Annexure No.	43 A
SCAA Dated	29.02.2008

#### **BHARATHIAR UNIVERSITY COIMBATORE – 641 046**

#### REGULATIONS FOR B.Sc. BIOTECHNOLOGY DEGREE COURSE WITH COMPULSORY DIPLOMA IN MEDICAL BIOTECHNOLOGY With semester system

(With effect from 2007 - 2008)

#### 1. Eligibility for admission to the course

Candidate for admission to the first year **B.Sc. Biotechnology** degree course shall be required to have passed the higher secondary examination with one of the subjects as Biology or Zoology/Botany (academic) conducted by the Govt. of Tamilnadu in the relevant subjects or other examinations accepted as equivalent there to by the syndicate, subject to such other conditions as may be prescribed therefore.

#### 2. Duration of the course

The course shall extend over a period of three years comprising of six semesters with two semesters in one academic year. There shall not be less than 90 working days for each semester. Examination shall be conducted at the end of every semester for the respective subjects, for the practicals at the end of the academic year

#### 3. Course of study

The course of study for the UG degree courses of all branches shall consist of the fallowing **a**) **Part I** 

Tamil or any one of the fallowing modern/classical languages i.e., Telugu, Kannada, Malayalam, Hindi, Sanskrit, French, German, Arabic, & Urdu.

The subject shall be offered during the first four semesters with one examination at the end of each semester

#### b)Part II: English

The subject shall be offered during the first four semesters with one examination at the end of each semester. During third semester part II English will be offered as communication skills **c)** Foundation course

The Foundation course shall comprise of two stages as follows;

Foundation course A: General awareness (I & II semesters)

Foundation course B: Environmental studies (III & IV semesters)

1. The syllabus and scheme of examination for the foundation course A. general awareness shall be apportioned as follows

From the printed material supplied by the University -75%Current affairs & who is who? -25%

The current affairs cover current developments in all aspects of general knowledge which are not covered in the printed material on this subject issued by the University.

2. The foundation course B shall comprise of only one paper which shall have environmental studies.

#### a. Part III

**Group** A: core subject – As prescribed in the scheme of examination. Examination will be conducted in the core subjects at the end of every semester

Group B: Allied subjects – two subjects – four papers

Examination shall be conducted in the allied subjects at the end of first four semesters.

Group C: application oriented subjects: Two subjects – Four papers.

The application-oriented subjects shall be offered during the last two semesters of study viz., V and VI semesters. Examination shall be conducted in the subjects at the end of V & VI semesters

Group D: Field work/Institutional training

Every student shall be required to undergo training in the biotechnology industry, related to the application –oriented subject for a period of not less than 2 weeks, conveniently arranged during the course of  $3^{rd}$  year and submit a training report based on his/her training. The principal of the college and the head of the department shall issue a certificate to the effect that the student had satisfactorily undergone the fieldwork/institutional training for the prescribed period.

#### **Diploma Programme:**

All the UG programmes shall offer compulsory diploma subjects and it shall be offered in four papers spread over each paper at the end of III, IV, V and VI semesters.

#### e) Co-curricular activities: NSS/NCC/Physical education:

Each student shall participate compulsorily for the period of not less than two years (4 semesters) in any one of the above pragrammes.

The above activities shall be conducted outside the regular working hours of the college. The principal shall furnish a certificate regarding the student's performance in the respective field and shall grade the student in the five point scale as follows

- A- Exemplary
- B- Very good
- C- Good
- D- Fair
- E- Satisfactory

This grading shall be incorporated in the mark sheet to be issued at the end of the appropriate semester  $(4^{th} \text{ or } 5^{th} \text{ or } 6^{th} \text{ semester})$ 

(Handicapped students who are unable to participate in any of the above activities shall be required to take a test in the theoretical aspects of any one of the above 3 field and be graded and certified accordingly).

### 4. **Requirement to appear for the examinations.**

- a) A candidate will be permitted to appear for the university examinations for any semester if
- i) He/she secures not less than 75% of attendance in the number of working days during the semester.

ii) He /she earns a progress certificate from the head of the institution, of having satisfactorily completed the course of study prescribed in the subjects as required by these regulations and iii) His / her conduct has been satisfactory.

Provided that it shall be open to the syndicate, or any authority delegated with such powers by the syndicate to grant exemption to a candidate who has failed to earn 75% of the attendance prescribed, for valid reasons, subject to usual conditions.

- b) A candidate who has secured less than 65 % but 55% and above attendance in any semester has to compensate the shortage in attendance in the subsequent semester besides, earning the required percentage of attendance in that semester and appear for both semester papers together at the end of the latter semester.
- c) A candidate who has secured less than 55% of attendance in any semester will not be permitted to appear for the regular examinations and to continue the study in the subsequent semester. He/she has to rejoin the semester in which the attendance is less than 55%.
- d) A candidate who has secured less than 65% of attendance in the final semester has to compensate his/her attendance shortage in a manner as decided by the concerned head of the department after rejoining the same course

#### 5. **Restriction to appear for the examinations**

- a) Any candidate having arrear paper(s) shall have the option to appear in any arrear paper along with the regular semester papers.
- b) Candidates who fail in any of the papers in part I, II & III of UG Degree examinations shall complete the paper concerned within 5 years from the date of admission to the said course, and should they fail to do so, they shall take the examination in the texts/revised syllabus prescribed for the immediate next batch of candidates. If there is no change in the texts/syllabus they shall appear for the examination in the paper with the syllabus in the vogue until there is a change in the texts or syllabus. In the event of removal of that paper consequent to change of regulation and/or curriculum after 5 year period, the candidates shall have to take up an equivalent paper in the revised syllabus as suggested by the chairman and fulfill the requirements as per regulation/curriculum for the award of the degree.

# 6. Medium of instruction and examinations

The medium of instruction and examinations for the papers of part I and part II shall be the language concerned. For part III subjects other than modern languages, the medium of instruction shall be either Tamil or English and the medium of examinations is in English/Tamil irrespective of the medium of instructions. For modern languages, the medium of instruction and examination will be in the languages concerned.

#### 7. Submission of record note book for practical examinations.

Candidates appearing for practical examinations should submit *bonafide* Record note books prescribed for practical examinations, otherwise the candidates will not be permitted to appear for the practical examinations. However, in genuine cases where the students, who could not submit the record note books, they may be permitted to appear for the practical examinations, provided the concerned Head of the department from the institution of the candidate certified that the candidate has performed the experiments prescribed for the course. For such candidates who do not submit record books, zero (0) marks will be awarded for record note books.

#### 8. Passing minimum

a) A candidate who secures not less than 40% of the total marks in any subject including the Diploma and Foundation courses (theory and practical) in the University examination shall be declared to have passed in the subject (theory and practical).

b) A candidate who passes the examination in all the subjects of part I, II and III (including the diploma and Foundation courses) shall be declared to have passed, the whole examination.

#### 9. Improvement of Marks in the subjects already passed

Candidates desirous of improving the marks awarded in a passed subject in their first attempt shall reappear once within a period of subject of subsequent two semesters. The improved marks shall be considered for the classification but not for ranking. When there is no improvement, there shall not be any change in the original marks already awarded.

#### **10. Classification of Successful candidates**

a) A candidate who passes all the Part II examinations in the First attempt within a period of three years securing 75% and above in the aggregate of Part III marks shall be declared to have passed B.A/B.sc,/B.Com./B.B.M., degree examination in <u>First Class with Distinction</u>

b) (i) A candidate who passes all the examination in Part I or Part II or Part III or Diploma securing not less than 60% of total marks for concerned Part shall be declared to have passed in <u>Fist Class</u>

(ii) A candidate who passed all the examination in Part I or Part II or Part III or Diploma securing not less than 50% but below 60% of total marks for concerned Part shall be declared to have passed in <u>Second Class.</u>

(iii) All other successful candidates shall be declared to have passed the Part I or Part II or Part II or Diploma examination in <u>Third Class.</u>

#### **11. Conferment of the Degree**

No candidate shall be eligible for conferment of the Degree unless he / she

i. Has undergone the prescribed course of study for a period of not less than six semesters in an institution approved by affiliated to the University or has been exempted from in the manner prescribed and has passed the examination as have been prescribed therefore.

ii. Has satisfactorily participates in either NSS or NCC or Physical Education as evidenced by a certificate issued by the Principle of the Institution.

iii. Has successfully completed the prescribed Field Work/ Industrial Training as evidenced by certificate issued by the Principal of the College.

# 12. Ranking

A candidate who qualifies for the UG Degree course passing all the examinations in the first attempt, within the minimum period prescribed for the course of study from the date of admission to the course and secures I or II class shall be eligible for ranking and such ranking will be confined to 10% of the total number of candidates qualified in that particular branch of study, subject to a maximum of 10 ranks.

The improved marks will not be taken into consideration for ranking

#### **13. Additional Degree**

Any candidate who wishes to obtain an additional UG degree not involving any practical shall be permitted to do so and such candidate shall join a college in the III year of the course and he/she will be permitted to appear for Part III alone by granting exemption from appearing Part I, Part II and common allied subjects (if any), already passed by the candidate. And a candidate desirous to obtain an additional UG degree involving practical shall be permitted to do so and such candidate shall join a college in the II year of the course and he/she be permitted to appear for Part II alone by granting exemption from appearing for Part I, Part II and the common allied subjects. If any, already passed. Such candidate should obtain exemption from the University by paying a fee of Rs 500/-.

#### **14. Evening College**

The above regulations shall be applicable for candidates under going the respective courses in evening Colleges also.

