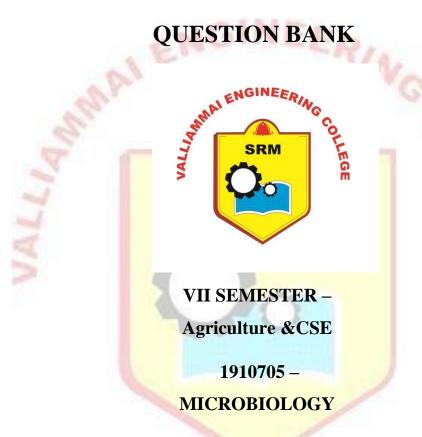
SRM VALLIAMMAI ENGINEERING COLLEGE

(An Autonomous Institution) SRM Nagar, Kattankulathur – 603 203

DEPARTMENT OF MEDICAL ELECTRONICS



Regulation – 2019 Academic Year: 2022 – 23 Odd Semester

Prepared by

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SRMVALLIAMMAI ENGINEERING COLLEGE

(An Autonomous Institution)





UNIT I – INTRODUCTION TO MICROBIOLOGY

Classification and nomenclature of microorganisms, microscopic examination of microorganisms: Light, Fluorescent, Dark field, Phase contrast, and Electron microscopy.

	PART – A		
Q.No	Questions	BT	Competence
	GINEE	Level	
1	Define the microorganism classification groups Protozoa & Slime moulds.	BTL1	Remembering
2	Identify the significance of Algae & Bacteria classification biological groups.	BTL1	Remembering
3	List the different sources for microorganism names.	BTL1	Remembering
4	Differentiate between classification and nomenclature.	BTL4	Analyzing
5	Write the various biological groups used for microorganism classification.	BTL3	Applying
6	Show the various taxonomic ranks used for classification.	BTL3	Applying
7	Enlist some cellular microorganisms based on clinical purpose.	BTL1	Remembering
8	When to use initials during naming the microorganisms?	BTL1	Remembering
9	Enumerate various rules used in nomenclature.	BTL1	Remembering
10	Categorize the steps used to name subspecies and serovars.	BTL4	Analyzing
11	Examine the nomenclature used during the naming of microorganism using	BTL4	Analyzing
	binary names.		
12	Employ the significance of classification of microorganism in clinical practice.	BTL3	Applying
13	Articulate the principal types of viruses causing human disease.	BTL3	Applying
14	Inspect the various direct methods used for identification of bacteria.	BTL4	Analyzing
15	Categorize various indirect methods used for identification of bacteria.	BTL4	Analyzing
16	Explain the importance of microscopy identification method.	BTL4	Understanding
17	Indicate the significance of the cultural characteristics method.	BTL2	Understanding
18	Review the kind of information you think that can be provided	BTL2	Understanding
	by microscopic observations of microorganisms.		
19	How did Pasteur and Tyndall finally settle the spontaneous generation controversy?	BTL2	Understanding
20	Extend the set of information that you think can be provided	BTL2	Understanding
	by isolating microorganisms from their natural environment and culturing		
	them in the laboratory.		
21	Why was the belief in spontaneous generation an obstacle to the development	BTL2	Understanding

