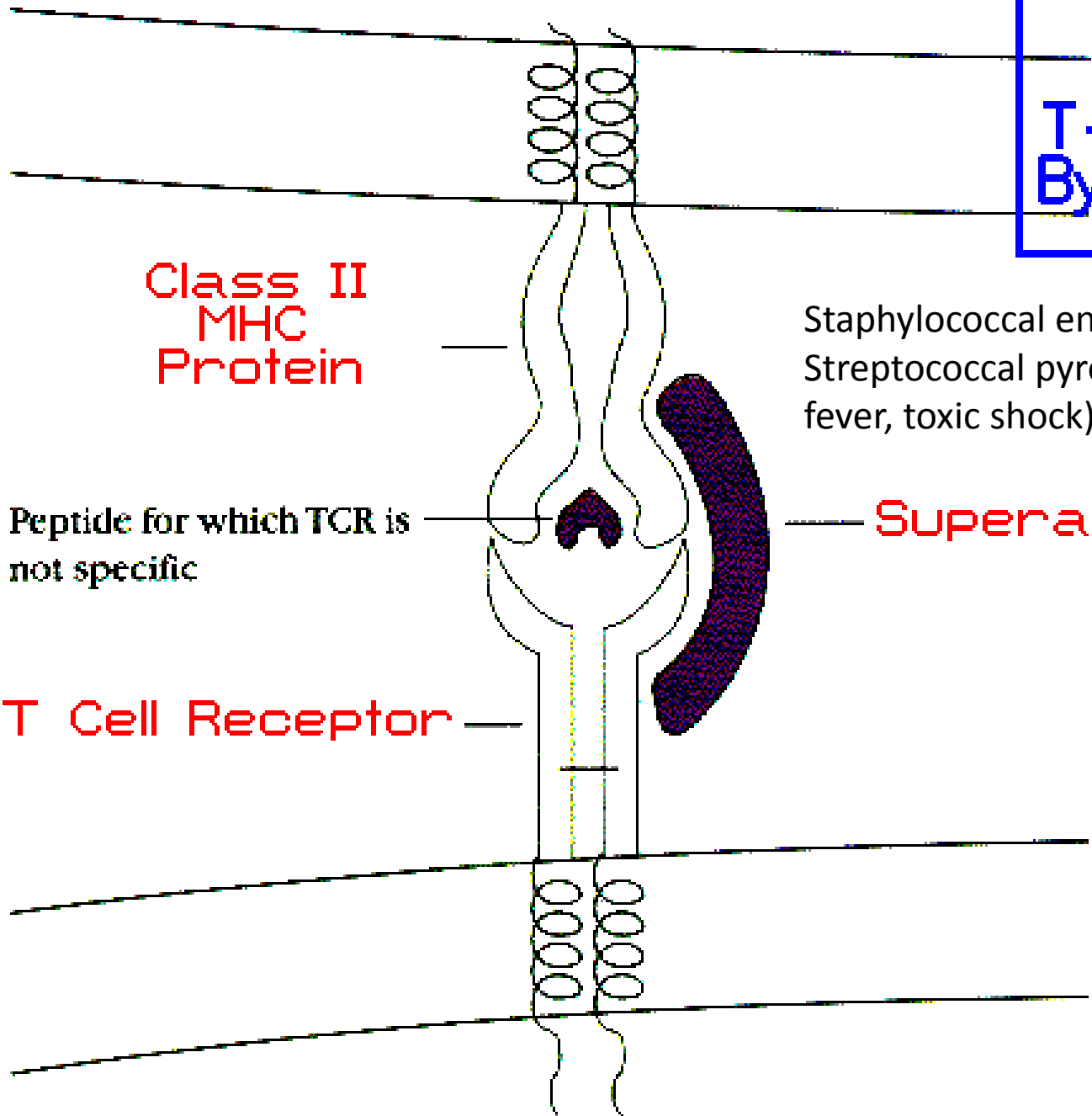
Peptide for which TCR is  
not specific

Superantigen

T Cell Receptor

SuperAg.pox  
Figure 4-15  
Kuby 2nd Ed

See Figure 11-6  
7<sup>th</sup> Edition p. 367





## Mitogens and Lectins

**Mitogens**: Stimulate mitosis and cell division non-specifically  
(non-specific therefore stimulate polyclonally)

**Lectins**: Proteins that bind to carbohydrate in glycoproteins  
(Can be potent mitogens)

**Glycoproteins**: Conjugated protein with covalently attached carbohydrate residues