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**Charting New Horizons in Education** 

Introduction to Immunity



Immunology

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## \*Learning objectives

• **1. Definitions and Characteristics** Antigen

Immunogen

Hapten Epitope

Adjuvant

- 2. Identify the factors that affect the immunogenicity
- 3. Understand the concept of antigen cross reactivity
- 4. Differentiate between active, passive, and adoptive immunity

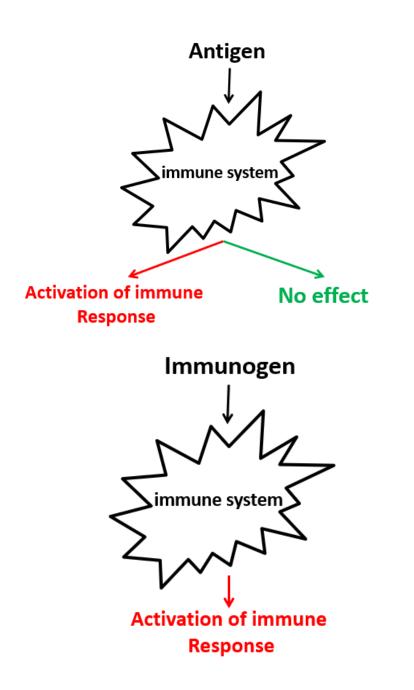
## **History of Immunology**

- The earliest known reference to immunity was during the plague of Athens in 430 BC
- In the 18th century, scientist made experiments with scorpion venom and observed that certain dogs and mice were immune to this venom.
- Immunology;
  - Study of the components and function of the immune system.
- Immune System;
  - Molecules, cells, tissues, and organs providing non-specific and specific protection against:
    - Microorganisms
    - Microbial toxins
    - Tumor cells

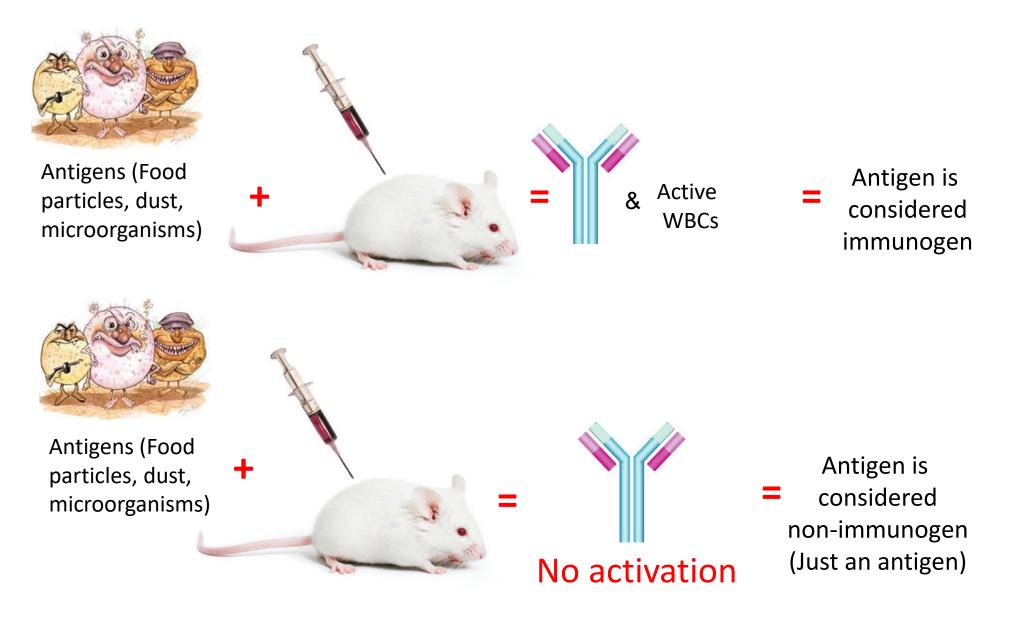


## Antigens & Immunogens

- Foreign substances that induce an immune response are of two types:
- a. Immunogen
  - A substance capable of inducing a specific immune response, resulting in the formation of antibodies or immune lymphocytes.
- b. Antigen
  - Any molecule that can bind to components of the immune system (lymphocytes, antibodies, and T cell receptors) without necessarily inducing an immune response.
- Immunogenic substances are always antigenic, whereas antigens are not necessarily immunogenic (e.g., autologous serum proteins).
- **Distinction;** The distinction between antigen and immunogen is functional.