	of microbiology as a scientific discipline?			
22	Use the various steps involved in scientific method used to conduct		BTL3	Applying
	experimental research.			PP-J8
23	Illustrate the scientific method used in experimental science. How does a theory differ from a hypothesis? Why is it important to have a control group	n?	BTL3	Applying
24		ρ.	BTL4	A 1
24	Criticize Koch's postulates. What is a pure culture? Why are pure cultures		B1L4	Analyzing
	important to Koch's postulates? PART – B			
1			BTL2	Undougtonding
1	(i) Explain the various stages involved in designing and conducting	(0)	DIL2	Understanding
	_ •	(9)		
	(ii) Describe the role of Koch's postulates to demonstrate that mycobacteriu			
_	tuberculosis is the causative agent of tuberculosis.	(4)	DTI 2	II J4 J:
2	Classify the five most important research areas to pursue	(12)	BTL2	Understanding
2		(13)	DTI A	Timed acceptance 10
3	Indicate the various aspects of resolution, numerical aperture, working	(12)	BTL2	Understanding
4		(13)		TI-dame.
4	Report the various rules of nomenclature used during naming of	(12)	BTL1	Understanding
		(13)	DOTE 4	- ·
5	Enlist the parts of a light microscope and describe their functions. (13)		BTL1	Remembering
6	Illustrate the following		BTL3	Applying
U	(i) refraction and refractive index.	(5)	DILS	Applying
	(ii) focal point.	(4)	140	
	(iii) focal length.	(4)	111	
7	Explain the importance of microscope resolution and numerical aperture wi	` '	BTL3	Applying
'	The state of the s	(13)	DILS	Applying
0		(13)	DEL 2	A 1 ·
8	Interpret a method to convert light microscope to a dark field microscope (13)		BTL3	Applying
0			DEL 2	A 1 •
9	Illustrate the following concept in detail		BTL3	Applying
	(i) The formation of a micrograph of the protozoan Amoeba proteus using			
	DIC microscope. (6)			
10	(ii) The condenser lens system in dark field microscope (7))	BTL1	Remembering
10	State in detail the principle of production of contrast in a phase-contrast microscopy.	13)	DILI	Kemembering
11	Model a microscope with the following details	13)	BTL4	Analyzing
11	(i) If a specimen is viewed using a 5X objective in a microscope with	ith	DIL4	Anaryzing
	a 15X eye piece. (7)			
	(ii) Why don't most light microscopes use 30X ocular lenses for			
	greater magnification? (6)			
12	Analyze the function of immersion oil.	(13)	BTL4	Analyzing
13	Define and identify various steps and stages involved in staining of		BTL2	Understanding
	microorganisms.	(13)		

14	Enumerate the characteristics of light and transmission electron microsc	opes in	BTL1	Remembering
	a tabular column.	(13)		
15	List the limits of microscopic resolution.	(13)	BTL1	Remembering
16	Examine an overview of TEM operation and compare it with the operation	on	BTL4	Analyzing
	of light microscope.	(13)		
17	Inspect the significance of lenses and bending of light and tabulate the		BTL4	Analyzing
	common units of measurement used in microscopy.	(13)		
	PART – C	•		
1	Identify the working of fluorescence microscope system		BTL4	Analyzing
1	with a neat diagram.	(15)	DIL	Anaiyzing
2	Infer the specimen shadowing and transmission electron micrograph fro	` /	BTL2	Understanding
	transmission electron microscope system.	(15)		
3	Under what circumstances would it be desirable to prepare specimens for	or	BTL3	Applying
	the TEM by use of negative staining? Shadowing? Freeze-etching?	(15)		
4	How does the scanning electron microscope operate and in what way	0	BTL1	Remembering
	does its function differ from that of the TEM? The SEM is used to study	1		
	which aspects of morphology?	(15)	-	
5	Explain the following:		BTL2	Understanding
	(i) Fixation, dye and chromophore.	(5)	1	
	(ii) Basic dye, acidic dye and simple staining.	(5)	TI	
	(iii) Differential staining, negative staining, and acid-fast staining	. (5)	-	

UNIT II – MICROBES- STRUCTURE AND REPRODUCTION

Structural organization and multiplication of bacteria, Viruses (TMV, Hepatitis B), Algae(cyanophyta, rhodophyta) and Fungi (Neurospora), Life history of actinomycetes (Streptomyces), Yeast (Saccharomyces), Mycoplasma (M. pneumoniae) and Bacteriophages (T4 phage, lphage)

	PART – A				
Q.	Questions	BT	Competence		
No		Level			
1	What characteristic shapes can bacteria assume? Describe the ways in	BTL1	Remembering		
	which bacterial cells cluster together.				
2	Label and draw a bacterial cell and label all important structures.	BTL1	Remembering		
3	List the functions of the procaryotic plasma membrane	BTL1	Remembering		
4	Prepare a diagram to represent the schematic of bacterial plasma membrane.	BTL3	Applying		
5	Write the structure of a gas vacuole to its function.	BTL3	Applying		
6	Illustrate how cell wall structure and sugar content are used to classify the	BTL3	Applying		
	actinomycetes				
7	Interpret seven steps are involved in the infection process and pathogenesis	BTL1	Remembering		
	of bacterial diseases?				
8	Mention the significance of nucleocapsid, capsid, icosahedral capsid, helical	BTL1	Remembering		
	capsid.				
9	Enumerate the major properties of the genus Streptomyces.	BTL1	Remembering		
10	Inspect the importance of gapped DNA in hepadnavirus.	BTL4	Analyzing		
11	Seperate three ways in which Streptomyces is of ecological importance.	BTL4	Analyzing		

