

8. What are the first cells that recognize a processed and presented T-dependent antigen?
 B lymphocytes Memory cells Macrophages T helper cells
9. Which of the following is not an assumption which can be derived from the clonal selection theory?
 Determined cells have begun to differentiate, but are not fully differentiated.
 Each genetically different type of lymphocyte expresses a single specificity.
 Any lymphocyte that could possibly mount a harmful response against self molecules is eliminated or suppressed
 Development of both T and B and T lymphocytes follow a similar pattern
10. All the following are characteristics of both MHC class I and class II molecules *except*:
 They are expressed codominantly.
 They are expressed constitutively on all nucleated cells.
 They are glycosylated polypeptides with domain structure.
 They are involved in presentation of antigen fragments to T cells.

SECTION "B"

[5 Q × 1 = 5 marks]

Fill in the blanks:

11. The ratio of T cells to B cells is _____.
12. The immunoglobulin class which is the least abundant in normal adult is _____.
13. _____ selection of thymocytes is necessary to produce a T-cell repertoire capable of interacting with self-MHC molecules.
14. Body's own cells are protected from membrane attack complex by surface glycoprotein called _____.
15. The specificity of an antibody is due to _____.

SECTION "C"

[5 Q × 1 = 5 marks]

Define the following:

16. F(ab')₂ fragment of antibody:
17. Discontinuous epitope:
18. Anergic B cells:
19. Linked recognition:
20. Granzymes:

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SECTION "D"

[7Q × 5 = 35 marks]

Attempt *ALL* questions.

1. A person receives a live attenuated vaccine for a certain disease. Discuss the antigen processing and presentation pathway that follows to develop immunity to that pathogen.
2. The innate immune system uses two different strategies to identify pathogens: recognition of non-self and recognition of self. Give examples of each and discuss how each example contributes to the ability of the organism to protect itself from infection.
3. Enlist the four main processes that generate the diversity in the lymphocyte repertoire and elaborate on somatic hypermutation.
4. Discuss the process through which double negative thymocytes develop into single positive thymocytes.
5. Elaborate on the role of antigen presenting cells in clonal expansion and differentiation of naïve T cells.
6. Discuss how positive and negative selection of thymocytes co-ordinate with each other for survival of T lymphocytes.
7. Discuss the role of inflammatory barrier in immunity

SECTION "E"

[5Q × 2 = 10 marks]

8. Give *TWO* major differences between (*ANY FIVE*)
 - a. Cell mediated and Humoral Immunity
 - b. Isotype switch and Receptor editing
 - c. Structure of MHCI and MHCII
 - d. Armed effector lymphocytes and Naïve lymphocytes
 - e. Thymus dependent and Thymus independent antigens
 - f. Dendritic cells and Macrophages

SECTION "F"
[5Q × 2 = 10 marks]

9. Explain WHY/HOW for *ANY FIVE* questions.
- a. Antibodies, which all have the same basic shape, recognize antigens of a wide variety of different shapes
 - b. B cell receptor and T cell receptor are similar in many aspects.
 - c. In the absence of Invariant chain (Ii), many MHC class II molecules are retained in the endoplasmic reticulum as complexes with misfolded proteins.
 - d. Spontaneous hydrolysis of the thioester bond in C3 occurs at a continuous low rate even in the absence of infection. However this does not cause damage to normal host cells.
 - e. Complement 3 deficiency is more serious clinically than complement 1 deficiency
 - f. Some microorganisms produce enzymes that can degrade the F_c portion of antibody molecules and this is advantageous for their survival.