## Antigens & Immunogens



## Antigens & Immunogens

Blood antigens in this case are considered Non-immunogens (Autologous donation) Blood group A Blood group B Blood antigens in this case are considered immunogens (Heterologous donation)

## \*Factor affecting Immunogenicity

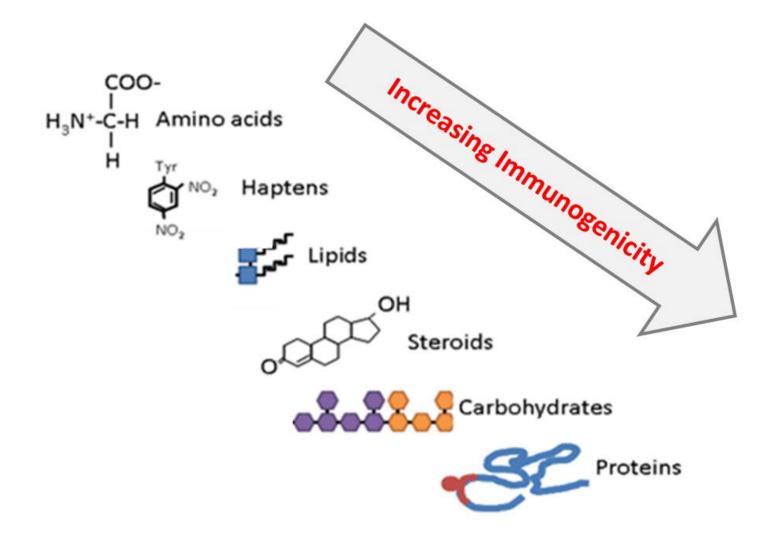
#### 1. Nature of the Immunogen;

- Foreignness: Animals typically do not respond to self-antigens. Compounds that are part of self are not immunogenic to the individual.
- Molecular Weight: A minimal molecular weight is required for immunogenicity.
  - <1000 daltons: Not immunogenic (e.g., penicillin, progesterone, aspirin).
  - 1000-6000 daltons: May or may not be immunogenic (e.g., insulin).
  - >6000 daltons: Are immunogenic (e.g., albumin, tetanus toxin).
- Chemical Structure Complexity: Heteropolymers containing two or more different amino acids are more immunogenic than homopolymers.

#### **2. Biological Factors**

- Dosage
- route of administration
- (Subcutaneous > Intravenous > Intragastric)
- individual genetic differences
- •the use of adjuvants.

## \*Factor affecting Immunogenicity



V:A

## **~**Adjuvants

#### 1. Definition:

- 1. Substances that enhance immunogenicity without altering chemical composition.
- 2. Substances which when mixed with an immunogen enhance the immune response against the **immunogen (Immunopotentiator or Immuno-booster)**.

#### 2. Examples:

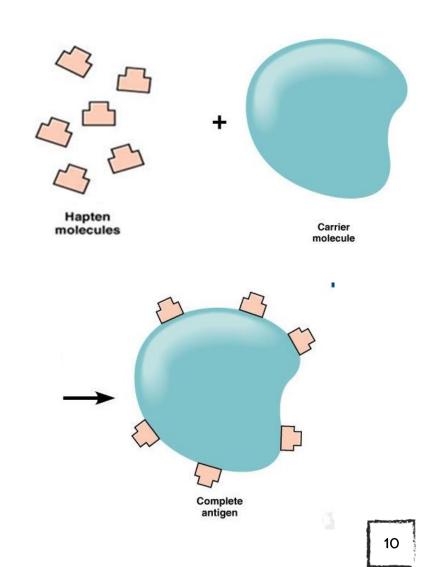
- 1. Inorganic Compounds: Alum, aluminum hydroxide
- 2. Mineral Oil: Paraffin oil
- 3. Bacterial Products: Killed bacteria (Bordetella pertussis, Mycobacterium bovis), toxoids
- 4. Freund's complete adjuvant, Freund's incomplete adjuvant

