## UNIVERSITY OF MUMBAL No. UG/ 104 of 2010

## CIRCULAR:

A reference is invited to the Ordinances, Regulations and syllabi relating to the Bachelor of Science (B.Sc.) degree course in Microbiology under the revised pattern vide this office Circular No. UG/151 of 2005 dated 27th April, 2005 and the Principals of the affiliated colleges in Science are hereby informed that the recommendation made by the Board of Studies in Microbiology at its meeting held on 17th February, 2010 has been accepted by the Academic Council at its meeting held on 3rd March, 2010 and vide Item No. 4.38 and that, in accordance therewith, the syllabus and pattern of question paper/s for the T.Y.B.Sc. examination in the subject of Microbiology is revised as per Appendix and that the same has been brought into force with effect from the academic year 2010-2011.

Mumbai-400 032 29th May, 2010

(L.R. Mane) Offg. Registrar

To,

The Principals of the affiliated colleges in Science.

#### A.C./ 4.38/ 03/03/2010

No.UG/104-A of 2010

MUMBAI-400 032

29th May, 2010

Copy forwarded with compliments for information to:-

- 1. The Dean, Faculty of Science,
- 2. The Chairperson, Board of Studies in Microbiology,
- 3. The Controller of Examinations,
- 4. The Co-ordinator, University Computerization Centre.

(D.N. Jadhav)

I/c. Deputy Registrar UG/PG Section

Copy to :-

The Director, Board of College and University Development, the Deputy Registrar (Eligibility and Migration Section), the Director, Board of College and University Development to the Vice-Chancellar, the Pro-Vice-Chancellor, the Registers of Students Welfare, the Personal Assistants to the Vice-Chancellar, the Pro-Vice-Chancellor, the Registrar and the Assistant Registrar, Administrative, Ratnagiri for information.

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# UNIVERSITY OF MUMBAI



Revised Syllabus and Paper Pattern at the T.Y.B.Sc. Examination M Microbiology

(With effect from the Academic year 2010-2011)

# REVISED SYLLABUS FOR TYB Sc Microbiology

## 2010-11 Onwards

Students opting for 6 Units of Microbiology (Major) at T Y B Sc level will study Papers I, II, IV of 100 marks each and four practicals based on these papers of 50 marks each.

Students opting for 3 Units of Microbiology at T Y B Sc level will study Papers I & II of 100 marks each and two practicals based on these papers of 50 marks each.

## Paper wise Units Summary

	Paper I	Paper II	Paper III	Paper IV
Title	Genetics, Bioinformatics, Molecular Biology & Virology	Medical microbiology and immunology	Microbial Biochemistry	Bioprocess Technology
Unit 1*	Classical genetics	Medical Microbiology	Solute transport Bioenergetics & Bioluminescence	Upstream Processing
Unit 2*	DNA replication, Mutation & Repair	Medical Microbiology Chemotherapy, Quality Assurance in Diagnostics	Methods & Carbohydrate metabolism	Downstream Processing
Unit 3*	Recombinant DNA technology	General immunology	Metabolism of Lipid, Protein, Nucleic acid & Aromatic compounds	Traditional Industrial Fermentations
Unit 4*	Virology ·	Immune system in Health & Disease	Metabolic regulation Photosynthesis & Inorganic metabolism	Advances in Bioprocess Technology

\* Note: Each Unit is of 30 lectures

## Paper I- Genetics, Bioinformatics, Molecular Biology & Virology (120 lectures)

Unit		
Offic	Topic	Lectures
	Classical genetics	° 30
11	DNA replication, Mutation & Repair	30
11	Recombinant DNA technology	30
IV	Virology	30

## I. Unit 1: - Classical Genetics (30)

#### 1. Branches of Genetics

- A. Transmission genetics
- B. Molecular genetics
- C. Population genetics
- D. Quantitative genetics.

#### 2. Model Organisms

[1]

- A. Characteristics of a model organism
- B. Examples of model organisms used in study
- C. Examples of studies undertaken using prokaryotic and eukaryotic model organisms.

#### Genetic Exchange

[18]

- A. Genetic analysis of bacteria
- B. Transformation
  - i. Introduction and History
  - ii. Types of transformation in prokaryotes--Natural transformation in Streptococcus pneumoniae, Haemophilus influenzae, and Bacillus subtilis
  - iii. Mapping of bacterial genes using transformation.
  - iv. Problems based on transformation.

#### C. Conjugation

- i. Discovery of conjugation in bacteria
- ii. Properties of F plasmid/Sex factor
- iii. The conjugation machinery
- iv. Hfr strains, their formation and mechanism of conjugation
- v. F' factor, origin and behavior of F' strains, Sexduction.
- vi. Mapping of bacterial genes using conjugation (Wolman and Jacob experiment).
- vii. Problems based on conjugation

#### D. Transduction

- i. Introduction and discovery
- ii. Generalised transduction
- iii. Use of Generalised transduction for mapping genes
- iv. Specialised transduction
- v. Problems based on transduction

	The second secon	
4	Plasmids A. Physical nature B. Detection and isolation of plasmids C. Plasmid incompatibility and Plasmid curing D. Cell to cell transfer of plasmids E. Types of plasmids i. Resistance Plasmids, ii. Plasmids encoding Toxins and other Virulence Characteristics iv. Degradative plasmids	[4]
5.	Transposable Elements in Prokaryotes  A. Insertion sequences B. Transposons i. Types ii. Structure and properties iii. Mechanism of transposition iv. Transposon mutagenesis C. Integrons	[4]
6.	Recombination in bacteria  A. General/Homologous recombination i. Molecular mechanism ii. Holliday model of recombination  B. Site –specific recombination  II. Unit 2: DNA replication, Mutation and Repair (30)	[2]
1.	The state of the s	lved in Initiation, mase, helicase, eins. , replicating the
2.	<ul> <li>Mutation</li> <li>A. Terminology: alleles, homozygous, heterozygous, genotype, phe mutation, Germline mutation, Gene mutation, Chromosome mutallag, hotspots and mutator genes</li> <li>B. Fluctuation test.</li> <li>C. Types of mutations: Point mutation, reverse mutation, support frameshift mutation, conditional lethal mutation, base pair substant transversion, missense mutation, nonsense mutation, silent mutation, pleiotropic mutations.</li> </ul>	ressor mutation,