#### 15. Revision of Regulations and Curriculum

The above Regulation and scheme of Examinations will be in vogue without any change for a minimum period of three years from the date of approval of the Regulations. The University may revise / amend/ change the regulations and scheme of Examinations, if found necessary.

#### **16.** Transitory provision

Candidates who have undergone the course of study prior to the academic year 2007-2008 will be permitted to take the Examination under those regulations for a period of four years i.e., up to and inclusive of April 2012. Thereafter they will be permitted to take the Examination only under the regulations in force at that time.

	part		Tanta di sal	University	
Sem		Subject & paper	Instructional	Dura-	Max
			nours / week	tion	marks
т	т	I on over an annar I	6	2	100
1	I TT	Language paper I	6	3	100
		English paper I	6	3	100
	111	Gr. A Core paper I Cell biology	4	3	100
		Gr. A .Core Paper II Biodiversity	4	3	100
		Gr. B. Allied A paper I Chemistry	4	3	15
		Core practical I	2	-	-
		Allied Practical	2	-	-
	FC	FCA	2	-	-
II	Ι	Language paper I	6	3	100
	II	English paper I	6	3	100
	III	Gr. A. Core paper III Biochemistry	5	3	100
		Core Practical I (cell Biology &	4	3	100
		biochemistry)			
		Gr B Allied Paper II chemistry	4	3	75
		Allied practical	3	3	50
	FC	Foundation course A	2	3	100
III	Ι	Language Paper III	6	3	100
	II	English Paper III	6	3	100
	III	Core Paper IV Microbiology	4	3	100
		Core Paper V Genetics	4	3	100
		Gr B Allied B Paper III	4	3	75
		Basic Mathematics			
		Practical II (Microbiology & Genetics)	2	-	-
		FCB	1	-	-
		Diploma Paper I Human physiology	3	3	100
IV	Ι	Language Paper IV	6	3	100
	II	English Paper IV	6	3	100
	III	Gr A Core Paper VI Bioinformatics	4	3	100
		Practical II (Microbiology & Genetics)	4	3	100
		Gr B Allied B paper II (Computer	4	3	75
		applications)	_		
		Allied Practical II	2	3	50
	FC	Foundation course B	1	3	100
		Diploma Paper II Human pathology	3	3	100

# SCHEME OF EXAMINATION FOR B. Sc. BIOTECHNOLOGY

V	III	Gr A Core paper VII Immunology	4	3	100
		Core Paper VIII Plant Biotechnology	4	3	100
		Core Paper IX Animal Biotechnology	4	3	100
		Core Practical III Applied	3	-	-
		Biotechnology			
		AOS practical IV:Lab in immunology	4	-	-
		and rDNA technology			
		Gr C (Application oriented subject A)	4	3	75
		Paper I Molecular Genetics			
		Gr C (Application oriented subject A)	4	3	75
		Paper II Biology of cloning Vectors			
		Diploma Paper III Diagnostic tools	3	3	100
VI	III	Gr A Core Paper X Microbial	5	3	100
		Biotechnology			
		Core Paper XI Environmental	5	3	100
		Biotechnology			
		Core Practical III applied biotechnology	4	6	100
		AOS practical IV:Lab in immunology	5	6	100
		and rDNA technology			
		Gr C (Application oriented subject B)	4	3	75
		Paper I rDNA technology & Bioethics			
		Gr C (Application oriented subject B)	4	3	75
		Paper II Biophysics &			
		Bioinstrumentation			
		Diploma Paper IV Pharmacology	3	3	100

# COMPULSORY DIPLOMA IN MEDICAL BIOTECHNOLOGY

Paper I	HUMAN PHYSIOLOGY	3	100
Paper II	HUMAN PATHOLOGY	3	100
Paper III	DIAGNOSTIC TOOLS	3	100
Paper IV	PHARMACOLOGY	3	100

# **CORE PAPER: I**

#### Subject Title: CELL BIOLOGY

**Subject description:** This course presents the types and structural details of the basic unit by which all the living things are made of (the cell).

Goals: To make the student to understand the concept of cell and their activities.

**Objectives:** On successful completion the subject student should have understand: Structural features, Organelles and the cellular mechanisms.

#### UNIT I

Cell as a basic unit: discovery of the cells, classification of cell types, development of cell theory, early chemical investigation in cell biology. Prokaryotic and Eukaryotic cell organization.

#### UNIT II

Cell transport phenomenon: membrane architecture. Active, Passive, diffusion and osmosis. Chemistry of carbohydrates, lipids, proteins and nucleic acids.

#### UNIT III

Structure and function of cytoplasmic compartments of the cell: ribosome and protein synthesis, energy flow through mitochondrion, chloroplast and photosynthesis, Golgi apparatus, lysozymes and micro bodies, endoplasmic reticulum, cytoskeleton, vacuoles, peroxysomes, lysozomes and Nuclear compartment. Heterochromatin and euchromatin, polytene chromosomes.

#### UNIT IV

Cell division in prokaryotes and eukaryotes: Cell cycle, mitosis, meiosis, crossing over and characteristics of cancer. Apoptosis, Stem cell. Prions.

#### UNIT V

Integrative and specialized cellular events, cell-cell signaling, specialized cells nerve cells, sperm cells, microfilaments, microtubules, muscle cells. Cells of vision, Nucleo-cytoplasmic interaction, cell cloning.

#### References

- 1. Cell and molecular biology, 3<sup>rd</sup> edition, Philip Sheeler, Donal E Bianchi, John Wiley
- 2. Molecular biology of cell, Alberts et al
- 3. Molecular cell biology, Lodish, Baltimore, Scientific American books, 1994
- 4. Molecular and cell biology, Stephen L Wolfe, Wordsworth Publishing company 1993
- 5. Cell biology. Sadava
- 6. Cell and Molecular Biology De Roberties

#### **CORE PAPER: II**

#### Subject Title: BIODIVERSITY

Subject description: This course presents the Divers nature of the organisms

Goals: To make the student to understand the diversity of the nature

**Objectives:** On successful completion the subject student should have understand: Species and genius on the earth, conservation and development of biological resources.

#### UNIT-I

General aspects of biodiversity-General introduction. Types of Biodiversity-Global biodiversity, biodiversity in India-Species biodiversity, Measures of biodiversity-Loss of biodiversity. Listing of threatened biodiversity-threatened animals, Plants, causes for the loss of biodiversity. Biodiversity-strategy and action plan.

#### UNIT II

Species concept; biological nomenclature. Theories of biological classification; structural, biochemical and molecular systematic. DNA fingerprinting, numerical taxonomy, magnitude and distribution of biodiversity, economic value, wild life biology,.

#### UNIT III

Broad outlines of classification and evolutionary trends among algae, fungi, bryophytes and terydophytes. Economic importance of algae, fungi and lichens. Biotic community-concept, structure, dominance, fluctuation and succession; N,P,C,S cycles in nature Principles of conservation, conservation strategy and sustainable development.

#### UNIT IV

Interaction between environment and biota; concept of habitat and ecological niches, limiting factor, energy flow, food chain, food web and tropic levels, ecological pyramids and recycling.. Ecosystem dynamics and management, stability and complexity of ecosystem, speciation and extinctions, environmental impact assessment.

#### UNIT V

Physiochemical properties of water, distribution and impact of experimental factors on the aquatic biota, productivity, mineral cycles and biodegradation in different ecosystem, biology and ecology of reservoirs.

#### References

- 1. Glimpses of Biodiversity- B.Blosetti.
- 2. Environmental biodiversity- P.R.Yadav
- 3. Biodiversity of microbial life- Stanely Reysenbach
- 4. Ecology & Env. Biology Sathyanarayana Books & Allied (P) Ltd

# **CORE PAPER: III**

# Subject title: BIOCHEMISTRY

**Subject description:** This course presents the chemical reactions or metabolic functions in the living system and their regulations.

Goals: To make the student to understand the concept of biochemical regulations

**Objectives:** On successful completion the subject student should have understand: Basic metabolism Enzymes and their kinetics Applications of metabolites

#### UNIT I

Structure of atoms and biomolecules: atomic theory, valancy, atomic weight, molecular weight, Molarity.

Chemical Bonding, properties of Water ionization p<sup>H</sup> and buffers

#### UNIT II

Introduction to Biochemistry: Fundamental Structures of proteins, nucleic acid, lipid and carbohydrates.

Thermodynamics in biology: Energy metabolism. Free energy, energy rich bonds, week interactions, coupled reactions, kinetics, association dissociation concepts, biological energy transducers.

#### UNIT III

Enzymes and co-enzymes, IUB classification and nomenclature of enzymes, enzyme kinetics (positive and negative cooperativity), regulation of enzyme activity, active sites, activators and inhibitors; allosteric enzymes

#### UNIT IV

Classifications and reactions of sugars. structural features of polysaccharides. Glycolysis, TCA cycle, glycogen breakdown and synthesis, gluconeogenesis, bioconversion of pentoses and hexoses, ETC, Oxidative and photo phospharylation

Classification and functions of lipids.

Metabolism of lipids; Fatty acid biosynthesis and oxidations,

#### UNIT V

Amino acids and peptides – classifications chemical reactions and physical properties, metabolisms of amino acids. Functions of proteins.Biosynthesis and degradation of nucleic acids (purines and pyrimidines) Integration of metabolism and regulations.