#### 3. Mechanisms of Action:

- 1. Extend antigen presence in blood
- 2. Aid in antigen absorption by presenting cells
- 3. Activate macrophages and lymphocytes
- 4. Support cytokine production

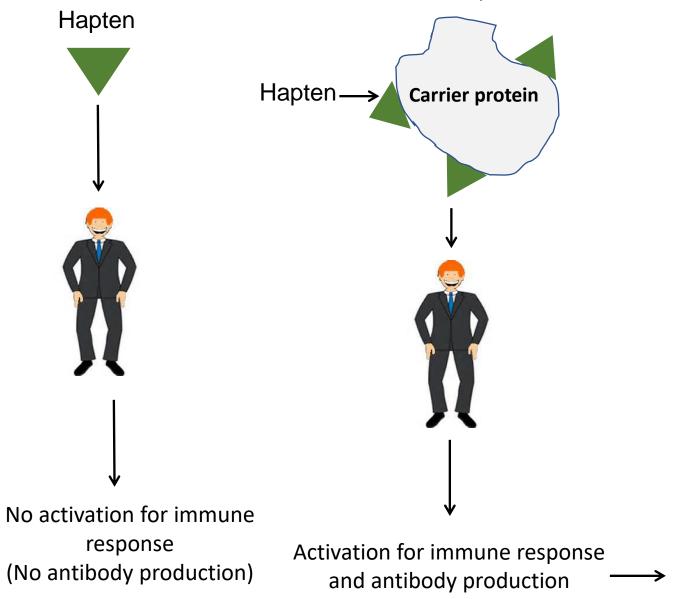
## **~**Haptens

- **1. Definition:** Nonantigenic molecules with low molecular weight (too small to be immunogenic) that cannot induce an immune response by themselves but react with products of that response.
- 2. Examples:
  - 1. Antibiotics
  - 2. Analgesics
  - 3. Other low-molecular weight compounds
- **3. Conversion to Antigen:** When a hapten is coupled to a larger carrier molecule (e.g., albumins, globulins, synthetic polypeptides), it becomes a complete antigen (immunogen).
- **4.** Clinical Significance: Certain haptens, such as penicillin, can cross-react with self-proteins, causing allergic responses.
- 5. Auto coupling Haptens:
  - Some haptens can form spontaneous covalent bonds with selfproteins, creating new antigens in vivo.
    This can lead to: Autoimmune diseases or Drug allergies



## **Haptens**

Exposure to hapten and carrier protein



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## Drug Allergy

VA

- **1. Definition:** Adverse immunological reactions to certain antigens, particularly antibiotics, which can pose significant medical problems.
- 2. Examples: Anaphylactic reactions to penicillin can be fatal.

#### 3. Mechanism of Reaction:

- Penicillin can form a hapten-carrier conjugate with self-proteins, acting as an immunogen that generates IgE antibodies.
- IgE antibodies bind to mast cells.
- During the second exposure, penicillin binds directly to IgE, activating mast cell degranulation.

#### 4. IgE Cross-Reactivity:

- Some anti-penicillin IgE antibodies cross-react with other antibiotics that have similar structures, such as cephalosporins and carbapenems.
- This complicates the treatment of bacterial infections in affected patients, as they may be unable to take necessary antibiotics.

## **~** Epitopes

**1. Definition:** Also known as antigenic determinants, epitopes are the sites on or within an antigen that antibodies react with.