- D. Causes of mutation:
  - i. Natural/spontaneous mutation--replication error, depurination, deamination.
  - ii. Induced mutation: principle and mechanism with illustrative diagrams for
    - a. Chemical mutagens- base analogues, nitrous acid, hydroxyl amine, intercalating agents and mechanism with illustrative diagrams of an intercalating agents and intercalating agents agents agents and intercalating agents agent intercalating agents and alkylating agents
    - b. Physical mutagen c. Biological mutagen
  - iii. Ames test.
  - iv. Detection of mutants
  - v. Complementation test.
- 3. DNA Repair
  - A. Mismatch repair,
  - B. Light repair
  - C. Repair of alkylation damage
  - D. Base excision repair
  - E. Nucleotide excision repair
  - F. SOS repair
  - G. Double strand break repair
  - H. Post replicative repair

## III. Unit 3: Recombinant DNA technology (30)

- Recombinant DNA technology
  - A. Basic steps in Gene Cloning.
  - B. Cutting and joining DNA molecules--Restriction and modification systems, restriction endonucleases, DNA ligases, Adaptors and linkers.
  - C. Vectors
    - i. Plasmids as cloning vectors. The plasmid vectors, pBR322 vector
    - Cloning genes into pBR322
    - iii. Phage as cloning vectors, cloning genes into phage vector
    - iv. Cosmids
    - v. Shuttle vectors
    - vi. Expression vectors
  - D. PCR- basic PCR and different types of PCR (Reverse transcriptase PCR, Real time quantitative PCR and Long accurate PCR)
  - E. Methods of transformation.
  - F. Construction of genomic and cDNA libraries.
  - G. Basic techniques
    - i. Southern, Northern and Western blotting.
    - ii. Preparation of radioactive and non-radioactive DNA probes
    - iii. Autoradiography.
  - H. Screening and selection methods for identification and isolation of recombinant cells.
  - Applications of recombinant DNA technology.

[5]

[22]

		[8]
2.	Bioinformatics  A. Introduction	•
,	Definition, aims, tasks and applications of Bioinformatics.     Database, tools and their uses	
	a. Importance, Types and classification of databases     b. Nucleic acid sequence databases- EMBL, DDBJ, GenBa     Ensembl and specialized Genomic resources.	nk, GSDB,
	c. Protein sequence databases-PIR, SWISS-PROT, TrEM 3D.Protein structure databases-SCOP, CATH, PROSITE, P	RINTS and
	BLOCKS. KEGG.  B. Brief introduction to Transcriptome, Metabolomics, Pharmac Phylogenetic analysis, Phylogenetic tree, Annotation, QSAR (quantitativativity relationship), Docking algorithms  C. Sequence alignment—global v/s local alignment, FASTA, BLAST.  D. Genomics—structural, functional and comparative genomics.  E. Proteomics—structural and functional proteomics.	ive structure
	IV. Unit 4: Virology (30)	
	A Contracture	[4]
	Viral architecture-  A. Capsid, viral genome and envelope  B. Structure of TMV, T4, Influenza virus, HIV.	
	B. Silucture of thirty of	[1]
2.	Viral classification	THE PROPERTY.
3.	The viral replication cycle- attachment, penetration, uncoating, types of v their replication, assembly, maturation and release. Life cycle of T4 Influenza Virus and HIV in detail	phage, TMV, [8]
		[3]
4.	Cultivation of viruses- cell culture techniques, embryonated egg, laboratory animals	
	to a provide a figure particles	٠ [7]
5.	Visualization and enumeration of virus particles  A. Measurement of infectious units	
	i. Plaque assay	
	ii. Fluorescent focus assay	
	oiii. Infectious center assay	
c	Fadraint dilution assay	
	B. Measurement of virus particles and their components	
	: Flastron microscony	
	" At-mis force microscopy	•
	::: Heamagalutination	ANTO
	iv Measurement of VITAL ENZYME activity.	
	C. Fate of the cells following virus infection	
6.	Regulation of lytic and lysogenic pathway of lambda phage	[3]
7.	Role of viruses in cancer	[2]
		[2]
8.	Prions and viroids	ردا

## ext books

1. Peter J. Russell (2006), "Genetics-A molecular approach", 2<sup>nd</sup> ed.

2. Benjamin A. Pierce (2008), "Genetics a conceptual approach", 3<sup>rd</sup> ed., W. H. Freeman and company.

3. R. H. Tamarin, (2004), "Principles of genetics", Tata McGraw Hill.

- 4. D. Nelson and M.Cox, (2005), "Lehninger's Principles of biochemistry", 4th ed., Macmillan worth Publishers.
- 5. M.Madigan, J.Martinko, J.Parkar, (2009), "Brock Biology of microorganisms", 12th ed., Pearson Education International.

6. Fairbanks and Anderson, (1999), "Genetics", Wadsworth Publishing Company.

- 7. Prescott, Harley and Klein, "Microbiology", 7th edition Mc Graw Hill international edition.

  8. Edward Wagner and Martinez Hewlett, (2005) "Basic Virology", 2<sup>nd</sup> edition, Blackwell
- Publishing
- g. Teri Shors, (2009), "Understanding viruses", Jones and Bartlett publishers.

10. S.Ignacimuthu, (2005), "Basic Bioinformatics", Narosa publishing house.

- 11. Robert Weaver, (), "Molecular biology", , 3rd edn. Mc Graw Hill international edition.
- 12. Nancy Trun and Janine Trempy, (2004), "Fundamental bacterial genetics", Blackwell Publishing
- 13. Primrose and Twyman, (2006), "Principles of gene manipulation and genomics", 7th ed, Wiley-Blackwell
- 14. Arthur Lesk, (2009), "Introduction to Bioinformatics", 3rd Edition, Oxford University Press

### Reference books:

1. Flint, Enquist, Racanillo and Skalka, "Principles of virology", 2<sup>nd</sup> edn. ASM press.

- 2. T. K. Attwood & D. J. Parry-Smith, (2003), "Introduction to bioinformatics", Pearson
- 3. Benjamin Lewin, (), "Genes IX", , Jones and Bartlett publishers.