Secondary metabolites -applications,

isoprenoids and hetero cyclic compounds

**REFERENCES**:

- 1. Boyer.R., (2002) Concepts in Biochemistry 2<sup>nd</sup> ed. Brooks/cole publishing company New York.
- 2. David L. Nelson and M. Cox (2003) Lehninger's Principles of Biochemistry, 3<sup>rd</sup> Ed, Worth publication New York
- 3. Voet & Voet (1995) Fundamentals of Biochemistry, 2<sup>nd</sup> Ed, John Wiley and sons inc., New York.
- 4. Geoffery L Zubay (1995) Principles of Biochemistry, WCB publishers, London
- 5. Murrey RK., D.K. Granner, P.A. Mayers and V.W. Rodwell, (2003) Harper's Biochemistry, Prentice –Hall Int, Boston
- 6. Outlines of Biochemistry Conn & Stumph
- 7. Biochemistry 3<sup>rd</sup> ed Sathyanarayana Books & Allied (P) Ltd

# **CORE PRACTICAL I:**

# LAB IN: CELL BIOLOGY AND BIOCHEMISTRY.

- 1. Microscopy
- 2. Cell Types --- Microbial, animal and Plant cells simple staining and visualization through microscope
- 3. Fraction of Cellular components—Demonstration.
- 4. Mitotic Preparation Onion Root Tip, Grasshopper Hepatic coacae
- 5. Meiotic Preparation from Grasshopper Testis and flower buds of Rheo discolor
- 6. Cell Staining Cytochemical Methods for Demonstration of Cellular and sub-cellular components.
- 7. Estimation of Protein Lowry's, Bradford's method.
- 8. Estimation of DNA--- DPA Method.
- 9. Estimation of RNA by Orcinol method
- 10. Estimation Of Sugars --- Anthrone and Benedict method
- 11. Estimation of total free amino acids --- Sulfovanicillin method.
- 12. Estimation of Lipids.
- 13. Analysis of Oils—Iodine Number---Saponification Value ---Acid Number.
- 14. Quantification of Vitamin C.
- 15. Thin Layer Chromatography.
- 16. Paper Chromatography.
- 17. Isolation of Enzymes.
- 18. Enzyme assay.
- 19. Determination of Km value.

## **CORE PAPER: IV**

#### Subject Title: MICROBIOLOGY

Subject description: This course presents the study of Micro organisms.

**Goals**: To make the student to understand Micro organisms and their participation in day to day activities.

**Objectives:** On successful completion the subject student should have understand: What are micro organisms? Their studying methods and their positive and negative on our lives

#### UNIT I:

Definition and scope of microbiology-- A general account on microbial diversity.Basic principles in microscopy, Types of microscopes- light, dark, phase contrast, fluorescent and electron microscope- (Transmission and Scanning electron)

#### UNIT II:

A detailed account of General structure, growth and reproduction of the various Bacteria, fungi and Viruses. Economic and industrial importance of yeast and moulds

#### UNIT III:

Microbiological Media: Types, preparation, methods of sterilization; enumeration of microorganisms in soil, water and air; isolation of microorganisms from Environment and infected tissue; Techniques of pure culture, maintenance and Preservation; Staining: stains and types of staining;

#### UNIT IV:

Physiology and biochemistry of microbes--Nutrition (Photo-autotrophs, Chemo-autotrophs, Parasitism, Saprophytism, Mutualism and Symbiosis, Commensalisms, endozoic microbes) - - microbial pathogens of plants, animals and Humans.

#### UNIT V:

Respiration and fermentation, Nitrogen metabolism including Nitrogen fixation (Symbiotic and asymbiotic), Lipid metabolism, Secondary metabolism, Production of enzymes and antibiotics--Role of microbes in biogeochemical cycles.

#### References:

1. Michael T. Madigan John M. Martin & Jack Parker, 1984, Biology of

Microorganisms Prentice Hall International, Inc., London.

2. Edward A. Birge, 1992, Modern Microbiology – Principles and application. Wm.C. Brown Publishers, Inc. U.S.A.

3. Gerard J. Tortora, Berdell R. Funke, Christine & L. Case, 2001, Microbiology -

An Introduction. Benjamin Cummings, U.S.A.

4. Danial Lim, 1998, Microbiology, McGraw-Hill Companies, New York.

5. Stephen A. Hill, 1984, Methods in Virology. Blackwell Scientific Publication, London.

## **CORE PAPER: V**

#### **Subject Title: GENETICS**

**Subject description:** This course presents the way characters get transferred through generations and methods to analyze and modify them **Goals**: To make the student to understand the concept of genes and their behaviour

**Objectives:** On successful completion the subject student should have understand: Basic genetics

Gene expression and regulation

#### UNIT I

History of Genetics - Mendelian Principles, Segregation, Independent Assortment, Dominance relations, Multiple alleles, Incomplete dominance, Over dominance,

#### UNIT II

Gene interaction, Epistasis, lethality and lethal genes, Sex determination and sex linkage in diploids, linkage and crossing over, gene mapping. Chromosomal theory of inheritance, maternal effects.

#### UNIT III

Chromosomal variation in number, Changes in Chromosomal structure, Chromosomal aberrations, Genetics of Heamoglobin, Transposable elements in prokaryotes and eukaryotes.

#### UNIT IV

Fine structure of Gene, cistron, recon, Structure of Eukaryotic gene, Experimental evidence for DNA as the genetic material, cytoplasmic genetic systems- mitochondria and chloroplast DNA, Plasmids- F, R and Col plasmids. Relation between genes and polypeptides.

#### UNIT V

Population genetics, calculating gene frequency, factors affecting gene frequency. Genetic control of Development in Drosophila and Arabidopsis. Genetic drift, Shift, Pedigree analysis and genetic counseling.

#### Reference:

- 1. Basic genetics by D.L.Hartl,1991,Jones and Bartett public.
- 2. Friedfelder 1987, Microbial genetics ,Jones and Bartett public.
- 3. Molecular Biology of the genes 4<sup>th</sup> Ed. Watson et.,al, the Benjamine /cummings coins 1987
- 4. Molecular cell biology, 1994.Lodish, Baltimore scientific American books,Inc.
- 5. Genetics Strickberger.M
- 6. Genetics by Goodenough

### **Diploma Paper I:**

#### Subject Title: HUMAN PHYSIOLOGY

**Subject description**: This course presents the various physiological activities in human being

**Goals:** To make the student to understand the human physiology

#### **Objectives**:

After the completion of the course the student should have understood varius systems in human and their activities

#### UNIT I

MUSCLE- skeletal muscles – composition – functions and properties of plain (smooth) and cardiac muscles – electromyography

NERVOUS SYSTEM – organization – basic functions of synapses and transmitter substances – sensory receptors – sense of hearing – taste and smell. Special senses – optics of vision – function of retina –cortical and brain stem control of motor function . Cerebellum – limbic and hypothalamus – states of brain activity cerebral blood flow, cerebrospinal and brain metabolism

#### UNIT II

BLOOD & BODY FLUID – blood cell –Haematosis – determination of coagulation – plasma proteins – platelets – leucocytes. Bone marrow – functions of tissue fluid – Lymph nodes

CADIO VASCULAR SYSTEM – Heart as pump – rhythmic excitation – electrocardiogram – cardiac arrhythmias. Circulation- functions of arterial and venous system – microcirculation and lymphatic system – rapid control of arterial pressure- hypertension – cardiac failure – heart sounds

RESPIRATORY SYSTEM- pulmonary ventilation – pulmonary circulation – gaseous exchange -  $O_2$  and  $CO_2$  transport in blood and body fluids – mechanism of breathing - ventilation

#### UNIT III

DIGESTIVE SYSTEM – digestive tract – gastrointestinal function – motility– secretory functions of alimentary tract – digestion and absorption.

EXCRETION – functions of kidney – renal associated mechanisms - extracellular and intracellular fluids – osmalality and sodium concentration – regulation of blood volume – excretion – regulation of urea, potassium – regulation of acid base balance. Micturition – skin – sweat

ENDOCRINES – pituitary hormones and their control by hypothalamus – thyroid metabolic hormones – adreno-cortical hormones – insulin, glucagons and Diabetes mellitus – parathyroid hormone, calcitonin. Gonadotrophic hormones –testosterone – estrogen – corpus leutem - progesterone – Endometrical and menstruation – puberty – menopause – pregnancy and lactation – fetal and neonatal physiology

References:

- 1. Text book of Medical physiology by Guyton . 8<sup>th</sup> edition . W B Saunders company. USA
- 2. Human physiology by Dr.C.Chatterjee I & II. Medical Allied Agency, Kolkatta.
- 3. Anthony's Text book of Anatomy and Physiology by Gary A. Thiodeare & Kevin T patton, 2<sup>nd</sup> edition . Moshi year book, New York
- 4. Anatomy and Physiology by Ross & Wilson 8<sup>th</sup> edition. Churchill livingstone
- 5. Human physiology by Sarada Subramaniam & K.MadhavanKutty. S.Chand and company, New Delhi
- 6. Human Physiology by Vander Sherman Luciano McGraw Hill NewYork.

# CORE PAPER: VI

# Subject Title: BIOINFORMATICS

Subject description: This course presents the fundamentals of Information processing

Goals: To make the student to understand the concept Informatics

**Objectives:** On successful completion the subject student should have understand: Methods to retrieve and submit data Genome data bases and other databases and their analysis

# UNIT I

Introduction and history of bioinformatics – Internet, World Wide Web, Web browser, EMB net, NCBI. File transfer protocol. Search engines

# UNIT II

Database- Definition, DBMS – Biological Databases – FASTA, Blast, Genbank, DNA sequence databases, Protein databases. Entry formats, carbohydrate databases, Enzyme databases, Pathway databases. Relational database model. Theory on RDBMS. SQL, introduction to access, making queries. Designing forms. Report design

# UNIT III

Genomic resources, Gene structure and DNA sequences. EST searches, gene hunting, gene finders, Expression analysis- SAGE, cDNA library, EST, Microarray – DNA sequencing and sequence alignment – RFLP, SNP, RAPD, Human Genome Project, RNA analysis.