#### 2. Comparison with Haptens: Epitopes and haptens are similar, but:

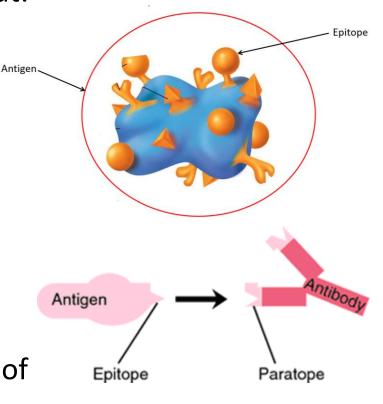
1. Haptens are artificially added to a molecule.

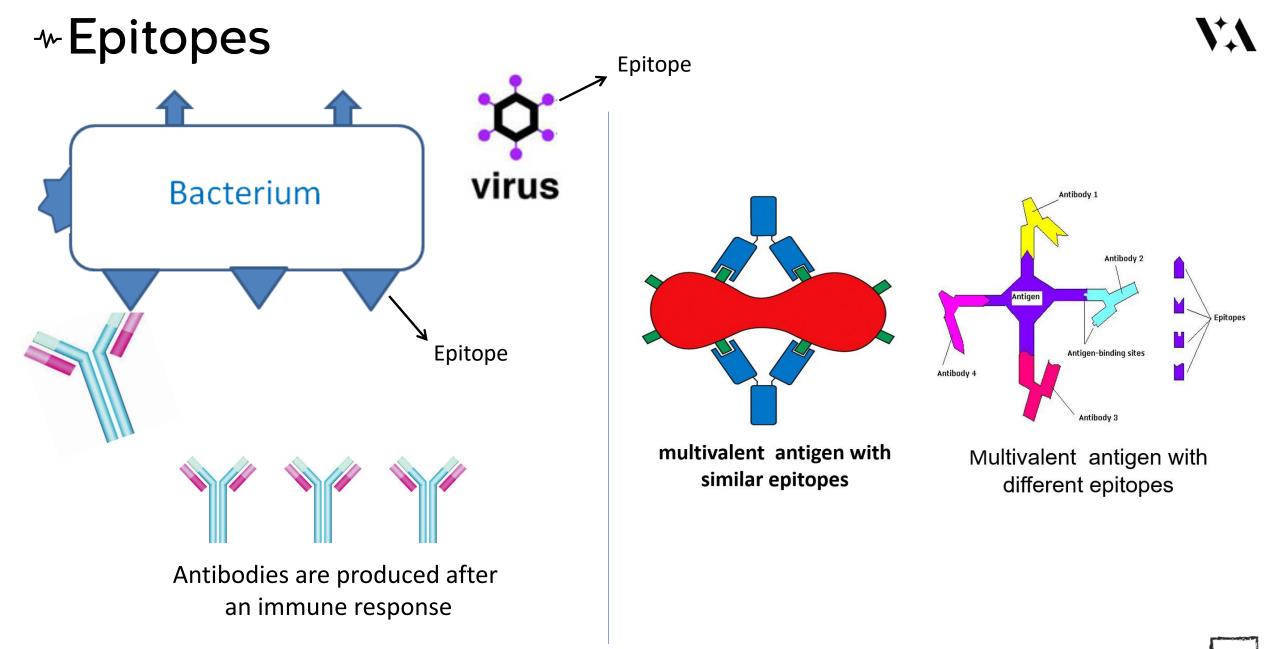
2. Epitopes are integral parts of the native molecule.

**3. Size:** Epitopes are very small, typically consisting of just four or five amino acid or monosaccharide residues.

#### 4. Types of Antigens:

- Antigens with many epitopes of different specificities are called polyvalent.
- Antigens with many epitopes of the same specificity are called multivalent.
- Diversity of Epitopes: An antigen molecule can carry a number of different epitopes, specifying different antibodies.