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12	Articulate the reproductive technique in Cyanophyta & Rhodophyta.		BTL2	Understanding
13	Identify the `general properties of actinomycetes.		BTL2	Understanding
14	Indicate with appropriate label and draw the helical model of TMV.		BTL2	Understanding
15	Recognize the process involved in reproduction of the virus.		BTL2	Understanding
16	Categorize the yeast and mold based on their differences.		BTL4	Analyzing
17	Describe the various types of reproduction fungi.		BTL2	Understanding
18	Review the various stages in lifecycle of Saccharomyces		BTL2	Understanding
	cerevisiae			
19	Prepare a table with various cellular organelles in a yeast cell.		BTL3	Applying
20	Interpret different characteristics of neurospora genus of ascomycete fun	ıgi.	BTL3	Applying
21	Point out the morphological structure of yeast with a diagram		BTL4	Analyzing
22	Outline the lifecycle of a bacteriophage		BTL4	Analyzing
23	Sketch various stages involved in the assembly of T4 Bacteriophage.		BTL3	Applying
24	Analyze the morphological structure of T4 Bacteriophage with a neat dia	agram	BTL4	Analyzing
	PART-B	3		
1	(i) Describe a bacterial cell and label all important structures.	(7)	BTL2	Understanding
	(ii) Indicate the various functions of prokaryotic structures		3	
	found in bacteria	(6)	-	
2	(i) Compare the following terms: nucleocapsid, capsid, icosa		BTL4	Analyzing
	capsid, helical capsid, complex virus, binal symmetry, pro-		1	
	capsomer, pentamer or penton, and hexamer or hexon. (ii) How do pentamers and hexamers associate to form a con-	(8) mplete	111	
	icosahedron; what determines helical capsid length	присс	(G)	
	and diameter?	(3)	177	
	(iii) Illustrate the TMV structure with a neat diagram	(2)		
3	Explain the importance of the Australian antigen and dane particle in		BTL2	Understanding
	discovery of morphological structure of Hepatitis B.	(13)		
4	Write the organization and reproduction of cyanoaphyta.	(13)	BTL1	Remembering
5	Predict the cell wall structure and sugar content are used to classify the		BTL3	Applying
	actinomycetes. Include a brief description of the four major wall types	(13)		
6	Illustrate the following with a neat diagram		BTL3	Applying
	(i) The Fluid Mosaic Model of Membrane Structure	(5)		
	(ii) The Structure of a Polar Membrane Lipid.	(4)		
	(iii) Bacterial membranes.	(4)		
7	Show three ways in which Streptomyces is of ecological importance.		BTL3	Applying
	Why do you think Streptomyces spp. produce antibiotics?	(13)		
8	Name the various the major properties of the genus Streptomyces.	(13)	BTL1	Remembering
9	Interpret the morphological significance of suborder streptomycineae	(13)	BTL3	Applying
10	Explain in detail the various stages in binary fission reproduction method	d	BTL1	Remembering
	with neat diagram.	(13)		
11	Describe the various types of bacteriophages.	(13)	BTL1	Remembering
		·		