# UNIT IV

Proteomics – proteome analysis – 2D gel electrophoresis, Mass spectrophotometry, protein – protein interaction, protein – DNA interaction. Enzyme – Substrate interaction, pathway analysis.

# UNIT V

Application aspects – target searchings – drug designing – E- cell, phylogenetic analysis, PERL, Chemoinformatics

References:

- 1. Introduction to Bioinformatics T.K.Altwood, D.J.Parry-smith (2004) Pearson Education
- 2. Bioinformatics for the beginners K.Mani & N.vijayaraj
- 3. Proteomics- Pennigton & Dunn (2002) Viva books publishers, New Delhi
- 4. Bioinformatics- A practical guide to the analysis of genes & protein 2<sup>nd</sup> ED Andreas, Baxevanis and Francis Ouellette.
- 5. The internet (1999) Christian Crumlish. BPB publications.

## CORE PRACTICAL II: LAB IN: MICROBIOLOGY AND GENETICS.

- 1. Laboratory rules and regulations of Microbiology
- 2. Media preparation and sterilization
- 3. Enumeration of microorganism from soil, water and spoiled food--- serial dilution technique.
- 4. Pure culture technique—Pour plate, spread plate and streak plate methods.
- 5. Isolation of single colonies of bacteria
- 6. Auxotrophic selection
- 7. Measurement of growth of bacteria.
- 8. Measurement of growth of Phage.
- 9. Staining of bacteria—Gram's; Spore, capsule, acid fast bacilli.
- 10. Fungal Staining --- Wet Mount technique.
- 11. Drosophila Morphology, Section culture and maintenance.
- 12. Identification of Mutants—Physical and Chemical Methods.
- 13. Experiments to determine Mendel's law.
- 14. Monohybrid and dihybrid cross using plants.
- 15. Salivary Gland chromosome of
- 16. Human karyotype --- demonstration.
- 17. Sex chromatin (buccal smear).

# **DIPLOMA PAPER II:**

# Subject Title: HUMAN PATHOLOGY

# Subject description: This course presents the diseases of human being

Goals: To make the student to understand the concept Pathology

**Objectives:** On successful completion the subject student should have understand: Pathogen, disease and metabolic disorders.

### UNIT I:

<u>An introduction to Biotechniques in clinical medicine</u>: sampling, analysis, reporting, and interpretation of results.<u>Disorders of Kidney</u>: acute renal failure, chronic renal failure, proteinuria and nephritic syndrome and urinary calculi. <u>Disorders of Liver</u>: Biochemical assessment of liver function. Liver diseases: Acute hepatitis, chronic hepatitis, acute liver failure, Cirrhosis, alcohol and liver. Inherited abnormalities of bilirubin metabolism: Gilbert's, Crigler-Najjar, Dubin-Johnson, Jaundice, and Rotor. Drugs and the Liver. Biotechnological approaches to liver diseases: Vaccine development and drug delivery.

# UNIT II:

<u>Disorders of Carbohydrate metabolism</u>: Diabetes mellitus - Etiology and pathogenesis, diagnosis and management. Metabolic complications of diabetes: Ketoacidosis, pathogenesis, non - ketotic hyperglycaemia, Lactic acidosis, diabetic nephropathy, Lipoprotein metabolism in diabetes, Diabetes in pregnancy, glycosuria, hypoglycaemia - diagnosis and management. Disorders of Plasma proteins and enzymes: Hypoalbuminaemia, hypogammaglobulinaemia, hypergammaglobulinaemia. Alkaline phosphatase, Creatine kinase, Haemoproteins -Haemoglobinopathies, and abnormal haemoglobin derivatives-Methaemoglobin, Carboxyhaemoglobin. Secondary hyper lipidaemia, Types of Primary hyperlipidaemias. Lipoprotein deficiency-abetalipproteinaemia. Diseases of Heart-Myocardial infarction, Heart failure and Hypertension. Inherited metabolic diseases: Glucose-6-phosphatase deficiency, Galactosaemia, Phenyl ketonuria, Cystic fibrosis

# UNIT III:

<u>Disorders of hypothalamus and pituitary</u>: Disorders of anterior pituitary hormones: Hypopituitarism, Anorexia nervosa, Growth hormone deficiency, Growth hormone excess: acromegaly and gigantism, Hyperprolactinaemia and Cushing's disease.

#### Disorders of Adrenal Glands:

Disorders of Adrenal cortex: Adrenal hypofunction (Addison's disease). Adrenal hyper function: Cushing's syndrome, conn's syndrome, congenital adrenal hyperplasia (CAH). Disorders of adrenal medulla: catacholamines.

Disorders of Thyroid gland: Hyperthyroidism, hypothyroidism, thyroiditis, goiter and thyroid cancer.

<u>Metabolic aspects of cancer</u>: Metabolic complications of cancer. Tumour markers:  $\alpha$ -Fetoprotein (AFP), Carcinoembryonic antigen (CEA), Para proteins, Human chorionic gonadotrophin (hCG), markers of prostatic cancer, enzymes as tumour markers and Carbohydrate antigen (CA) markers.

# **REFERENCES**:

- 1. Clinical Chemistry by Willium J.Marshall (Fifth edition, Mosby Publications).
- 2. An Illustrated color text of Clinical Biochemistry by Allen Gaw, Robert A.Cowan, illustrated by Robert Britton (1999, second edition, Churchill Living stone press).
- 3. Harper's Illustrated Biochemistry (27th Edition) by Robert K. Murray, Daryl K. Granner, Victor W. Rodwell.
- 4. Lippincott's Illustrated reviews: Biochemistry (Lippincott press, Third Edition) by Richard Harvey and Pamela C.Champe.

- **5.** Medical Microbiology by PanickerMedical Microbiology by Roitt Medical Parasitology by Panicker
- 6. Color Atlas of Biochemistry (second edition, Thieme Publications, revised and enlarged) by Jan Koolman and Klaus-Heinrich Roehm.
- 7. Marks' Basic Medical Biochemistry: A Clinical Approach (2nd Edition), by Colleen M. Smith, Allan D. Marks and Michael A. Lieberman.
- 8. Medical Microbiology by Jawetz.

# CORE PAPER: VII

# Subject Title: IMMUNOLOGY

Subject description: This course presents the basic defense mechanism of animals

Goals: To make the student to understand the concept immunology

**Objectives:** On successful completion the subject student should have understand: Immunity, Antigen, Antibody, Cells of immune system and their function and regulations

#### UNIT I:

Introduction- Historical Development in Immunology. Immunity-. Humoral and Cell mediated response, Primary and Secondary immune response. Cells involved in immune response. Innate and Acquired Immunity. Mechanisms of defense.

#### UNIT II:

Antigen- Types and classifications. Antibody – Structure, Types, properties and their biological functions, poly clonal sera, Monoclonal antibody. Primary and Secondary lymphoid organs – Thymus, Bone marrow, Lymph nodes and Spleen. Lymphocytes traffic and regulation, CD molecules

#### UNIT III:

Hematopoiesis and development of B and T lymphocytes. Immunoglobulin Gene expression B cell and T cell activation. MHC molecules Response of B cells to antigens. Plasma Cells, Memory Cells.

#### Unit IV

Complement – activation and regulation. Cytokines- structure and functions, Interferons and interleukins. Immuno regulation: Tolerance. Suppression, Autoimmunity and hypersensitivity reactions .Primary and secondary Immuno deficiency disorders.

#### UNIT V:

Transplantation, HLA Typing; Mechanism of Graft rejection. Tumor immunology.Immuno surveillance- mechanisms.

Antigen – Antibody Interactions. Immunodiffussion and Immunoelectrophoresis. Principle and Applications of RIA, ELISA, Fluorescent Antibody techniques. I

#### **REFERENCES**:

- 1. Immunology Kuby., J  $-5^{th}$  Edition
- 2. Immunology Tizard
- 3. Immunology Ivan M. Roitt Third Edition
- 4. Immunobiology Janeway and Travers 5<sup>th</sup> Edition

# CORE PAPER: VIII Subject Title: PLANT BIOTECHNOLOGY

Subject description: This course presents the application of Plants in Biotechnology

**Goals**: To make the student to understand usage of Plants and methods to improve their utility

**Objectives:** On successful completion the subject student should have understand: Crop development, Callus culture, Biotechnological applications of plants

#### UNIT I

Conventional methods of crop improvement- Selection, mutation, polyploidy and clonal selection.

#### UNIT II

Plant genome organization, gene families in plant. Organization of chloroplast genome, nucleus- encoded and chloroplast encoded genes for chloroplast proteins. Organization of mitochondrial genome- nuclear and mitochondrial encoded genes for mitochondrial proteins, cytoplasmic male sterility. Seed storage proteins and heat shock proteins.

#### UNIT III

Plant tissue culture. Callus culture, organogenesis, meristem culture, anther, pollen, embryo culture and their applications .somatic hybridization Somatic embryogenesis, cybrids.

#### UNIT IV

Symbiotic nitrogen fixation in legumes -Biochemistry and molecular biology, gene rearrangement and nitrogen fixation in cyanophytes. Agrobacterium and Crown gall tumors. Ti plasmid vectors for plant transformation, agro-infection. Classification of plant viruses, molecular biology of plant stress response.

