

The Immune System 免疫系統

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PowerPoint Lectures for

Biology: Concepts and Connections, Fifth Edition

- Campbell, Reece, Taylor, and Simon

Understand:

- how innate defenses help our body act against infection.

- how acquired immunity plays its role in our immune responses.

- what common disorders of the immune system are.

An AIDS Uproar

- Acquired immune deficiency syndrome (AIDS)
 - Is epidemic throughout much of the world
- Thousands of people are infected every day



- HIV, the virus that causes AIDS, attacks the immune system
 - And eventually destroys the body's ability to fight infection



Innate Defense Against Infection

24.1 Innate defenses against infection include the <u>skin</u> and <u>mucous membranes</u>, <u>phagocytic</u> <u>cells</u>, and <u>antimicrobial proteins</u>

- Innate immunity (先天性免疫)
 - Is present and effective long before exposure to pathogens
 - 與生俱來。
 - 不需要病原菌刺激即存在。
 - 反應快速,但持續性低。
 - 專一性差,且無記憶性。

- Microbes that breach the body's external defenses
 - Are engulfed and destroyed by macrophages



Colorized SEM 3,800×

• An overview of animal immune system

Innate immunity (24.1-3) Response is the same whether or not pathogen has been previously encountered		Acquired immunity (24.4-15) Found only in vertebrates; previous exposure to pathogen enhances immune response
External barriers	Internal defenses	•Antibodies (24.8-10) •Lymphocytes (24.11-14)
 Skin/exoskeleton Secretions Mucous membranes 	Skin/exoskeleton Secretions• Phagocytic cells • NK cells • Defensive proteins • Inflammatory response (24.2)• Lyr	
	The lymphatic system (24.3)	

- Interferons are proteins produced by virusinfected cells
 - That help other cells resist viruses



24.2 The inflammatory response mobilizes nonspecific defense forces

Tissue damage triggers the inflammatory response



- (Puposes) The inflammatory response
 - Can disinfect tissues, and
 - Can limit further infection from occurring
 - Can help heal the damaged tissues



24.3 The <u>lymphatic system</u> becomes a crucial battleground during infection

- The lymphatic system
 - Is a network of lymphatic vessels and organs



- The vessels collect fluid from body tissues
 - And return it as <u>lymph</u> to the blood
- Lymph organs such as the <u>spleen</u> and <u>lymph</u> <u>nodes</u>
 - Are packed with white blood cells that fight infections

Lymphatic system = Lymphatic organs + lymphatic vessels + lymph

Circulation system

= Heart + blood vessels + blood

Acquired Immunity (獲得性/後天性免疫)

24.4 The immune response counters specific invaders

- Our immune system
 - Responds to foreign molecules called <u>antigens</u> (Ags)
- The immune system reacts to antigens



- We can temporarily acquire passive immunity
 - By receiving "premade" antibodies
- Infection or vaccination
 - Triggers active immunity



Figure 24.4

24.5 Lymphocytes mount a dual defense

- Two kinds of lymphocytes carry out the immune response
 - B cells secrete antibodies that attack antigens
 - T cells attack cells infected with pathogens



- Millions of kinds of B cells and T cells, each with *different* membrane receptors
 - Wait in the lymphatic system, where they may respond to invaders



Figure 24.5B

24.6 Antigens have specific regions where antibodies bind to them

- Antigenic determinants ('epitopes')
 - Are the specific regions on an antigen to which antibodies bind



24.7 <u>Clonal selection</u> musters defensive forces against specific antigens

- When an antigen enters the body
 - It activates only a small subset of lymphocytes with complementary receptors

A single progenitor cell gives rise to a large number of lymphocytes, each with a different specificity

- The selected lymphocyte cells multiply into clones of short-lived effector cells
 - Specialized for defending against the antigen that triggered the response



The Steps of Clonal Selection

- In the primary immune response, clonal selection
 - Produces <u>effector cells</u> and <u>memory cells</u> that may confer lifelong immunity
- In the secondary immune response
 - Memory cells are activated by a second exposure to the same antigen, which initiates a faster and more massive response