4. JD Watson, "Molecular biology of the gene", , 5th edn.

5. Snustad, Simmons, "Principles of genetics", 3rd edn. John Wiley & sons, Inc.

#### PRACTICALS BASED ON PAPER 1

Enrichment of coliphages, phage assay (pilot & proper).

2. UV survival curve - determination of exposure time leading to 90% reduction

Isolation of mutants using UV mutagenesis

4. Replica plate technique for selection & characterization of mutants - auxotroph & antibiotic resistant

Isolation and detection of plasmid DNA.

Preparation of competent cells and transformation

Restriction analysis.

8. Isolation of genomic DNA of E. coli

9. PCR (Demo)

- 10. Western Blot.(Demo)
- Genetics problems.
- 12. Bioinformatics practical

A. Off Line Practical

- ii. Installation of representative software for off line use SPDBV and Bioedit
- iii. Visualizing and manipulating Protein structure database files using SPDBV
- iv. Sequence Alignment, dot plot, phylogenetic tree building exercise using Bioedit

- B. On Line Practical
  - i. Visiting NCBI and EMBL websites & list services available, software tools ii. Visiting & exploring various databases mentioned in syllabus and
  - - b. Using BLAST and FASTA for sequence analysis c. Fish out homologs for given specific sequences (by teacher – decide sequence of some relevance to their syllabus and related to some biological problem e.g. evolution of a specific protein in bacteria, predicting function of unknown protein from a new organism based on its homology – list can be really long and should be generated with the help of students while teaching
    - topics in genetics, biochemistry, bioinformatics through out the year illustrating power of informatics tools in biology)
    - d. Understand every item mentioned in the report generated, its significance and use in interpretation of results as well as limitations of the results.
- 3. Animal cell culture (demo)

# PAPER II: MEDICAL MICROBIOLOGY AND IMMUNOLOGY (120 Lectures)

Unit	Brief Topic Description	T
		No of Lectures
1	Medical Microbiology	30
2	Medical Microbiology, Chemotherapy, Quality Assurance in Diagnostics	30
3	General Immunology	30
4	Immune system in Health & Disease	30
Dag	Total	120

## I. UNIT1: MEDICAL MICROBIOLOGY (30)

All infections are to be covered with respect to all details with emphasis on Etiology, Transmission, Pathogenesis, Clinical Manifestations, Lab Diagnosis, Prophylaxis, and Treatment.

Respiratory tract Infections

[11]

A. Upper respiratory tract:

i. Streptococcal Pharyngitis, Diphtheria

- ii. Common Cold , Oral Candidiasis , Measles- Rubeola, Rubella, Mumps , Chicken pox, Shingles
- B. Lower Respiratory tract:
  - i Tuberculosis Influenza
  - Bacterial pneumonia. Whooping cough
- Urinary Tract Infections Pathogens & Factors Involved

[3]

Gastro – Intestinal Infections

[9]

- A. Infectious diseases caused by
  - i. Salmonella, Shigella, Vibrio
    - ii. E.coli, Helicobacter pylori, Campylobacter, Rota virus, Hepatitis A, E. histolytica
- B. Food Poisoning: Staphylococcal, Botulism

. Central Nervous System Infections

[7]

- i. Tetanus, Polio, Rabies
- ii. Meningitis: viral, bacterial -Meningococcal, Pneumococcal and Haemophilus

## II UNIT 2: MEDICAL MICROBIOLOGY, CHEMOTHERAPY, QUALITY ASSURANCE (30)

Μ	<ul> <li>All infections are to be covered with respect to all details with emphasi Etiology, Transmission, Pathogenesis, Clinical Manifestations, Lab Dis Prophylaxis, and Treatment</li> </ul>	[15] s on agnosis,
- D	Sexually transmitted Infections  i. HIV infection, Syphilis  ii. Gonorrhea, Herpes, Hepatitis B  Skin Infections	
	Pyogenic Staphylococcal, Streptococcal, Leprosy, Malaria Candidiasis, Dermatophytosis, Pseudomonas Emerging Infections - SARS, H1N1, Avian flu, Leptospirosis, Dengue	
BOD	hemotherapy i. History and Development of Chemotherapy ii. General Properties of antimicrobial agents iii. Attributes of an ideal antimicrobial agent Drug Resistance: Origin, Mechanisms and Transmission Selection & Testing Principal Groups of Antibacterial Agents and Mechanism of Action i. Cell Wall Inhibitors ii. Inhibitors of Protein Synthesis iii. Inhibitors of Nucleic Acid Synthesis iv. Cell Membrane Disruptors v. Antimetabolites Anti-mycobacterial, Antifungal, Antiviral - Tabulation of Examples Quality Assurance in Diagnostics - Concepts of Quality Assurance in Diagn	[12]
, c	Quality Assurance in Diagnostics - Concepts of the [3]	
	III UNIT 3: GENERAL IMMUNOLOGY (30)	[1]
. (	Cells of the immune system- T-cells, B-cells, NK-cells	[2]
:. C	Cytokines  A. Properties and Functions  B. Cytokines secreted by Th1 andTh2 cells  Cytokines secreted by Th1 andTh2 cells  Antigen Presenting Cells - Antigen presentation and processing pathways  and Endocytic pathway)	(Cytosolic
		[4]
ł. N	MHC complex and MHC Molecules  A. Organization of MHC genes  B. Structure of class I and class II molecules  C. Polymorphism and Polygenism  C. T cell antigen receptors and MHC molecules.  Tests for MHC specificity.	

