26.1 Defence against disease

Name: _____ Class: _____ Date: _____ **Total: 28 marks**

KEY WORDS pathogen • antigen • antibody • phagocyte • memory cell • herd immunity

Objective

Build the skills to answer exam questions on **defence against disease** : pathogens, barriers and clotting, innate and adaptive immunity, antibodies, vaccines, HIV and antibiotics.

You must be able to:

- describe pathogens and the first line of defence, including blood clotting
- compare innate and adaptive immunity, and explain the roles of phagocytes and lymphocytes
- explain how antibodies and memory cells are made, and how vaccines and herd immunity work
- explain the effect of HIV on the immune system, and why antibiotic resistance is a problem
- explain the evolution of antibiotic resistance and the risk of zoonoses, using data

1 Key knowledge and worked examples

■ Defence against disease

Pathogens are organisms or viruses that cause disease: bacteria, viruses, fungi, protists. **First line:** the **skin** (tough outer layer of dead cells) and **mucous membranes** (sticky mucus traps microbes). **Blood clotting** seals cuts: platelets release clotting factors, which start a cascade ending with **thrombin** converting **fibrinogen** into insoluble **fibrin** threads that trap blood cells.

innate immunity	**adaptive immunity**
present from birth, non-specific	develops after exposure, specific
phagocytes engulf pathogens	**lymphocytes** respond to particular **antigens**
the same response each time	faster and stronger the second time (**memory cells**)

Antibody production: a phagocyte presents the antigen; a **helper T-cell** activates the matching **B-cell**; the B-cell divides (**clonal selection**) into **plasma cells**, which secrete antibodies, and **memory cells**. **Antibodies** bind antigens, making pathogens clump and marking them for phagocytes.

HIV infects helper T-cells; as their number falls, B-cells cannot be activated, so the body cannot fight infections (**AIDS**). HIV spreads in blood, sexual fluids and from mother to baby.

Vaccines contain antigens (weakened or killed pathogens, or mRNA coding for an antigen); they cause memory cells to form. **Herd immunity:** when enough people are immune, the pathogen cannot spread, which protects those who cannot be vaccinated. **Antibiotics** kill bacteria or stop their growth but do **not** affect viruses.

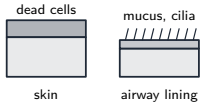
HL extension: antibiotic **resistance** evolves by natural selection: a random mutation gives resistance; antibiotics kill the others; resistant bacteria multiply. **Zoonoses** are diseases that pass from animals to humans (e.g. COVID-19, rabies).

Worked example. If each infected person infects $R_0 = 4$ others, about $1 - \frac{1}{4} = \mathbf{75}\%$ of people must be immune for herd immunity.

Answer shapes: antigen → helper T-cell → B-cell clones → plasma cells (antibodies) and memory cells.

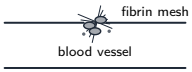
2 Practice

2.1 Describe how the skin and mucous membranes protect the body against pathogens. [2]

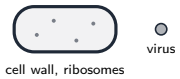


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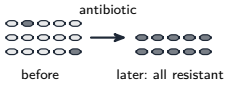
2.2 Explain how blood clots at a cut. [3]



2.3 Explain why antibiotics can treat bacterial infections but not viral infections. [3]



2.4 Explain how antibiotic resistance spreads in a bacterial population. [2]



2.5 State what is meant by a zoonosis, with one example. [2]



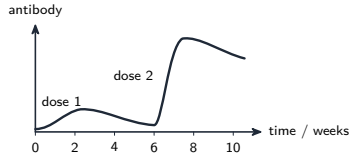
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3 Exam-style questions

3.1 The graph shows the concentration of antibodies in the blood after two doses of a vaccine.

(a) Compare the responses to the first and second doses. [2]

(b) Explain why the response to the second dose is different. [3]



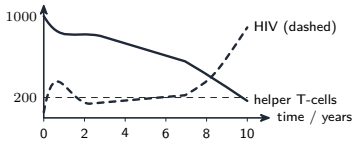
(c) State what a vaccine contains. [1]

(d) Explain how herd immunity protects people who cannot be vaccinated. [2]

3.2 HIV infects helper T-cells. The graph shows the number of helper T-cells and the amount of HIV in a patient over 10 years without treatment.

(a) Describe the change in helper T-cell count. [2]

(b) Explain why the patient becomes ill with other infections when the count is very low. [3]



(c) State two ways in which HIV is transmitted. [2]

(d) Distinguish between innate and adaptive immunity. [1]

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