

MCBACOR12T-MICROBIOLOGY (CC12)

Time Allotted: 2 Hours

The figures in the margin indicate full marks. Candidates should answer in their own words and adhere to the word limit as practicable. All symbols are of usual significance.

Question No. 1 is compulsory and answer any *four* from the rest

1.	Answer	any four	questions	from	the	follow	ing
1.	1 1115 W CI	any jour	questions	nom	une	10110 W	mg.

- (a) Define allotypic and idiotypic determinants.
- (b) Define the function of bone marrow stromal cells in haematopoiesis.
- (c) What are T1 antigens? Give two examples of T1 antigens.
- (d) What do you mean by clonal anergy?
- (e) What is chimeric antibody?
- (f) What are exogeneous and endogeneous antigens?
- (g) What are NKT cells?
- (h) Mention any two biological consequences of complement activation.

2.	(a)	'The central event of complement activation is proteolysis of C3 molecule'. — Justify this statement.	3
	(b)	Explain why the red blood cells of an individual are not normally destroyed as a result of innocent-bystander lysis by complements.	1
	(c)	Briefly explain the mechanism of action of the following regulatory proteins of the complement activation pathways:	1+1
		(i) C1 inhibitor (C1 Inh), (ii) Factor H	
	(d)	Define affinity and avidity in relation to antigen-antibody interactions.	2
3.	(a)	Draw the structure of IgG with proper labeling.	1
	(b)	Explain the function of hinge region.	1
	(c)	Explain the process of 'ADCC'.	2
	(d)	Describe the principle of selection of monoclonal antibody-producing B-cells.	4
4.	(a)	What is meant by sensitized mast cell in Type I hypersensitivity?	1
	(b)	Name any two pre-formed and any two newly-synthesized mediators involved in Type I hypersensitivity.	2

Full Marks: 40

 $2 \times 4 = 8$

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	(c)	Write a short note on Erythroblastosis Fetalis.	3
	(d)	Define prozone effect.	2
5.	(a)	Mention the proposed mechanisms involved in development of autoimmune diseases.	3
	(b)	Name any two immunodeficiency diseases.	1
	(c)	What is central tolerance? How does it differ from peripheral tolerance?	2
	(d)	What are tumor antigens?	1
	(e)	What do you mean by paratope and agretope?	1
6.	(a)	Explain the ELISPOT technique.	3
	(b)	Distinguish between primary immune response and secondary immune response.	2
	(c)	What is the principle behind immunofluorescence technique?	2
	(d)	What do you mean by passive agglutination?	1
7.	(a)	Mention any one contribution of (i) Rodney Porter and (ii) Karl Landsteiner in the field of Immunology.	1
	(b)	What are Anaphylatoxins?	1
	(c)	Explain the antigen presentation process by the cytosolic pathway.	2
	(d)	Why are circulating IgM unable to activate the complement pathway?	2
	(e)	What are Superantigens?	2
8.	(a)	Define: (i) Haptens, (ii) Adjuvants.	2
	(b)	Following pair of antigens listed below are:-	2
		(i) Native bovine serum albumin (BSA)	
		(ii) Heat – denatured BSA	
		Which one of these above is a better immunogen? Justify your answer.	
	(c)	What are T & B cell epitopes?	2
	(d)	What are the differences between activated and resting lymphocytes?	2
9.	(a)	Mention one application of Flow cytometry.	1
	(b)	What are granzymes?	1
	(c)	How do cytotoxic T lymphocytes kill the target cells? Explain diagrammatically.	3
	(d)	What do you mean by co-stimulatory signal?	2
	(e)	What do you mean by bacterial agglutination?	1

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N.B.: Students have to complete submission of their Answer Scripts through E-mail / Whatsapp to their own respective colleges on the same day / date of examination within 1 hour after end of exam. University / College authorities will not be held responsible for wrong submission (at in proper address). Students are strongly advised not to submit multiple copies of the same answer script.