## II UNIT 2: MEDICAL MICROBIOLOGY, CHEMOTHERAPY, QUALITY ASSURANCE (30)

A. Sexually transmitted Infections	
<ul><li>i. HIV infection, Syphilis</li><li>ii. Gonorrhea, Herpes, Hepatitis B</li><li>B. Skin Infections</li></ul>	
Pyogenic Staphylococcal, Streptococcal, Leprosy, Malaria Candidiasis, Dermatophytosis, Pseudomonas  C. Emerging Infections - SARS, H1N1, Avian flu, Leptospirosis, Dengue	9
<ol> <li>Chemotherapy         <ol> <li>Basics of Chemotherapy</li> <li>History and Development of Chemotherapy</li> <li>General Properties of antimicrobial agents</li> <li>Attributes of an ideal antimicrobial agent</li> </ol> </li> <li>Drug Resistance: Origin, Mechanisms and Transmission</li> <li>Selection &amp; Testing</li> <li>Principal Groups of Antibacterial Agents and Mechanism of Action</li> <ol> <li>Cell Wall Inhibitors</li> <li>Inhibitors of Protein Synthesis</li> <li>Inhibitors of Nucleic Acid Synthesis</li> <li>Cell Membrane Disruptors</li> <li>Antimetabolites</li> </ol> <li>Anti-mycobacterial, Antifungal, Antiviral - Tabulation of Examples</li> <li>Quality Assurance in Diagnostics - Concepts of Quality Assurance in [3]</li> </ol>	[12] Diagnostics
III UNIT 3: GENERAL IMMUNOLOGY (30)	
1. Cells of the immune system- T-cells, B-cells, NK-cells	[1]
<ul> <li>2. Cytokines</li> <li>A. Properties and Functions</li> <li>B. Cytokines secreted by Th1 andTh2 cells</li> </ul>	[2]
<ol> <li>Antigen Presenting Cells - Antigen presentation and processing path and Endocytic pathway)</li> </ol>	nways, (Cytosolic [2]
<ul> <li>4. MHC complex and MHC Molecules</li> <li>A. Organization of MHC genes</li> <li>B. Structure of class I and class II molecules</li> <li>C. Polymorphism and Polygenism</li> <li>D. T cell antigen receptors and MHC molecules.</li> <li>E. Tests for MHC specificity.</li> </ul>	[4]

	<ul> <li>7 cells</li> <li>A. Receptors, structure and organization</li> <li>B. T cell development and maturation, positive and negative selection</li> </ul>	[7]
,	C. T cell activation TCR coupled signaling pathway Costimulatory signals Superantigen induced T cell activation.	
	<ul> <li>D. T cell differentiation</li> <li>i. Generation of effector and memory cells.</li> <li>ii. Cell death and T cell population.</li> <li>iii. Functions of peripheral αβ and γδ cells</li> </ul>	
6	Receptorsstructure & organization  B cell development and maturation	[5]
	B cell activation & differentiation  i. Thymus dependent and independent antigens.  ii. B cell activating signals  iii. Role of Th cells in humoral response, formation of T-B conjugates CI	040/CD40L
-	Interaction, The cell cytokine signals.	[3]
7.	<ul> <li>A. Induction of Humoral response, Primary and secondary responses,</li> <li>B. Germinal centers and antigen induced B cell differentiation</li> <li>C. Outline of Organization and expression of Ig genes, gene rearrangement</li> <li>D. Affinity maturation and somatic hyper mutation, Ig diversity, class switchi</li> <li>E. Generation of plasma cells and memory cells, synthesis, assembly and simmunoglobulins.</li> <li>F. Evaluation of humoral response.</li> </ul>	secretion of
8.	Cell mediated effector response  A. Generation and target destruction by Cytotoxic T cells.  B. Killing mechanism of NK cells.  C. Antibody dependent cell cytotoxicity (ADCC)  D. Experimental assessment of CM cytotoxicity.	[2]
9.	Complement system  A. Complement components and notations  B. Complement activation (classical pathway, Alternate pathway, Lectin particle C. Biological consequences of complement activation.  D. Regulation of complement pathways.	[4] thway)
	IV UNIT 4: IMMUNE SYSTEM IN HEALTH AND DISEASE (30)	Cerollar
	Antigen- Antibody reactions Precipitation, agglutination, passive agglutination, agglutination inhibition, C Fixation, Radioimmunoassays (RIA), Enzyme immunoassays (EIA), Immunofluorescence, Flow cytometry, western blot technique, immunoelect microscopy, Toxin antitoxin assays.	[5] omplement ron
2.	Monoclonal antibodies - Preparation, applications, Engineered antibodies.	[2]

		· saines	[10]
3	3. ) I	Vaccines A. Active and passive immunization B. Types of vaccines - Killed and attenuated vaccines, Whole organisms	m vaccines,
		purified macromolecules as vaccines, recombinant vector vaccines	S,DIVA V
		- use of adjuvants in vaccine	
		New vaccine strategies	
		Ideal vaccine Route of vaccine administration, Vaccination schedule, Failures in	vaccination.
		A STATE OF THE PARTY OF THE PAR	[4]
4	. !	mmunohematology  A. Human blood group systems, ABO, secretors and non secretors, E group. Rhesus system and list of other blood group systems.	3ombay Blood
		group. Wresde system and list of other blood group systems.	
	(	Blood Transfusion, Major and Minor Cross matching, transfusion re	eactions
		Hypersensitivity –	[4]
5		A Coombs and Gells classification	
	í	3. Type I to Type IV hypersensitivity, Mechanism and manifestation.	
			[2]
6	i. /	Autoimmunity  A. Definition of immune tolerance,	
	- 1	n Immune suppression and autoimmunity	
		c Spectrum of autoimmune diseases.	
		D. Mechanism and treatment of autoimmune diseases.	
		Transplantation immunology	[3]
7		A leadinglodical basis of draft rejection.	
	-	Types of graft rejection, Clinical manifestation of graft rejection,	
		C. General and specific immunosuppresive therapy	
Text bo		Ananthanaravan, R. & Pallikel, OK Jayaram, (2000), Termina	icrobiology", 8th
		r Pitam Universities Press (India) PVL LIU	14
2		Order Mine et al. (2004) "Medical Microbiology", 3rd Edition Mosby	
3		Prescott, Harley, Klein, "Microbiology", 6th Edition McGraw Hill WC Winn Jr., S D Allen, W M Janda, Er W Koneman, P C Schrecker	nberger, G
4	١. '	W C Winn Jr., S D Allen, W W Janda, El W Konsman, W W Procop, G L Woods, "Koneman's Color Atlas and Textbook of Diag	gnostic
5			ett Publishers
6	S. 1	Teri Shors Jones, (2009), "Understanding Viruses, edition of the Edition." Richard A. Goldsby, Janis Kuby, "Immunology", 6th and 7th Edition.	vv. 11. i recilian an
-	. (	company. Fahim Halim Khan, (2009), "The elements of Immunology",. Pearson	Education.
7	′. l	Fahim Halim Khan, (2009), The elements of mindulology probable and Fundamental", 2nd Pathak, S., Palan U, "Immunology Essential and Fundamental", 2nd	Edition. Capital
8	),   		
_	۱ ۱	Publishing company les B. Tizard "Immunology. An Introduction", 4 <sup>th</sup> - Edition, Saunders	college publishing

