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Reg. No.:					

# **Question Paper Code: 94B05**

## B.E. / B.Tech. DEGREE EXAMINATION, NOV 2023

#### Fourth Semester

## **Biomedical Engineering**

## 19UBM405- PATHOLOGY AND MICROBIOLOGY

(Regulation 2019)

Duration: Three hours Maximum: 100 Marks

#### **Answer ALL Questions**

	Allswei ALL Questions					
	PART A - $(10 \times 2 = 20 \text{ Marks})$					
1.	Investigate cell injury and cell death.					
2.	How do you implement tissue processing for tumour using microtome?					
3.	How do you analyze clotting time using capillary tubes?	CO3-Ana				
4.	How do you differentiate Leukemia patients with normal human?					
5.	How do you distinguish moist Heat and cold killing of microbes.					
6.	Draw a schematic diagram of TEM.					
7.	Give a list of cancer causing chemical and physical mutagens.	CO3-Ana				
8.	Draw a schematic diagram of operon model?	CO3-Ana				
9.	Define phagocytosis.	CO1-U				
10.	How antibodies are more specific to antigens?	CO3-Ana				
	$PART - C (5 \times 16 = 80 \text{ Marks})$					
11.	(a) Give a brief note on intracellular accumulations and analyse its various types due to the accumulation process?  Or	CO3- Ana (16)				

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(b) Give a brief notes on cellular adaptations. Comparative study of CO3- Ana (16) hypertrophy with hyperplasia.

12. (a) Describe Hematological disorders and its causing factors in CO3-Ana (16)human. Demonstrate of bleeding time for analysis of the fluid in the human body. Or (b) Describe Bleeding disorders and its impact on human health. CO3- Ana (16)Demonstrate of bleeding time for analysis of the fluid in the human body. 13. (a) How do you isolate pure bacterial strains from mixed population CO3- Ana (16)using spread and streak plate technique? Give your suggestion for the simple and suitable method. Or (b) Describe media preparation and sterilization process. CO3- Ana (16)Comparative analysis of physical and chemical techniques for sterilization. Give a brief notes on operon concepts. Investigate operon model CO1- U 14. (a) (16)to regulate gene expression in Gram negative bacteria? (b) Explain in detail on Bacterial genetic system. Distinguish CO1-U (16)transformation and transduction techniques with examples. 15 (a) Give a brief note on Immunological techniques. How do you use CO1- U (16)mono-clonal antibody technology for the bacterial treatment?

(b) Define mono-clonal antibody technology. How do you use mono- CO1- U

clonal antibody technology for the cancer treatment?

(16)