#### UNIT V

Direct transformation of plants by using physical methods Genetic engineering in plants, selectable markers, reporter genes and promoters used in plant vectors.. Genetic engineering of plants for virus resistance, pest resistance, herbicide tolerance, delay of fruit

ripening, resistance to fungi and bacteria. Production of antibodies, viral antigens and peptide hormones in plants. Importance of RFLP in plant breeding. Management aspect of plant genetic engineering, tagging and cloning of plant genes..

#### REFERENCE

- 1. An introduction to genetic engineering in plants, Mantel. S. H, Mathews. J. A, Mickee. R. A
- 2. Revolution on biotechnology, Marks. J.L.
- 3. Plant genetic engineering, Dodds J.H.
- 4. Plant molecule biology, Grierson and S.V. Convey
- 5. Molecular biotechnology, Principle and applications of recombinant DNA technology, Bernard R Glick.
- 6. Plant Biotechnology-Monica Hughes.

#### **CORE PAPER: IX**

#### Subject Title: ANIMAL BIOTECHNOLOGY

**Subject description:** This course presents the usage of Animals in biotechnology and their improvement

**Goals**: To make the student to understand the Animal products and exploitation of them in Biotechnology

**Objectives:** On successful completion the subject student should have understand: Animal tissue culture, Animal products, production & improvement of them.

#### UNIT I

Animal cell culture: Fundamentals. facilities and applications. Media for Animal cells. Biology of cultured cells, measurement of growth, cell synchronization, senescence and apoptosis

#### UNIT II

Types of cell culture: Primary cell culture, secondary culture, cell transformation, cell lines, stem cell cultures, cell viability and cytotoxicity. Organ culture. Cryopreservation Insect cell lines

#### UNIT III

Genetic engineering in animals: methods of DNA transfer into animal cells- calciumphosphate co precipitation, micro-injection, electroporation, Liposome encapsulation, Billogical vectors. Hybridoma technology, Vaccine production.

#### UNIT IV

Embyology: Collection and preservation of embryo, culture of embryos, culture of embryonic stem cells and its applications. Gametogenesis and fertilization in animals, Molecular events during fertilization, genetic regulations in embryonic development.

#### UNIT V

Transgenics: Transgenic animals. Production and recovery of products from animal tissue cultures: cytokines, Plasminogen activators, Blood clotting factors, Growth harmones.

#### **REFERENCES**:

- 1. Animal cell culture a practical approach, 4<sup>th</sup> ED., Freshney. John Wiley Pub.,
- 2. Methods in Cell Biology. VOL 57 Animal methods, ED Mather & Barnes, Academic Press.
- 3. Mammalian Cell Biotechnology- A practical approach. ED Butler. Oxford UNI Press.
- 4. Exploring Genetic mechanisms. ED Singer & Berg.

## Gr: C (Application Oriented Subject A) Paper I Subject Title: MOLECULAR GENETICS

Subject description: This course presents the genetics at molecular level

**Goals:** To make the student to understand the molecular genetics

#### UNIT- I

DNA as genetic material; Organization of genome – Structure and function of DNA and RNA; DNA replication – conservative, semi conservative, unidirectional, bidirectional replication; Enzymology of replication; *in-vitro* DNA synthesis

#### UNIT – II

Gene as the unit of expression; Colinearity: Transcription and gene regulation in prokaryotes and eukaryotes; elucidation of genetic code.

#### UNIT – III

Translation of protein – post translational modifications and folding of newly assembled polypeptides; transit peptide and signal sequences – protein export

#### UNIT - IV

Gene mutation – Biochemical basis of mutations – types of mutations- spontaneous and induced mutations; Ames test for mutation; DNA damage – types of DNA repair and mechanisms – photo reactivation excision repair, post replication recombinant repair, SOS repair

#### UNIT - V

Genetic exchange – bacterial transformation, transduction, conjugation and their mapping, linkage and chromosome mapping, crossing over, gene targeting.

#### REFERENCES

- 1. Basic Genetics by D.L. Hartl 1991, Jones & Bartett publications.
- 2. Microbail Genetics, Friefelder 1987 Jones & Bartnett publications
- 3. Molecular Biology of thegene 4<sup>th</sup> edition by Watson et al, The Benjamin / Cummings co
- 4. Molecular Cell Biology by Lodish 1994, Baltimore Scientific American Brocks

# Gr: C (Application Oriented Subject A) Paper II

# Subject Title: BIOLOGY OF CLONING VECTORS

Subject description: This course presents the types of cloning vehicle

**Goals**: To make the student to understand the concept of vector preparation, gene manipulation and gene transfer technologies

**Objectives:** On successful completion the subject student should have understand: Biology of plasmid, usability of plasmid and viral particles as vectors

#### UNIT-I

Introduction to cloning vectors: Plasmid Biology. *E.coli* vector; properties of plasmid (plasmids in gene transfer) plasmid compatibility, copy number control, PBR<sup>322</sup>, BAC and expression vectors in prokaryotes. Site - directed mutagenesis, m RNA isolation, cDNA synthesis.Genomic and cDNA liobraries.

#### UNIT II

Molecular biology of lambda, Lambda vectors; cosmid, phagemid. *in-vitro* packaging, M13 and other viral vectors of prokaryotes.

#### UNIT-III

Cloning in Yeast: genetics of S.cerevisiae, identification of Yeast genes, Yeast vectors, YAC. Cloning in Bacillus. Plasmids and vectors, inducible promoters. Cloning in Streptomyces.

#### UNIT-IV

Animal vectors; Selectable markers, SV40 Vectors, papiloma virus, Retero virus, Vaccinia virus.Bacculo virus

Ti plasmid as gene vector, Caulimo viruses, Gemini viruses, Transposable elements, RNA viruses, viroids

#### UNIT-V

Manipulation of genes for the Safety r DNA research. Laboratory and industrial applications. Reproductive engineering, Human genetic diseases, gene therapy, genetic manipulation of germ cells

#### **REFERENCES**:

- 1. Ernst.L.Winnacker, (2003) from genes to clones, 2nd edition, Panima publishing corporation, NewDelhi.
- 2. Benjamin Lewin (2004) Genes VIII, Pearson Education corporation, New Jersy
- 3. Alberts B, (1994) molecula biology of the cell, Garland publishing Inc New York
- 4. Friedfielder.D, (1987), Molecular biology II Ed., Narosa publishing house, New Delhi.
- 5. J.d.Watson (2001) Recombinant DNA technology, 2<sup>nd</sup> Ed WH Freeman and Company, New York
- 6. Brown T.A (1998) Introduction to gene cloning 3<sup>rd</sup> ED Stanley Thomas Pub ltd, Germany
- 7. Primbrose S.B (2003) Principles of gene manipulation 6<sup>th</sup> Ed Black well Sci ltd, Germany.

# **DIPLOMA PAPER III**

# Subject Title: DIAGNOSTIC TOOLS

UNIT I **Subject description:** This course presents the Diagnostic methods of diseases **Goals**: To make the student to understand the concept of Diagnostic methods

**Objectives:** On successful completion the subject student should have understand: Examination of Blood, Urine and CSF.

#### UNIT I

Blood examination – anticoagulant, hemoglobin, RBC, Packed cell volume, ESR, WBC total, differential normal and abnormal hematopathies – anemia, bone marrow smear, leukemia and myelodysplastic syndromes, diagnostic significance of PB smear, hemorrhagic disorder, L.E. cell phenomenon.

#### UNIT II

 $\label{eq:unine} Urine \ analysis - collection - physical, \ chemical \ and \ microscopic \ examination \ of \ urine - CSF$ 

Parasite analysis

#### UNIT III

Histopathology Biochemical analysis of Blood, Blood banking, Transplantation, AIDS, Lab safety, ELISA, RIA, FACS, PCR, Computers in lab. Quality control.

### **REFERENCE**:

- 1. Handbook of medical lab technology Ed; V.H.Talib, CBS publication
- 2. Clinical Chemistry by Willium J.Marshall (Fifth edition, Mosby Publications).
- 3. An Illustrated color text of Clinical Biochemistry by Allen Gaw, Robert A.Cowan, illustrated by Robert Britton (1999, second edition, Churchill Living stone press).
- 4. Marks' Basic Medical Biochemistry: A Clinical Approach (2nd Edition), by Colleen M. Smith, Allan D. Marks and Michael A. Lieberman.
- 5. Medical Microbiology by Jawetz.

#### CORE PAPER: X Subject Title: MICROBIAL BIOTECHNOLOGY

**Subject description:** This course presents the utility of Microbes **Goals**: To make the student to understand the applications of Microbes

**Objectives:** On successful completion the subject student should have understand: Fermentation, Microbial products, Vaccine and antibiotics.

#### UNIT I:

Microbial Biotechnology: Scope and application-horizons of microbial Technology, public concern about the microbial biotechnology and Economics of microbial biotechnology.

# UNIT II:

Microbes: Living factories for macromolecules-Production of proteins in Bacteria and yeast; recombinant and synthetic vaccines; microbial insecticides (*Bacillus.thuringiensis, B.spaerinus, B.papilliae* and Baculo-Viruses); microbial enzymes application in starch processing, textile designing, detergents, cheese making;polysaccharides and polyesters.

# UNIT III:

Microorganisms in fermentation-Ethanol from feed stocks to fermentable Sugars, from sugars to alcholos, clostridial fermentation, lactic acid fermentation, acetic acid production and industrial production of various milk products.

# UNIT IV:

Metabolites from microorganisms-amino acids; antibiotics-antibacterial agents (□-lactams, tetracyclines, peptides, amino glycosides), antifungal agents, anti-tumor antibodies.

# UNIT V:

Application of microbial biotechnology in sewage and wastewater treatment, degradation of xenobiotics, mineral recovery, removal of heavy metals from aqueous effluents, production of biofertilizers (nitrogen fixing Bacteria, single cell protein, mycorrizha and phosphate solubilizing Bacteria).