## Practical Syllabus Based on Paper II

1. Schematic /diagrammatic representation of each system as per the theory syllabus (Respiratory, Urinary, Gastro-intestinal, Central Nervous Systems, Skin)

2. "Diagnostic Cycle" of any one infection of each of the above systems (viz., in upper

respiratory tract: Pharyngitis)

- 3. Samples of various forms/procedures used for diagnostic tests e.g. Request forms, QC slips, for samples, reagents, stains, media, equipment validation, Test-reports, (Results, Panic report, alert report) to be drawn or attached in the journal. 4. Tabulation of:
  - A. Types of samples, containers, specimens, with reference to the symptoms/infections.

B. Transport media with reference to samples/suspected pathogen.

C. Collection and Processing of samples in various infections.

D. Primary isolation of suspected pathogens in different infections with reference to pathological samples

E. Rapid tests for identification of pathogens e.g. oxidase, catalase, stainings (Acid

fast, Metachromatic granules, Capsule), Germ tube formation.

- F. Minimum biochemical media for identification of the pathogens listed in the syllabus i.e. S. aureus, S. pyogenes, Corynebacterium diphtheriae, E. coli, Klebsiella spp., Salmonella spp., Shigella spp., Vibrio spp., Proteus spp., Pseudomonas spp.
- G. List of samples to be used with the above:

i. URT: Nasal swab, pus.

ii. GIT: Faeces, Rectal swab.

iii. UTI: Urine,

iv. Bacteraemia: Blood.

v. CNS: CSF.

5. Case study and problem solving for identification of the pathogen and antibiotic sensitivity with reference to each of the infections (Include approach writing, suspected organisms, requirements for the identification tests and their justification rapid tests, AST reports.)

6. Perform quality control tests of media, reagents, strains and equipment used in the

7. Kirby-Bauer method and Stokes method for AST.

8. Synergistic activity of antibiotics.

E test.(Demonstration)

- 10. Agar cup method for determination of antibiotic levels in body fluids (serum)
- 11. Detection of β-lactamase producer by Acidometric/Iodometric method
- 12. Differential Blood Count, Blood Grouping, Direct & Reverse Typing

13. Determination of Isoagglutinin titre

14. Coombs test - direct & indirect method

Compatibility test – cross matching.

16. Preparation of Typhoid vaccine and sterility checking

17. Antigen – Antibody Reactions: Agglutination – Widal ( Demonstration); VDRL Qualitative and Quantitative (Demonstration); Immuno diffusion - Single- Oudin's; Double -Ouchterlony; SRID

18. Separation of lymphocytes and staining (Demonstration)

19. Pregnancy test - ELISA (Demonstration)

20. Rheumatoid arthritis test (Demonstration)

## Paper III - Microbial Biochemistry . (120 Lectures)

Unit	T	
	A. Solute transport	Lectures
	B. Bioenergetics	30
11	A) Methods for studying metabolism     B) Čarbohydrate metabolism	30
III	Metabolism of Lipid, Protein Nucleic acid & Aromatic compounds	
	A. Metabolic result is	30
IV	A. Metabolic regulation     B. Photosynthesis     C. Inorganic metabolism	30
,	Total	120

# I. Unit 1: Solute Transport, Bioenergetics & Bioluminescence (30)

#### Solute transport:

- A. Methods of studying solute transport
- B. Role of membrane in solute transport
- C. Mechanism for uptake of solutes
  - Passive diffusion
  - ii. Facilitated diffusion
  - iii. Active transport
    - a. Primary active transport Histidine uptake model (Shock sensitive system), Maltose uptake
    - b. Secondary active transport- Uniport, Antiport, Symport
    - Active transport linked to phosphate bond energy
  - iv. Group translocation
  - v. Other examples of solute transport
    - a. Iron transport : A special problem
    - b. Transport through outer membrane (Porin and special transport protein)
    - c. Assembly of proteins in to membranes and protein export

#### Bioenergetics

A. Electron transport chain: components, complexes and functions

- Mitochondrial ETC
- Prokaryotic ETC
  - a. Organotroph E. Coli
  - b. Lithotroph Nitrosomonas& Nitrobater (Only schematic)
- B. Oxidative phosphorylation: by Chemiosmotic coupling hypothesis
- C. Structure and mechanism of ATP synthase
  - Structure of
    - a. bacterial ATP synthase
    - b. Mitochodrial ATP synthase
  - ii. Mechanism Rotational catalysis

[13]

[15]

Page 15 of 23 D. Other modes of generation of electrochemical energy i. Oxalate formate exchange ii. Decarboxylases dependent ion transport
iii. End product efflux
iv. iv. ATP hydrolysis E. Calculation of energetics of glycolysis, TCA and Beta oxidation of fatty acid (palmitic acid) - balance sheet to be given with efficiency calculation F. Bacteriorhodopsin: Photo cycle and significance

3. Bioluminescence - Introduction, ETC, Significance / Application

[2]

# Unit II – Methods & Carbohydrate Metabolism (30)

Methods of studying metabolism

- A. Use of biochemical mutants, Isotopic labeling (Including radiorespirometry with reference to EMP&ED), sequential induction technique
- B. Modern methods based on biochemical genetics, molecular biological and computational techniques, concept of metabolome and its uses in the study of
- 2. Metabolism of Carbohydrates