## 

- The immune response has several generalized features that distinguish it from other body systems, such as the respiratory and reproductive systems.
- Key Features:

a. **Specificity:** The ability to discriminate among different molecular entities rather than making a random, undifferentiated response.

b. **Discrimination:** The ability to differentiate between "self" and "nonself" antigens.

c. **Memory:** The ability to recall previous contact with foreign molecules and respond to them in a learned manner, resulting in a more rapid and larger response.

## **~**Types of Immunity

- **1. Acquired Immunity:** 
  - a. Natural Acquired Immunity:
  - 1. Active Immunity:
    - Antigens enter the body naturally, triggering responses from both the innate and adaptive immune systems.
    - Provides long-term protection.

#### 2. Passive Immunity:

- Antibodies pass from mother to fetus across the placenta.
- Infants receive antibodies through breast milk.
- Provides immediate short-term protection.

#### b. Artificial Acquired Immunity:

#### **1. Active Immunity:**

- Antigens enter the body through vaccination, prompting responses from innate and adaptive immune systems.
- Provides long-term protection.

#### 2. Passive Immunity:

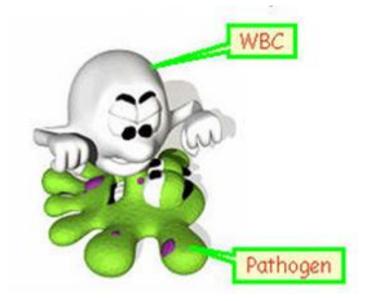
- Transfer of antibodies from immune individuals to a recipient.
- Provides immediate short-term protection.

#### 2. Adoptive Immunity:

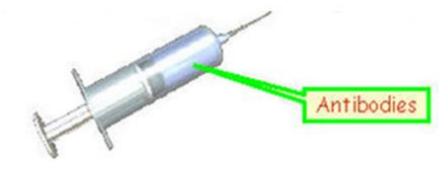
Refers to the transfer of immunity through the transfer of immune cells.

## Routes of acquiring immunity

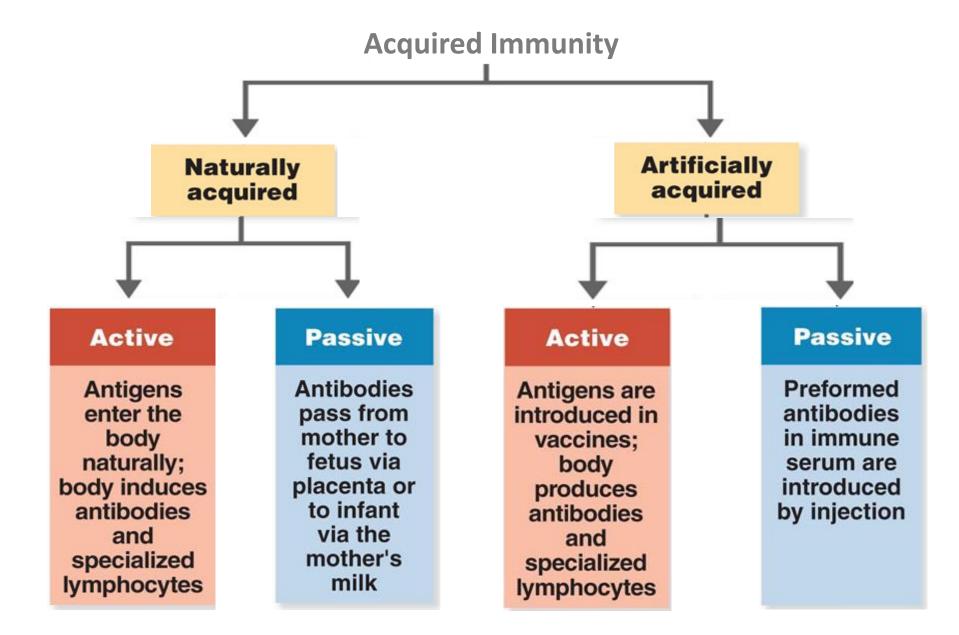
Active Immunity: Reaction of your own immune system



#### **Passive Immunity:** Borrow immune agents from other person









# Thank you



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