12	Enumerate the significance of different stages in cell cycle of E coli. (13)	BTL2	Understanding
13	Review the cell reproduction cycle of mycoplasma. (13)	BTL2	Understanding
14	Illustrate the relationship between Mycoplasma genome size and growth requirements. (13)	BTL3	Applying
15	Categorize the various stages involved in the life cycle of the yeast Saccharomyces. (13)	BTL4	Analyzing
16	Analyze the Asexual reproduction in the fungi and some representative spores with diagrammatic illustrations. (13)		Analyzing
17	Examine the morphology of t4 phage with a neat diagram. (13)	BTL4	Analyzing
	PART – C		
1	Analyze the various stages in the life cycle of the Bacteriophage. Describe briefly the lytic and lysogenic cycle. (15)	BTL4	Analyzing
2	Describe the endocytic pathway and the three routes	BTL2	Understanding
	that deliver materials to lysosomes for digestion. Which type of endocytosis	1	
	does not deliver ingested material to lysosomes? (15)	1	
3	Write the significance of the following	BTL1	Remembering
	(iii) Rough & smooth EPR. (5)	G	
	(ii) Golgi apparatus. (5)	-	
4	(iii) Mitochondria. (5) Illustrate the structure of the nucleus. What are euchromatin and	BTL3	Applying
7	heterochromatin? What is the role of the pores in the nuclear envelope? (15)		Applying
5	What is meiosis, how does it take place, and what is its role in the microbial life cycle? (15)	BTL3	Applying

UNIT III - MICROBIAL NUTRITION, GROWTH AND METABOLISM

Nutritional classification of microorganisms based on carbon, Energy and electron sources . Definition of growth, Balanced and unbalanced growth, Growth curve and different methods to quantify bacterial growth:(counting chamber, viable count method, counting without equipment),Different media used for bacterial culture (defined, complex, selective, differential, enriched),The mathematics of growth-generation time, Specific growth rate.

DA	\mathbf{DT}		A
PA	KI	_	A

Q. No	Questions	BT Level	Competence
1	Define heterotroph and autotroph.	BTL1	Remembering

2	What are nutrients? On what basis are they divided into macroelements and trace element	BTL1	Remembering
3	What are the six most important macroelements? How do cells use them?	BTL1	Remembering
4	What are growth factors? What are vitamins?	BTL3	Applying
5	How can humans put to use a microbe with a specific growth factor Requirement?	BTL3	Applying
6	Illustrate the growth factors that microorganisms produce industrially.	BTL3	Applying
7	Why do you think amino acids, purines, and pyrimidines are often growth factors, whereas glucose is not?	BTL1	Remembering
8	Mention the two pathways function during the procaryotic cell cycle.	BTL1	Remembering
9	How does the procaryotic cell cycle compare with the eucaryotic cell cycle?	BTL1	Remembering
10	Tabulate the sources of carbon, energy, and electrons.	BTL4	Analyzing
11	Categorize two trace elements. How do cells use them?	BTL4	Analyzing
12	Review the four phases of the growth curve in a closed system.	BTL2	Understanding
13	Differentiate the principles of turbidity and microbial mass measurement.	BTL4	Analyzing
14	Inspect the role of ATP in metabolism.	BTL4	Analyzing
15	Analyze the equations for the endergonic or exergonic reactions.	BTL4	Analyzing
16	Do living cells increase entropy within themselves? Do they increase entropy in the environment?	BTL4	Analyzing
17	What are cardinal temperatures?	BTL2	Understanding
18	Infer the significance between the dilution rate and maintenance energy	BTL2	Understanding
19	Mention the importance of growth rate rise with increasing temperature and then fall again at higher temperatures.	BTL2	Understanding
20	Indicate the significance of growth in living organisms.	BTL2	Understanding
21	Estimate the generation or doubling time and the mean growth rate constant.	BTL2	Understanding
22	Interpret the phenomenon of entropy and enthalpy.	BTL3	Applying
23	Employ a generalized method to write equation for a redox reaction.	BTL3	Applying
24	Show the significance of energy cycle and ATP's role in it.	BTL3	Applying
1	PART – B	DTT 4	IIndot 1°
1	Review the ways in which microorganisms are classified based on their requirements for energy, carbon, and electrons. (13)	BTL2	Understanding
2	requirements for energy, carbon, and electrons. (13) Indicate the nutritional requirements of the major nutritional groups	BTL2	Understanding
	and give some microbial examples of each. (13)	V 1 L/2	o naci stanting
3	Explain facilitated diffusion, active transport, and group translocation in terms	BTL2	Understanding
	of their distinctive characteristics and mechanisms. (13)	·	
4	Describe the significance of measurement of cell numbers. (13)	BTL1	Remembering
5	Distinguish between turbidity and microbial mass measurement. (13)	BTL2	Understanding