[28]

- A. Catabolism
  - i., Breakdown of polysaccharides glycogen, starch, cellulose
  - ii. Breakdown of oligosaccharides lactose, maltose, sucrose
  - iii. Utilization of monosaccharides fructose, galactose, mannose
  - iv. Major pathways:
    - a. Glycolysis (EMP)
    - b. HMP Shunt
    - c. ED pathway
    - d. Phosphoketolase pathway (pentose phosphoketolase), hexose Bifidobacterium pathway
    - e. Other modes of fermentations in microorganisms: alcohol, mixed acid, butanediol, butyric acid, butanol-acetone, propionic acid (randomizing & nonrandomizing pathway)
    - f. Citric acid cycle, anaplerotic reactions, glyoxylate bypass, Incomplete TCA in anaerobic bacteria
    - g. Amphibolic pathways: role of EMP and TCA cycle
- B. Anabolism
  - i. Gluconeogenesis
  - ii. Biosynthesis of glycogen
  - iii. Biosynthesis of Peptidoglycan and Lipopolysaccharide

## III Unit 3: Lipid, Proteins, Nucleic Acids & Aromatic Compound Metabolism (30)

Lipid metabolism

[9]

- A. Catabolism
  - i. Oxidation of saturated fatty acid-β oxidation pathway
  - ii. Oxidation of propionic acid
  - iii. Oxidation of saturated aliphatic hydrocarbon (n-alkane)-Omega oxidation pathway- Pathway in Corynebacterium and yeast, Pathway in Pseudomonas
  - iv. Degradation of poly beta Hydroxyl butyrate

#### B. Anabolism

- Biosynthesis of straight chain even carbon saturated fatty acid (palmitic acid)
- protein metabolism

#### A. Catabolism

[10]

- i. Enzymatic degradation of proteins
- ii. Metabolic fate of amino acids (schematic only) iii. Metabolism of single amino acids –Deamination, decarboxylation, and
- iv. Fermentation of single amino acids
- v. Glutamic acid, Alanine by Clostridium propionicum
- vi. Fermentation of pair of amino acids (Stickland reaction)
- B. Anabolism
  - Schematic representation of amino acid families
  - ii. Synthesis of amino acids of Serine family- Examples serine, cysteine, glycine

### 3. Nucleic acid metabolism

[8]

- A. Catabolism
  - i. Degradation of purine and pyrimidine nucleotides up to uric acid formation
  - ii. Recycling of purines and pyrimidines nucleotides by salvage pathway-
- B. Anabolism Synthesis of ribonucleotides and deoxyribonucleotides
- 4. Metabolism of aromatic compounds

- A. Schematic representation for conversion of various aromatic compounds to catechol and protocatechuic acid
- B. Catabolism of catechol and protocatechuic acid by ortho and Meta cleavage

### IV Unit 4: Metabolic Regulation (30)

Metabolic regulation

[15]

- A. Cellular control mechanism acting at various levels of metabolism
- B. Allosteric proteins Role as enzymes and regulatory proteins
- C. RNA's as regulatory molecules
- D. Regulation of gene expression
  - Regulation in bacteria Operon model criteria for negative / positive types and inducible / repressible types
  - Regulation of enzyme synthesis (Enzyme induction/repression)

Mechanism of control of transcription

- a. By DNA- Binding proteins
  - ⇒ Lac operon (Negative control of enzyme induction)
  - ⇒ Ara operon (Positive control of enzyme induction)
  - ⇒ Catabolite repression
- b. Enzyme Repression in Branched Biosynthetic Pathways
- c. By Attenuation
  - ⇒ Trp operon (End-Product Repression)
- d. By Multiple Sigma Factors

- E. Regulation of enzyme activity (Enzyme inhibition/activation) i Mechanism of End-Product Inhibition
  - a. Patterns of regulation-- End-Product Inhibition in branched pathways, Isofunctional enzymes, concerted feedback Inhibition, sequential feedback Inhibition, sequential feedback inhibition, Cumulative Feedback Inhibition, sequential too and inhibition
  - b. Covalent modification of regulatory of enzymes Glutamate synthetase

c. Regulation by proteolytic cleavage

F. Regulation of EMP, TCA and ortho cleavage pathway of aromatic compounds

2. Prokaryotic photosynthesis

A. The phototrophic prokaryotes

i. Oxygenic phototrophs

ii. Anoxygenic phototrophs

B. Photosynthetic pigments and photosynthetic apparatus

C. Light reactions of purple photosynthetic bacteria, green sulphur bacteria and cyanobacteria

- D. Dark reaction: Calvin Benson cycle and reductive TCA
- 3. Inorganic metabolism

[8]

[7]

- A. Assimilatory pathways
  - i. Assimilation of nitrate
  - ii. Ammonia fixation
  - iii. Biological nitrogen fixation Mechanism for N2 fixation and protection of nitrogenase
  - iv. Assimilation of sulphate

B. Dissimilatory pathways

i. Nitrate as an electron acceptor (Denitrification in Paracoccus denitrificans)

ii. Sulphate as an electron acceptor

C. Lithotrophy - Enlist organisms and products formed during oxidation of Hydrogen, carbon monoxide, Ammonia, Nitrite, Sulphur, Iron.

#### Practical based on Paper III

- 1. Isolation of Phenol degraders and estimation of residual phenol by 4-amino antipyrene
- 2. Estimation of β-galactodidase activity in induced and non- induced cells of Escherichia coli
- 3. To study catabolite repression by diauxic growth curve

4. Protein estimation by Lowry's method

5. Isolation of bioluminiscent bacteria from fish

6. Study of biochemical pathway and study of end products of enzymes in characterization of micro-organisms.

A. Detection of lysine decarboxylase enzyme.

B. Oxidative and fermentative utilization of glucose by microbes.

C. Phosphatase activity detection-qualitative and quantitative.

D. Detection of Penicillinase activity.

E. Detection of homo and mixed acid fermentation.

Estimation of uric acid.