• The primary and secondary immune responses **Primary immune** Antigen response molecules Antigen receptor (antibody on cell B cells with different surface) response First exposure to antigen Cell activation: growth, division, Antibody and differentiation molecules Endoplasmic Clone of memory cells reticulum Antigen Plasma (effector) cells secreting antibodies molecules **Second exposure** Antibody Endoplasmic to the same antigen molecules reticulum **Secondary immune** response Clone of memory cells Figure 24.7A Plasma (effector) cells secreting antibodies

Primary vs. Secondary Immune Response

- The primary immune response
 - Is slower than the secondary immune response



24.8 Antibodies are the weapons of humoral immunity

- Antibody molecules
 - Are secreted by plasma (effector) B cells





- An antibody molecule
 - Has antigen-binding sites specific to the antigenic determinants that elicited its secretion



24.9 Antibodies mark antigens for elimination

- Antibodies promote antigen elimination
 - Through several mechanisms



CONNECTION

24.10 Monoclonal antibodies are powerful tools in the lab and clinic

- Monoclonal antibodies
 - Are produced by fusing B cells specific for a single antigenic determinant with easy to grow tumor cells
 - Tumor cell \rightarrow immortality
 - B cells (speen) → make Abs



- Monoclonal antibodies (mAbs)
 - are useful in research, diagnosis, and treatment of certain cancers

24.11 Helper T cells stimulate humoral and cellmediated immunity

- Helper T cells and cytotoxic T cells
 - Are the main effectors of cell-mediated immunity
- Helper T cells
 - Also stimulate the humoral responses

- In cell-mediated immunity, an <u>antigen-</u> presenting <u>cell (APC)</u>
 - Displays a foreign antigen and one of the body's own self proteins to a <u>helper T cell</u>





- The helper T (Th) cell's receptors (TCR)
 - Recognize the self-nonself complexes and the interaction activates the helper
 T cells
- The helper T cell
 - Can then activate cytotoxic T (Tc) cells and B cells
 L Antigen is up-taked by APC



• The activation of a helper T cell and its roles in immunity

Self-nonself complex = MHC + Ag fragment



CONNECTION

24.12 HIV destroys helper T cells, compromising the body's defenses

- The AIDS virus attacks helper T (Th) Cells
 - Opening the way for opportunistic infection



A human helper T cell (green) under attach by HIV (red dots).

Figure 24.13

24.13 Cytotoxic T (Tc) cells destroy infected body cells

- Cytotoxic T cells
 - Bind to infected body cells and destroy them



Figure 24.12

24.14 Cytotoxic T (Tc) cells may help prevent cancer

- Cytotoxic T cells may attack cancer cells
 - Which have abnormal surface molecules



24.15 The immune system depends on our molecular fingerprints

- The immune system
 - Normally reacts <u>only</u> against nonself substances, not against self
 - May reject transplanted organs because these cells lack the unique "fingerprint" of the recipient's self proteins

DISORDERS OF THE IMMUNE SYSTEM CONNECTION

- 24.16 Malfunction or failure of the immune system causes disease
 - In autoimmune diseases
 - The system turns against the body's own molecules
 - In immunodeficiency diseases
 - Immune components are lacking, and frequent infections recur
 - Physical and emotional stress
 - May weaken the immune system

24.17 Allergies are <u>overreactions</u> to certain environmental antigens

- Allergies
 - Are abnormal sensitivities to antigens (allergens) in the surroundings



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Figure 24.17

In mammals, there are 5 Ab types - IgG, IgM, IgA, IgD, IgE

Summary

- Innate defenses include the skin and mucous membranes, phagocytic cells, and anti-microbial proteins.
- The inflammation mobilizes nonspecific defense forces.
- The lymphatic system is a crucial system during infection.
- Antigens have specific regions where antibodies can bind.
- Helper T cells help stimulate humoral and cell-mediated immunity, whereas cytotoxic T cells destroy infected body cells.
- Allergies are overreactions to certain environmental antigens.

End of Chapter

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