#### **REFERENCES** :

1. Glazer, A.N. and Nikaido, H. 1995. Microbial biotechnology. W.H.Freeman & Co., New York

2. Encyclopedia of Microbiology. 1992. Vols.1-4. Academic Press.

3. Preve et al. 1987. Fundamentals of Biotechnology. VCH Publ.

4. Stanbury, P.F. Whittaker, A, Hall, S.J. 1995. Principles of fermentation

technology.Butterworth Heinemann.

5. Prescott, L.M. Harley, J.P. and Klein, D.A. 1999. Microbiology. McGraw Hill Co.

6. Glick, B.R. and Pasternak, J.J. 1998. Molecular Biotechnology. Washington D.C. ASM Press.

7. Stainer, R.Y. Ingraham, J.L., Wheelis, M.L. and Painter, P.R. 1987. General Microbiology. Macmillan Co.

8. Lancini, G. Parenti, F. and Gallo, G.G. 1995. Antibiotics-A multidisciplinary Approach. Plenum Press, New York.

9. Gunasekaran.P. 1995. Laboratory manual in microbiology. New Age International Limited. New Delhi.

# CORE PAPER: XI Subject Title: ENVIRONMENTAL BIOTECHNOLOGY

**Subject description:** This course presents the Study and the Management of the Environment

Goals: To make the student to understand Ecology and Conservation of the Environment

**Objectives:** On successful completion the subject student should have understand: Ecosystem, Natural cycles. Diversity.

#### UNIT I:

Scope – Branches of ecology – Abiotic factors – water – soil – temperature – light. Biotic factors – Animal relationship – symbiosis – commensalisms – mutalism –Antagonism – Antibiosis – Parasitism – Predation – competition.

#### UNIT II:

Ecosystem –Definition –structure – pond ecosystem – primary production –secondary production – food chain – food web – trophic levels – energy flow – pyramid of biomass– pyramid of energy. Biogeochemical cycle: Nitrogen and Phosphorous.

#### UNIT III:

Pollution – types – sources – effects – Air-water – land – Noise – Thermal – Pesticide – Radioactive – green house effect, ozone and its importance – global warming – Acid rain– Bio accumulation – Bio magnification. Biological control. Principles of environment Impact. Assessment and environmental monitoring.

#### UNIT IV:

Uses and values of Biodiversity -A very general account on uses of Bioresources-plant uses: food, timber, medicinal ornamental and other uses- animal uses: food animals (terrestrial and aquatic), non food uses of animals, Domestic livestock-uses of microbes. Valuing Biodiversity-Instrumental (Goods, Services, and Information and Psychospiritual values) and Inherent or Intrinsic values, ethical and aesthetic values-An outline account on methods of valuing biodiversity. A general account on multilateral treaties- the role of CBD, IUCN, GEF, IBPGR, NBPGR, WWF, FAO, UNESCO and CITES-Bioresources, Biotechnology and Intellectual property rights:

# UNIT V:

Conservation of Biodiversity - Current Practices in conservation - Habitat or ecosystem approaches – Speciesbased approaches - Social approaches - Chipko movement - *In situ* (Afforestation, Social Forestry, Agro forestry, Botanical Gardens, Zoos, Biosphere Reserves, National Parks, Sanctuaries, Sacred Groves and Sthalavrikshas) and *Ex situ* (Cryopreservation, Gene Banks, Seed Banks, Pollen Banks, Sperms Banks, DNA Banks, Tissue Culture and Biotechnological Strategies), ecorestoration, environmental education.

# **REFERENCES**:

1. Groombridge, B (Ed.) 1992. Global Biodiversity – Status of the Earth's Living Resources. Chapman & Hall, London.

2. UNEP, 1995, Global Biodiversity Assessment, Cambridge Univ. Press, Cambridge.

3. Virchow, D. 1998. Conservation & Genetic Resources, Springer - Verlag, Berlin.

4. Gary K.Meffe & .Ronald Carroll ,C.1994. Principles of Conservation Biology, SinauerAssociates, Inc., Massachusetts.

5. Clarke, G.L. 1954, Elements of ecology, John Wiley & sons. N.Y.

6. Kendeigh, S.c. 1961. Animal Ecology. Prentice Hall.

7. Odum, E.P. 1971. Fundamentals of Ecology. W.B.Saunders company, Philadelphia.

8. Rastogi, V.B. and M.S. Jayaraj, 1989. Animal ecology and distribution of animals, Kedamath Ramnath.

9. Sharma, P.D. 1990. Ecology and environment. Rsatogi publications, Meerut.

10. Southwick, C.H. 1976. Ecology and the quality of environment D.Van.Nostrand Co.,

11. Verma P.S. and V.K. Agarwal. 1996. Principles of Ecology S.Chand. & co., New Delhi.

# Gr C (Application Oriented Subject B) Paper I Subject Title: RECOMBINANT DNA TECHNOLOGY AND BIOETHICS

Subject description: This course presents the mechanism of gene manipulation

**Goals**: To make the student to understand the concept of gene manipulation and gene transfer technologies

**Objectives:** On successful completion the subject student should have understand: Manipulation of genes Transfer techniques Expression systems and methods of selection

#### UNIT I:

Restriction and Modification systems of Bacteria. Restriction enzyme: DNA Polymerases, DNALigase, methylase, Taq polymerase, polynucleotide kinase, alkaline phosphotase, revese transcriptase, DNasel, S1nuclease, RnaseH, terminal deoxynucleotidyl transferase, RNA polymerase.

#### UNIT II:

Types and methods in probe construction, methods of labeling gene probes, identification of recombinant DNA. Construction of DNA libraries and genomic libraries, protein engineering.

#### UNITIII:

Introduction of cloned genes into the host cells: Transformation, transduction, Particle gun, electroporation, liposome mediated and co-cultivation.

#### UNIT IV:

Recombinant DNA techniques: Anti sense technology, terminator gene technology, site directed mutagenesis, Human genome project, hybridization techniques-southern, Western and Northern blotting, Chromosome walking. PCR, RFLP, RAPD, DNA finger printing, Micro array and sequencing, gene therapy, DNA sequencing.

#### UNIT V:

Public acceptance issues for biotechnology: Case studies/experiences from developing and developed countries. Biotechnology and hunger: Challenges for the Indian Biotechnological research and industries. The Cartagena protocol on biosafety. Biosafety management: Key to the environmentally responsible use of biotechnology. Ethical implications of biotechnological products and techniques. Social and ethical implications of biological weapons.

#### **REFERENCES:**

- 1. Ernst.L.Winnacker, (2003) from genes to clones, 2nd edition, Panima publishing corporation, NewDelhi.
- 2. James.D.Watson(2001) Recombinant DNA technology,2<sup>nd</sup> edition, WH Freeman and company, New York.
- 3. Glick and Pasternak,(1996),Molecular biotechnology, Panima publishing corporation.NewDelhi.
- 4. BrownT.A., (1998) Introduction to gene cloning, 3rd edition, Stanley Thomas Publishing Ltd, London.
- 5. PrimroseS.B., (2003) Principles of gene manipulation,6<sup>th</sup> edition, Blackwell Science Ltd, Germany.
- 6. Cartagena Protocol on Biosafety, January 2000.

- 7. Biological Warfare in the 21st century, by M.R. Dano, Brassies London, 1994.
- 8. Safety Considerations for Biotechnology, Paris, OECD, 1992 and latest publications.

# Gr C (Application Oriented Subject B) Paper II

# Subject Title: BIOPHYSICS AND BIOINSTRUMENTATION

**Subject description:** This course presents study of Biophysics and Instrument of Biological Importantce.

**Goals**: To make the student to understand the methods to analyze Biomolecules. **Objectives:** On successful completion the subject student should have understand: Analytical methods and Molecular structures.

#### UNIT I:

Conformation of Biological Macromolecules – <u>Structure of Proteins</u> – Primary, Secondary, Tertiary and Quaternary – Composition of proteins – Amino acids – Properties – <u>Structure of</u> <u>Nucleic Acids</u> – Primary, Secondary, Tertiary and Quaternary – Composition of nucleic acids – Nucleoside and Nucleotides – Properties of nucleic acids – Polysaccharides – Lipids.

#### UNIT II:

Forces that stabilizes the Macromolecules – Proteins and Nucleic acids – Hydrogen Bonding – Hydrophobic interactions – Ionic interactions – Disulfide Bonds – Glycosidic Bonds.

#### UNIT III:

Techniques for the study of biological structure – Spectroscopy – Principles and applications of UV, Visible, NMR, Infra red and Raman Spectroscopy. X - Ray Scattering and Diffraction.

#### UNIT IV:

Separation Methods – Chromatography – Affinity, Column, Paper, Thin Layer, Ion Exchange, HPLC, Gel Filtration and GC.

#### UNIT V:

Separation Methods – Sedimentation – Centrifugation – Ultracentrifugation – Diffusion – Macromolecular Diffusion - Electrophoresis – Agarose gel, Native PAGE and SDS – PAGE.

#### **REFERENCES**:

- 1. Biophysical Chemistry Cantor and Schimmel W.H. Freeman and Company, 2001
- Practical Biochemistry Keith Wilson and John Walker Cambridge University Press 5<sup>th</sup> edition, 2003
- 3. Basic Biophysics for Biologist M.Daniel Agrobios (India), 2005.