8. Isolation of mitochondria and assay for ETC activity

## Text books

- 1. Stanier.R.Y., Ingrahm, J.L., Wheelis, M.L., Painter, R.R., (1987) General Microbiology, 5<sup>th</sup>
- 2. Conn , Stmpf, P. K., Bruening, G. R. H.(1987) Outlines of Biochemistry, 5th edition, John
- 3. Gottschalk,G., (1985), Bacterial Metabolism, 2<sup>nd</sup> edition, Springer Verlag Gottschait, C., (1995), The Physiology and Biochemistry of Prokaryotes, 2<sup>nd</sup> edition, Oxford
- 5. Nelson, D, Cox, M,(2005), Lehninger Principles of biochemistry,4<sup>th</sup> edition, W. H.

## Reference books

- 1. Voet, D & Voet, J. G., (2004), Biochemistry, 3rd edition, John Wiley& Sons Inc.
- 2. Zubey, G. L (1996), Biochemistry, 4<sup>th</sup> edition, Wm. C. Brown publishers
- 3. Zubey, G. L (1996), Principles of Biochemistry, Wm. C. Brown publishers

## PAPER IV: BIOPROCESS TECHNOLOGY (120 Lectures)

Unit	Brief Topic Description	
1	Linette en D	No of Lectures
'	Opstream Processing	30
- 11	Downstream Processing	
111	Traditional Industrial Fermentations	30
- "	A discontinuity of the second	30
IV	Advances in Bioprocess Technology	30
-	Total	
	, otal	120

## I. UNIT 1: UPSTREAM PROCESSING (30)

1. Industrial Strains

- A. Strain improvement (One example of each method of strain improvement for primary and secondary metabolite)
- B. Preservation of industrial strains

2. Fermentation Media Design

[3]

- A. Buffers, precursors, steering agents, inducers, inhibitors, antifoam agents, trace elements, Animal cell culture media
- B. Media Optimizations general principles
- Fermentation Equipments

[10]

- A. Construction material
- B. Scale of operation (Lab, Bench scale, pilot plant, production level)
- C. Mode of operation (Batch, fed-batch, semi-continuous, continuous, SSF)
  - D. Power Input for mixing (mechanical, hydrodynamic and pneumatic)
  - E. Types of fermentors typical constructional features and their importance in the specific processes, brief review of other supporting services or equipments used for process operations
    - i. Mechanical Waldhof fermenter, rotating disc fermenter, trickling generator,
    - ii. Hydrodynamic- deep-jet fermenter
    - iii. Pneumatic air-lift fermenter, bubble-cap fermenter, cylindroconical vessels, acetator, cavitator.
    - Animal cell culture reactors.
    - v. Photo-bioreactor, tower and packed tower fermenters, cyclone column.
- 4. Fermentation Process Operations

[10]

- F. Aseptic operation and containment
- A. Sterilization & maintenance of aseptic conditions vessels, medium, additives, air
- B. Aseptic transfer of inoculum
- C. Process parameter monitoring and control
  - i. Temperature, flow, pressure, dissolved oxygen, foam, inlet and exit gases, pH
  - ii. Control systems manual and automatic (only list)

# II UNIT 2 - DOWNSTREAM PROCESSING (30)

1.	Fermentation Product Recovery  A. Criteria for choice of recovery process  B. Biomass separation from fermentation media  i. Precipitation  ii. Filtration, filter aids, plate frame and rotary vacuum filters  iii. Centrifugation - Cell aggregation and flocculation, types of centrifuge  C. Cell Disruption for intracellular products  D. Solvent extraction and recovery  E. Chromatography  F. Membrane processes  G. Drying  H. Crystallization  I. Whole broth processing	[10]
2.	Industrial Effluent Treatment A. Distillery Effluents B. Pharmaceutical Effluents	[3]
3.	Fermentation Economics - Isolation, strain improvement, market potential, media, air sterilization, temperature control, aeration and agitation, recoverecycling, effluent treatment.  [2]	equipment, ry, water
4.		[5]
	DefinitionsGMP, QA, QC QC of raw materials, in-process items, finished products, packaging materials	riale labels
	Sterility assurance and testing	ilais, labels
	Microbiological Assays	
5.	Bioinstrumentation - Principles, working and applications of A. Spectroscopic techniques B. Electron spin resonance (ESR) spectroscopy C. Nuclear magnetic resonance (NMR) spectroscopy D. Circular dichroism (CD) spectroscopy E. Mass spectroscopy F. Spectrophotometry (U.V., Visible, I. R) G. Fluorimetry H. Flame photometry I. Radioisotopes and autoradiography	[10]
	III UNIT 3: TRADITIONAL INDUSTRIAL FERMENTATIONS (3	0)
3	<ol> <li>Beer</li> <li>Wine</li> <li>Alcohol from molasses</li> <li>Vinegar (acetator</li> <li>Penicillins and semisynthetic penicillins</li> <li>Streptomycin</li> <li>Vitamin B12 from Propionibacterium</li> <li>Glutamic Acid (direct)</li> <li>Baker's and Brewer's yeast</li> </ol>	[2] [2] [1] [1] [3] [1] [2] [1]

12 13 14 15 16	Citric acid - Stationary culture Mushrooms (Agaricus bisporus) Biotransformation of Steroids (List of organisms and steroids transformed) Vaccines -General Manufacturing aspects and quality control Ergot Alkaloids Probiotic foods & neutraceuticals Amylase enzyme production (Solid state fermentation) Microbial polysaccharides	[2] [2] [1] [3] [2] [2] [1]
	IV. UNIT 4: ADVANCES IN BIOPROCESSES (30)	
1.	Enzyme Technology  A. Enzyme Immobilization methods  B. Applications in therapeutic uses, Analytical uses and Industrial uses	[6]
2.	Animal Cell Lines  Methods of cultivation and establishment of cell lines  Large scale cultivation procedures  Applications in production of tPA, Blood facor viii and erythropoietin	[6]
3.	Plant Tissue Culture Methods of cultivation of organ culture, callus culture and cell suspension culture Application in  Agriculture (Disease resistant plants, virus free plants) Horticulture (Micropropagation) Industry (secondary metabolites production) Transgenic plant (Insect resistant plants)	[6] ulture
4.	Commercial Products from Recombinant Microorganisms - Indigo, Bioflavor Biopolymer, Polyhydroxyalkanoate, Rubber, Recombinant proteins of high v	urs, Melanin, ⁄alue [5]
5.	Synthesis of Nanomaterials by Biological methods Applications in biotechnology and medical field	[2]
6.	Intellectual Property Rights  A. Introduction to IPR – What is intellectual property? Genesis of IPR (WIFTRIPs)  B. Types of intellectual property – a. Patents b. Copyright c. Trademark d. Trade secret e. Plant varieties (C. Patents – a. Patent system terminologies b. Categories of patents c. Preparation of patent i. Criteria for patenting ii. Patent specification – standard format iii. Typical patenting procedure	
	iv. Rights of a patentee  d. Uses of patent system  D. Recent trends of biotechnological and microbiological patents - Patent forms (viz. multicellular organisms, DNA sequences)	ing of life