6	Examine the concept of membrane filtration procedure	(13)	BTL4	Analyzing
7	Interpret the concept of specific growth rate and apply it to a real-time scenario.	(13)	BTL3	Applying
8	Predict functional importance between chemostat and turbidostat continuiture system.	nuous (13)	BTL3	Applying
9	Analyze the influence of solutes and water activity on growth.	(13)	BTL4	Analyzing
10	State in detail the effect of pH on microbial growth.	(13)	BTL1	Remembering
11	List the different phases of microbial growth curve in a closed system.	(13)	BTL1	Remembering
12	Analyze the overall schema of metabolism.	(13)	BTL4	Analyzing
13	Describe the flow of carbon and energy in an ecosystem.	(13)	BTL1	Remembering
14	Categorize the laws of thermodynamics with appropriate interpretations	s. (13)	BTL4	Analyzing
15	Illustrate the importance of loss of viability caused in a irreparable cell.	(13)	BTL3	Applying
16	Break down the process of counting and identifying microorganisms in natural Environments.	(13)	BTL4	Analyzing
17	Apply the principle the cell-cell communication within microbial populations.	(13)	BTL3	Applying
	PART – C	1	0	
1	Model the growth rates in microorganisms by fitting exponential and curves. Use appropriate test data to simulate the results. From the sin data use conservative approach to Compute generation time microorganisms.	nulated	Green	Analyzing
2	Explain the structure and classification aspects of enzymes with emphalock and key model of enzyme function.	sis on (15)	BTL2	Understanding
3	Apply the following environmental factors on the growth rate (i) Temperature. (ii) Oxygen Concentration. (iii) Pressure.	(5) (5) (5)	BTL3	Applying
4	Illustrate different approaches and methods used to quantify bacterial growth.	(15)	BTL3	Applying
5	Enlist different media used for bacterial culture.	(15)	BTL1	Remembering

UNIT - IV CONTROL OF MICROORGANISMS

Physical and chemical control of microorganisms, Definition of sterilization, Dry and moist heat, Pasteurization, Tyndalization, Radiation, Ultrasonication, Filtration. Disinfections anitization, Antiseptics sterilants and fumigation. Mode of action and resistance to antibiotics, Clinically important microorganisms.

PART – A

Q. No	Questions	BT Level	Competence
1	Mention the significance of sterilization and sterilant.	BTL1	Remembering
2	Define disinfection and disinfectant.	BTL1	Remembering
3	Describe the principles of sanitization, antisepsis and antiseptic.	BTL1	Remembering
4	How one decides whether microorganisms are actually dead?	BTL3	Applying
5	Employ graphically the pattern of microbial death.	BTL3	Applying
6	Illustrate the significance of population size and composition.	BTL3	Applying
7	Recognize the concepts of chemotherapy, germicide, bactericide and bacteriostatic.	BTL1	Remembering
8	State the significance of heat in controlling microbial growth.	BTL1	Remembering
9	Outline the phenomenon of thermal death time.	BTL1	Remembering
10	Categorize the approximate conditions for moist heat killing.	BTL4	Analyzing
11	Point out the importance of dry heat sterilization	BTL4	Analyzing
12	Examine the importance of filtration in controlling microbial growth.	BTL4	Analyzing
13	Inspect the use of chemical agents in control of microbial growth.	BTL4	Analyzing
14	Indicate the importance of depth filters and membrane filters	BTL2	Understanding
15	Examine the advantages and disadvantages of ultraviolet light and ionizing Radiation.	BTL4	Analyzing
16	Analyze the significance of pasteurization in the past for death of infants.	BTL4	Analyzing
17	Interpret the need of phenol coefficient test.	BTL3	Applying
18	How can low temperatures be used to control microorganisms?	BTL2	Understanding
19	Why is boiling milk over prolonged periods not a desirable method for controlling spoilage and spread of milk-borne pathogens?	BTL2	Understanding
20	What three things one must do when operating an autoclave to help ensure success?	BTL2	Understanding
21	Infer the Dans Z values for some food-borne pathogens.	BTL2	Understanding
22	Express the longitudinal cross section of a typical autoclave showing some of its parts and the pathway of steam in a schematic representation.	BTL2	Understanding
23	Interpret the activity levels of selected germicides	BTL3	Applying
24	Prepare a schematic design of an EtO sterilizer.	BTL3	Applying