# **CORE PRACTICAL III:**

#### **APPLIED BIOTECHNOLOGY**

- 1. Fermentor Design and Working Principle
- 2. Organic acid Production -- Citric acid (solid state or submerged).
- 3. Production and assay of extra cellular enzyme Protease—submerged.
- 4. Wine Production.
- 5. Antibacterial sensitivity test.
- 6. Production of an Antibiotic
- 7. Preservation and maintenance of cells.
- 8. Identification of fungal spoilers. (Aspergillums, Mucor).
- 9. Plant tissue culture media preparations.
- 10. In-vitro germination of seeds.
- 11. Callus induction and differentiation.
- 12. Embryo culture.
- 13. Somatic embryogenesis.
- 14. Isolation and fusion of protoplast.
- 15. Artificial seed production.
- 16. Meristem culture.
- 17. Micro propagation.
- 18. Agrobacterium mediated gene transfer --- Demonstration.

# AOS PRACTICAL IV: LAB IN IMMUNOLOGY AND rDNA TECHNOLOGY.

- 1. Preparation of Antibodies.
- 2. Antigen- Antibody Reactions.
- 3. Immuno diffusion. (Single radial, double and rocket)
- 4. Blood grouping.
- 5. Preparation of serum from blood.
- 6. Method of immunization and bleeding.
- 7. ELISA- Demonstration.
- 8. Immuno assay and Typhoid antibodies.
- 9. Salt precipitation of Immunoglobulin.
- 10. Agarose gel Electrophoresis.
- 11. Isolation of Genomic DNA-Bacteria, Plant and Animal.
- 12. Isolation of Plasmid DNA.
- 13. Isolation of RNA.
- 14. Restriction Digestion.
- 15. Isolation of Phage DNA.
- 16. Transformation.
- 17. Southern blotting --- Demonstration.
- 18. Northern blotting --- Demonstration.
- 19. Western blotting --- Demonstration.
- 20. PCR-. Demonstration.
- 21. Construction of restriction Map of plasmid DNA --- Demonstration.

#### **DIPLOMA PAPER IV**

#### Subject Title: PHARMACOLOGY

**Subject description:** This course presents Medicines for different disease **Goals**: To make the student to understand the concept therapy.

**Objectives:** On successful completion the subject student should have understand: Drug administration, drug metabolism and allergy.

#### UNIT I

Pharmacology – origins and antecedents – Pharmacology in the 20<sup>th</sup> century – Drugs – Sources, dosage forms and routes of administration. Absorption, factors modifying drug absorption, distribution, metabolism – Phase I, II reactions, action of cytochrome P450

#### UNIT II

Targets for drug action, receptor proteins, ion channel and drug targets, control of receptor expression, assay of drug potency: Chemical, bioassay and immunoassay-Drug tolerance and drug dependence. Principles of basic Pharmacokinetics, Adverse response to drugs, drug intolerance, drug allergy, tachyphylaxis, drug abuse, vaccination against infection, factors modifying drug action and effect.

#### UNIT III

Mechanism of action of drugs used in therapy of

- a) Respiratory systems cough, bronchial asthma, pulmonary tuberculosis
- b) Cancer chemotherapy
- c) Antimicrobial drugs sulfonamide, trimethoprim, penicillins, aminoglycosides and bacterial resistance.
- d) Thyroid and anti thyroid drugs, insulin and anti diabetic drugs, anti fertility and ovulation inducing drugs.

#### References:

- 1. Pharmacology 5<sup>th</sup> edition H.P.Rang, M.M.Dale, J.M.Ritter, P.K.Moore
- 2. The Pharmacology, Volume I and II Goodman and Gillman
- 3. Basic Pharmacology Foxter Cox. Butterworth's 1980
- 4. Pharmacology and Pharmacotherapeutics R.S.Satoskar, S.D. Bhandhakam and S.S. Alinapure
- 5. Principles of Medicinal Chemistry William O.Foye B.I. Waverks. Pvt. Ltd, New Delhi

# **B.Sc., BIOTECHNOLOGY-SAMPLE QUESTION PAPERS**

	I semeste Core paper I CELI	r BIOLOGV	
Duration: 3 hrs	Core paper 1 CELL	marks: 100	
Section –A	Answer all	(10XI=10)	
<ol> <li>Histones</li> <li>Proteins</li> <li>lysozyme</li> <li>Erythrocyte</li> <li>Chiasma.</li> <li>Inner membrane</li> </ol>	folding towards the matrix in m	itochondria is called	7.the
proteinous part of e	enzyme is called		
8. Micro tubules ar 9 Mitosis provides	e made up of proteins called equilibrium in the amount of	and	in the cell
10. Choromosomes	are shortest and thickest at	phase.	in the cen.
Section B	Answer all	(5*6=30)	
11. a) give a detaile	ed account of nucleic acid preser	nt in cells?	
b) What are the	(or) chemical bonds present in Bior	nolecules?	
12.a) draw a neat la	abeled diagram of mitochondria (or)	and discuss its features	
b) Give a detaile	ed account of nuclear componen	ts with diagram	
13.a) explain heter	ochromatin & polytene chromos	ome	
b) Describe the f Chromosome	(or) eatures of crossing over taking j	place in homologous	
14. a) what is cell-o	cell signaling?		
b) Give a detail	or ed account of microtubules in co	ells with diagram.	
15.a) what is photo	synthesis and explain about the	sit of the synthesis?	
b) What is endo	plasmic reticulum? Explain with	ı diagram.	
Section C	Answer all	(5x12=60)	
16. a) Discuss abou	nt cell cycle?		

or

- b) Describe the characteristic features of cancer.
- 17. a)highlight the importance of B lymphocytes in human body? (or)
  - b) Discuss how protein synthesis occurs in cell?
- 18. a) difference between prokaryotic and eukaryotic cell?
  - (or)
  - b) Discuss on development of cell theory?
- 19. a) explain: i) Proteins ii) carbohydrates iii) lipid
  - (or)
  - b) Give a detailed account of red blood cells?
- 20. a) what is immunity and explain the term with lymphocytes in blood (or)b) What is Apoptosis and how it is related to cell functioning?

		Semester I	
	Core pa	per II	BIODIVERSITY
	Duration 3 hrs		Marks 100
	SECTION: A.	ANSWER ALL	(10 x 1 =10)
1. 2. 3.	Variation of Genes can I When the new spec	within population or within sp be defined as mating of individ ies evolves in geographic isola	becies duals related by common ancestry ation from the parent species is
4.	the preservation of a resource	resents the value of the future rce	information made available through
5.	The process involvi there is degradation of ener	ng energy transformation will gy is called	not occur spontaneously unless
6.	Diatomic		
7.	Succession		
8.	Nitrogen cycle		
9.	Buoyancy		
10.	Biodegradation		
	Section B	answer all	(5 x 6 = 30)
	11 a) Give an account of gl	obal diversity	
		or	

b) Write a note on loss of biodiversity

12 a) Theories of biological classification
b) Write a note on conservation strategies
13 a) Write about classification of bryophytes
b) Write about economic importance of lichens
14 a) Write short notes on i) Energy flow, ii) food chain, iii) food web
b) Write a note on speciation
15 a) Give an account on physio-chemical properties of water
b) Write about biodegradation in ecosystem
Section C Answer All $(5 \times 12 = 60)$
Section CAnswer All $(5 \ge 12 = 60)$ 16 a) Define: biodiversity. Explain the different types of biodiversity.
Section C Answer All (5 x 12 = 60) 16 a) Define: biodiversity. Explain the different types of biodiversity. or b) Write notes on Western Ghats
Section C Answer All (5 x 12 = 60) 16 a) Define: biodiversity. Explain the different types of biodiversity. or b) Write notes on Western Ghats 17 a) Write the methods of DNA finger printing or
Section C Answer All (5 x 12 = 60) 16 a) Define: biodiversity. Explain the different types of biodiversity. or b) Write notes on Western Ghats 17 a) Write the methods of DNA finger printing or b) Write about economic values of biodiversity
Section C Answer All (5 x 12 = 60) 16 a) Define: biodiversity. Explain the different types of biodiversity. or b) Write notes on Western Ghats 17 a) Write the methods of DNA finger printing or b) Write about economic values of biodiversity 18 a) Give broad outlines of classification of Fungi
Section C Answer All (5 x 12 = 60) 16 a) Define: biodiversity. Explain the different types of biodiversity. or b) Write notes on Western Ghats 17 a) Write the methods of DNA finger printing or b) Write about economic values of biodiversity 18 a) Give broad outlines of classification of Fungi or b) Write about Economic importance of Fungi
<ul> <li>Section C Answer All (5 x 12 = 60)</li> <li>16 a) Define: biodiversity. Explain the different types of biodiversity. or</li> <li>b) Write notes on Western Ghats</li> <li>17 a) Write the methods of DNA finger printing or</li> <li>b) Write about economic values of biodiversity</li> <li>18 a) Give broad outlines of classification of Fungi or</li> <li>b) Write about Economic importance of Fungi</li> <li>19 a) Write about interaction between environment and biota</li> </ul>
<ul> <li>Section C Answer All (5 x 12 = 60)</li> <li>16 a) Define: biodiversity. Explain the different types of biodiversity. or</li> <li>b) Write notes on Western Ghats</li> <li>17 a) Write the methods of DNA finger printing or</li> <li>b) Write about economic values of biodiversity</li> <li>18 a) Give broad outlines of classification of Fungi or</li> <li>b) Write about Economic importance of Fungi</li> <li>19 a) Write about interaction between environment and biota or</li> <li>b) Explain about Nitrogen and Sulpher cycle</li> </ul>
Section C Answer All (5 x 12 = 60) 16 a) Define: biodiversity. Explain the different types of biodiversity. or b) Write notes on Western Ghats 17 a) Write the methods of DNA finger printing or b) Write about economic values of biodiversity 18 a) Give broad outlines of classification of Fungi or b) Write about Economic importance of Fungi 19 a) Write about interaction between environment and biota or b) Explain about Nitrogen and Sulpher cycle 20 a) Write about impact of environmental factors on aquatic biota