**Text Books** 

Casida L. E., "Industrial Microbiology" 2009 Reprint, New Age International (P) Ltd,

2. Glick B.R. & Pasternak J. J., 2003, "Molecular Biotechnology, Principles and Applications of Recombinant DNA", 3rd Edition, ASM Press, Washington, USA

3. Stanbury P. F., Whitaker A. & Hall--S. J., 1997, "Principles of Fermentation Technology", 2nd Edition, Aditya Books Pvt. Ltd, New Delhi.

4. Crueger W. and Crueger A. 2000 "Biotechnology -"A Textbook of Industrial Microbiology", 2nd Edition, Panima Publishing Corporation, New Delhi.

- 5. Prescott and Dunn's "Industrial Microbiology". 1982 4th Edition, McMillan Publishers 6. Ratledge & B. Kristinsen 2<sup>nd</sup> edn 2006. "Basic Biotechnology". Cambridge University
- 7. R. C. Dubey, 2005 A Textbook of "Biotechnology" S. Chand and Company, New Delhi

8. Indu Shekar Thakur 2006 "Industrial Biotechnology" Problems and Remedies, I K International Pvt Ltd

9. S. K. Kulkarni, Nanotechnology: Principles and Practices, Capital Publishing Co.

#### Reference Books

1. Peppler, H. J. and Perlman, D. (1979), "Microbial Technology". Vol 1 & 2, Academic Press

2. U. Satyanarayana 2005. "Biotechnology". Books and Allied (P) Ltd

3. Agrawal A. K. and P. Parihar 2005. "Industrial Microbiology"- Fundamentals and Application AGRIBIOS (India)

4. H. A. Modi, 2009. "Fermentation Technology" Vols 1 & 2, Pointer Publications, india

5. Okafor Nkuda 2007 "Modern Industrial Microbiology and Biotechnology", Science Publications Enfield, NH, USA.

#### Practical Syllabus Based On Paper IV

1. Comparison of amylase activity of Aspergillus culture grown in liquid medium and on solid substrate.

2. Chemical and Bioassay of Penicillin

3. Bioautography and Bioassay of Vitamin B12.

Isolation of lactic acid bacteria from Probiotic foods.

5. Sugar and alcohol tolerance of Saccharomyces cerevesiae.

6. Ethanol production from jaggery, chemical estimation of sugar by Cole's Method, alcohol produced by dichromate method, efficiency of fermentation

7. Sterility testing of injectable (D/W ampoules)

8. Estimation of BOD and COD from distillery effluent.

- 9. Immobilization of enzyme---preparation of alginate-enzyme/culture beads, qualitative and quantitative activity estimation, viable count of bead culture.
- 10. Plant tissue culture (Demonstration)
- 11. Visits
  - A. Antibiotic production plant or Pharmaceutical Industry

B. Vaccine Production Plant (Animal/ Human)

C. Application of Recombinant DNA in Industrial Production

# Suggested Examination Pattern

- 1. Students opting for 6 Units of Microbiology (Major) at T Y B Sc level will study Papers I, III. IV of 100 marks each.
- II, III, IV of 100 marks each and 4 practicals based on these papers of 50 marks each. 2. Students opting for 3 Units of Microbiology at T Y B Sc level will study Papers I & II of 100 marks each and two practicals based on these papers of 50 marks each.
- 3. Quiz and spots shall be based exclusively on the practical syllabus. 4. Theory Examination - Four papers of 100 marks each of 3 hr duration as per the prescribed university pattern for B.Sc. should be followed. Each paper should cover entire syllabus in proportionate manner, using number of lectures assigned to the topic
- 5. Practical Examination
  - A. <u>3 Units:</u> -As per university directives, should be of 100 marks; two practicals of 50 marks each. The following pattern of practical exam is suggested.

	D		
20	Practical 2		
10	Medical	40	
10			
10	Rapid diagnostics	10	
50 *	Total	50	
	10 10	10 Medical 10 Rapid diagnostics	

- Techniques and chemical estimation shall be based on all relevant practicals including demonstrations / group experiments based on two papers.
- Practical examination will be held on 3 consecutive days between 10.00 a.m. to 4.00 p.m. with half hour lunch break.
- iii. Laboratory journal is to be duly certified by the Head, Department of Microbiology. Examiners are required to sign the journal and report at the end of examination.
- B. 6 Units As per university directives, should be of 200 marks; four practical of 50 marks each. The following pattern of practical exam is suggested.

Practical 1 Technique [40]		Practical 2		Practical 3		Practical 4	
		Medical	40	Chemical	30	Bioassay	40
Problem 1	25		10	Estimation	30	Dioassay	40
Problem 2 15	15	Rapid diagnostics		Journal	16	Quiz	10
Journal	10			Spots	10		
Total	50	Total	50	Total	50	Total	50

- Techniques and chemical estimation will be based on all relevant practical including demonstrations / group experiments based on all 4 paper practical.
- ii. Practical examination will be held on 3 consecutive days between 10.00 a.m. to 5.00 p.m. with half hour lunch break.
- iii. Laboratory journal is to be duly certified by the Head, Department of Microbiology. Examiners are required to sign the journal and report at the end of examination.