	PART – B			
1	Explain the various microbial control methods in an hierarchical tree structure.	(13)		Understanding
2	Complete the pattern of microbial death and fit the assessment with an exponential curve.	(13)	BTL3	Applying
3	Review the various aspects of use of heat as physical methods in contro microbial growth.	l of (13)		Understanding
4	Examine the significance of a membrane filters in sterilization process.	(13)		Remembering
5	Indicate the sterilization process used in automatically controlled autocl system.	ave (13)	BTL2	Understanding
6	Illustrate the following conditions influencing the effectiveness of antimicrobial agents (i) Concentration. (ii) Duration of exposure. (iii) Local environment.	(5) (4) (4)	BTL3	Applying
7	Review the working principle of UV treatment system for disinfection of water.	(13)	BTL2	Understanding
8	Interpret the use of following physical methods in control of microbial growth (i) Radiation. (ii) Low temperatures.	(7) (6)	BTL3	Applying
9	Apply the working principle of sterilization system during the onset of ionization radiation. State in detail the various universal precautions for microbiology	(13)	BTL3 BTL1	Applying Remembering
11	Laboratories. Recognize the functioning of an ethylene oxide sterilizer.	(13)		Remembering
12	Describe the operation of a biological safety cabinet.	(13)		Understanding
13 14	Name the use of phenolics and alcohols in control of microbial growth. Categorize the various tests involved to check the effectiveness of	(13)	BTL1 BTL4	Remembering Analyzing
15	antimicrobial agent. Examine the working principle of dry heat incineration.	(13)	BTL4	Analyzing
16	Inspect different steps in the mechanism of antimicrobial resistance to antibiotics.	(13)	BTL4	Analyzing
17	Analyze the use of heavy metals and sterilizing gases in control of microfrowth.	` ′	BTL4	Analyzing
	PART –C	,		1

1.	Explain various different systems that use heat as a physical method to control microbial growth. (15)	BTL2	Understanding
2.	Describe the structures of some frequently used disinfectants and antiseptics and explain their significance. (15)	BTL2	Understanding
3.	Recognize the use of following chemical agents in microbial control (i) Sterilizing gases. (ii) Chemotherapeutic agents. (5)	BTL1	Remembering
4.	Integrate various different systems that use filtration and radiation as a physical method to control microbial growth. (15)	BTL4	Analyzing
5.	Suppose hospital custodians have been assigned the task of cleaning all showerheads in patient rooms in order to prevent the spread of infectious disease. What two factors would have the greatest impact on the effectiveness of the disinfectant the custodians use? Explain what that impact would be. Also briefly describe about other conditions. (15)	BTL3	Applying

UNIT V - INDUSTRIAL MICROBIOLOGY

Microbes involved in preservation (Lactobacillus, bacteriocins), Spoilage of food and food borne pathogens (E.coli, S.aureus, Bacillus, Clostridium). Industrial use of microbes (production of penicillin, alcohol, vitamin B-12); Biogas; Bioremediation (oil spillage leaching of ores by microorganisms, pollution control); Biofertilizers, Biopesticides. Biosensors. Quality assurance — Quality control — Practice of cGMP — Schedule M — USFDA.

PART – A				
Q. No	Questions	BT Level	Competence	
1	What steps are usually taken to purify drinking water?	BTL1	Remembering	
2	Why is chlorination, although beneficial in terms of bacterial pathogen control, of environmental concern?	BTL1	Remembering	
3	Which important waterborne pathogens are not controlled reliably by chlorination?	BTL1	Remembering	
4	Interpret the significance of coagulation.	BTL3	Applying	
5	Show the various steps involved in water purification process.	BTL3	Applying	
6	Illustrate the various criteria required for "ideal" indicator organism.	BTL3	Applying	
7	Define coliform. How does this definition relate to presumptive, confirmed, and completed tests?	BTL1	Remembering	
8	In what type of environment is it better to use fecal enterococci rather than fecal coliforms as an indicator organism? Why?	BTL1	Remembering	
9	Why has the defined substrate test with ONPG and MUG been accepted as a test of drinking water quality?	BTL1	Remembering	
10	Differentiate the various metrics used to assess water quality.	BTL4	Analyzing	

11	Differentiate between coliforms and fecal coliforms in the laboratory.		BTL4	Analyzing
12	Outline the advantages and disadvantages of membrane filters for microbiological examinations of water.		BTL2	Understanding
13	Categorize the components that should limit the reactions in a BOD test.		BTL4	Analyzing
14	Identify the components in excess and limiting at the end of incubation per	iod	BTL2	Understanding
15	Inspect the factors that can lead to a nitrogen oxygen demand (NOD) in wa	ter.	BTL4	Analyzing
16	Analyze the parameters that can be monitored in a modern, large-scale industrial fermentation.		BTL4	Analyzing
17	Select the minerals can contribute to eutrophication		BTL2	Understanding
18	Review the critical limiting factors are used in the penicillin and streptomycin fermentations.		BTL2	Understanding
19	Describe the major uses for biopolymers and biosurfactants.	-	BTL2	Understanding
20	Use alternative definitions for the term biodegradation.	1	BTL3	Applying
21	Illustrate the components that should not limit reaction rates in a BOD test.	- 1	BTL3	Applying
22	What are biosensors and how do they detect substances?		BTL2	Understanding
23	Illustrate the significance of GMP Schedule M.		BTL3	Applying
24	Categorize the five main components of good manufacturing practice.		BTL4	Analyzing
	PART – B		(4)	
1	Indicate the various steps involved in water purification steps with a neat Diagram.	(13)	BTL2	Understanding
2	Enlist different stages involved in the multiple-tube fermentation test.	(13)	BTL1	Remembering
3	Enumerate the different metrics used to evaluate the quality of water.	(13)	BTL1	Remembering
4	State the significance of major steps in primary, secondary, and tertiary treatment of wastes.	(13)	BTL1	Remembering
5	Analyze the steps of organic matter processing that occur in anaerobic Digestion. Mention the significance acetogenesis step.	(13)	BTL4	Analyzing
6	Illustrate the following with a neat diagrams	(- /	BTL3	Applying
	(i) Constructed Wetland for Wastewater Treatment.	(7)		
	(ii) The Conventional Septic Tank Home Treatment System.	(6)		
7	Recognize the working principle of industrial stirred fermenters.	(13)	BTL2	Understanding
8	Infer the following products of industrial microbiology		BTL2	Understanding
	(i) Antibiotics.	(7)		
	(ii) Amino acids.	(6)		
9	Analyze the alternate methods for mass culture.	(13)	BTL4	Analyzing

10	Recognize the phenomenon of stimulating biodegradation.	(13)	BTL1	Remembering
11	Explain in detail the following		BTL2	Understanding
	(i) Metal Bioleaching.	(4)		
	(ii) Biopesticides.	(4)		
	(iii) Bio augmentation.	(5)		
12	Analyze the significance of a subsurface engineered bioremediation		BTL4	Analyzing
	System.	(13)		
13	Categorize the components used to design a biosensor used in industrial		BTL4	Analyzing
	microbiology.	(13)		
14	Inspect the phenomenon of phytoremediation with a neat diagram.	(13)	BTL4	Analyzing
15	Show the major organic acids produced by microbial processes.	(13)	BTL3	Applying
16	Interpret the major microbial products and processes of Interest in Indust	trial	BTL3	Applying
	Microbiology.	(13)		
17	Practice and use the growth of microorganisms in an industrial setting.	(13)	BTL3	Applying
	PART – C	0		
1	Integrate different sectors in a waste water treatment process to build a a		BTL4	Analyzing
	secondary sewage treatment.	(15)	DIL	7 mary zmg
2	Employ appropriate principle used for recombinant vaccine production.	(15)	BTL3	Applying
3	Recognize the Streptavidin-Biotin binding systems and enumerate the va	rious	BTL1	Remembering
	application of such systems.	(15)	111	
4	Inspect the mode of action of the Bacillus thuringiensis toxin.	(15)	BTL4	Analyzing
	Describe the use of microorg <mark>anisms</mark> in the fields of industrial microbiolog		BTL2	Understanding
	Biotechnology.	(15)	111	