# Semester II Core paper III BIOCHEMSTRY

#### MODEL QUESTION PAPER

TIME: 3 HOURS

#### MAX. MARKS: 100

(10 X 1 = 10)

# SECTION A CHOOSE THE BEST ANSWER

1. The end of hydrolysis of starch by amylase is			
(a) soluble starch	(b) glucose		
(c) dextrins	(d) maltose		
2. The following carbohydrate is ca	lled animal starch		
(a) glucose	(b) fructose		
(c) sucrose	(d) glycogen		
3. Iodine of an oil shows the extent	of		
(a) polymerization	(b) unsaturation		
(c) molecular size	(d) estrification		
4. One of the following is an unsatu	reated fatty acid		
(a) arachidonic acid	(b) palmitic acid		
(c) stearic acid	(d) acetic acid		
5. Maximum enzyme activity is obs	served at		
(a) acidic p H	(b) neutral pH		
(c) basic p H	(d) optimum p H		
6. The following reaction is charact	eristic of what type of enzymes		
$2 H_2 O \rightarrow 2 H_2 O$	+ O <sub>2</sub>		
(a) Peroxidases	(b) catalases		
(c) dehydrogenases	(d) copper containing oxidases		
7. Which among the following is a	basic amino acid		
(a) asparagine	(b) arginine		
(c) proline	(d) alanine		
8. Flavoprotein contains			
(a) triboflavin	(b) thiamine		
(c) niacin	(d) pyridoxin		
9. A nucleoside is a mixture of			
(a) base + ribose	(b) base + ribose + phosphoric acid		
(c) base + phosphoric acid	(d) ribose + phosphoric acid		
10. Protein synthesis is called			
(a) Replication	(b) transcription		
(c) Translation	(d) termination		

#### SECTION B

(5 X 6 = 30)

ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

11. (a) Explain the structure of sucrose and their properties.

- (b) Briefly describe about the classification of carbohydrate.
- 12. (a) what is PUFA? What are their roles in living system?
  - Or
  - (b) Explain  $\Box$  oxidation of fatty acids.
- 13. (a) differentiate co enzymes and cofactors with an example.

Or

- (b) Derived line weaver Burke equation.
- 14. (a) give an account of the classification of amino acids.

Or

- (b) Write short on zwitter ionic properties of amino acids.
- 15. (a) describe genetic code with a suitable example .
  - Or
  - (b) With a suitable diagram explain the structure and properties of t RNA.
    - SECTION C  $(5 \times 12 = 60)$

# ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

16. (a) Give an detailed account on the structure and the properties of starch.

Or

- (b) Describe the synthesis of purine and pyrimidines .
- 17. (a) What are saturated and unsaturated and describe about the biosysthesis of fatty acids .

Or

- (b) Discuss about the degradation of lipids by alpha, beta and omega oxidation.
- 18 (a) what are the various factors which influence the enzyme activity.
  - Or
  - (b) Derived michelis manton equation and the transfor4mation of minchel menton kinetics , line weaver burke plot?

19. (a) Give an detailed account on the structures and properties of amino acids and write a note on zwitter ionic properties .

#### Or

(b) Describe the different structures of protein and amino acids sequencing of protein.20. (a) Describe the structures of DNA and explain the biosynthesis pathway of purine and pyrimidine.

Or

(b) Give a detailed account on transcription process.

# Semester III Core paper IV MICROBIOLOGY

#### TIME: 3 HOURS

MARKS: 100

# SECTION - A $(10 \times 1 = 10 \text{ MARKS})$

#### Define the following

- 1. TEM
- 2. Oogamy
- 3. Sterilant
- 4. Nutrient Agar
- 5. Resolution
- 6. The penicillin antibiotic is produced by ------
- 7. Ethyl Alcohol is produced by ------
- 8. N<sub>2</sub> fixation is an example for -----
- 9. β-oxidation is the general path way for-----metabolism
- 10. Blood Agar media is an example for -----media

SECTION - B (5 X 6 = 30 MARKS)

11 a) Write short notes on various types of light microscopes

or

- b) Write a general account of microbial diversity
- 12 a) Write short notes on general structure of Bacteria

or

- b) Explain the Economic importance of Yeasts and Moulds
- 13 a) Write a note on chemoautotrophs

or

- b) Write about Symbiosis.
- 14 a) Write about the different types of media

or

- b) Write short notes on different types staining techniques.
- 15 a) Explain fixation of Nitrogen

or

b) Explain about Fermentation.

SECTION - C (5 X12 = 60 marks)

- 16 a) Describe about the different types of electron microscopes. or
  - b) Write about the definition and scope of Microbiology.

17 a) Describe about the growth and reproduction of fungi.

- b) Write an essay on growth and reproduction in viruses.
- 18 a) Write about the microbial pathogens of plants.

or

or

- b) Write an essay on microbial pathogens of animals and humans
- 19 a) Write explanatory notes on different methods of chemical sterilization

r

- b) Describe the structure and composition of bacterial cell wall.
- 20 a) Write about Carbon and Phosphorous cycle.

or

b) Write an essay on production of enzymes and antibiotics.

# Semester III Core Paper V GENETICS

SECTION- A ( $10 \times 1 = 10 \text{ MARKS}$ )

TIME: 3 HOURS

#### MARKS: 100

Define the following

- 1. Col plasmid
- 2. Transposons
- 3. Hardy-Weinberg's law
- 4. Genetic counseling
- 5. Chromosomal aberrations
- 6. The pink flower color of 4'o clock plant is an example for the phenomenon of ------
- 7. The Epistatic ratio is ------
- 8. IS elements is an example of -----
- 9. The unit of recombination in a gene is called------
- 10. ------ is the genetic material.

SECTION -  $(5 \times 6 = 30 \text{ MARKS})$ 

11 a) Write short notes on Independent Assortment

- b) Explain about the Incomplete dominance
- 12 a) Write short notes on Epistasis

#### or

- b) Explain lethal genes and lethality
- 13 a) Write a note on chromosomal variation in number

or

b) Explain genetics of hemoglobin

- 14 a) Write about the fine structure of gene.
  - or
  - b) Write short notes on Experimental evidence for DNA as the genetic material
- 15 a) Explain Genetic counseling
  - or
  - b) Explain about the genetic drift

SECTION - C

(5 X 12 = 60 marks)

16 a) Describe about the chromosomal theory of inheritance

or

- b) Describe about the sex determination mechanism in animals
- 17 a) Describe about the Mendelian principles
  - or
  - b) Write an essay on Multiple allelism
- 18 a) Write about the Transposable elements in prokaryotes
  - or
  - b) Write an essay on chromosomal abberations
- 19 a) Write explanatory notes on relation between genes and polypeptides or
  - b) Describe the F, Col and R plasmids in bacteria
- 20 a) Write about regulation of development in Arabidopsis

or

b) Write an essay on Factors affecting gene frequency

# Semester III Diploma Paper I Human Physiology Model Question Paper

# TIME: 3 HOURS

MARKS: 100

Section A:

Answer All

(10 \* 1 = 10)

Define the following:

- 1. Myosin
- 2. Acetyl choline
- 3. Rhodopsin
- 4. Coklea
- 5. Stem cell
- 6. Plasminogen
- 7. Lymph node
- 8. Assimilation
- 9. Threshold level
- 10. Adrenaline

Section B:	Answer all	(5 * 6 = 30)
11a) Write a b	orief note on muscle contraction or	
b) With fig	gures -Discuss the muscle types	
12 a) With dia	agram discuss the structure of Ear or	
b) Discuss	blood clotting	
13 a) Discuss	the functions of Heart	
b) Discuss	or $P^{H}$ maintenance in the blood system	
14 a) Discuss	lymphatic system with diagrams	
b) With ne	or at diagram explain the digestive system	
15 a) Discuss	the types of Hormones	
b) Discuss	or the functions of insulin	
Section C:	Answer all	(5 * 12 = 60)
16 a) Explain	neuro – muclular junction	
b) Explain	how signals are conducted in neurons.	
17 a) Explain		
	the functions of Brain	
b) Discuss	the functions of Brain or erythropoisis	
b) Discuss 18 a) Discuss	the functions of Brain or erythropoisis ECG	
<ul><li>b) Discuss</li><li>18 a) Discuss</li><li>b) Discuss</li></ul>	the functions of Brain or erythropoisis ECG or gas exchange in the respiratory system	
<ul><li>b) Discuss</li><li>18 a) Discuss</li><li>b) Discuss</li><li>19 a) Discuss</li></ul>	the functions of Brain or erythropoisis ECG or gas exchange in the respiratory system the absorption of carbohydrates in digestive system or	
<ul> <li>b) Discuss</li> <li>18 a) Discuss</li> <li>b) Discuss</li> <li>19 a) Discuss</li> <li>b) Discuss</li> </ul>	the functions of Brain or erythropoisis ECG or gas exchange in the respiratory system the absorption of carbohydrates in digestive system or the functions of Kidney	
<ul> <li>b) Discuss</li> <li>18 a) Discuss</li> <li>b) Discuss</li> <li>19 a) Discuss</li> <li>b) Discuss</li> <li>20 a) Discuss</li> </ul>	the functions of Brain or erythropoisis ECG or gas exchange in the respiratory system the absorption of carbohydrates in digestive system or the functions of Kidney the functions of hypothalamus or	

	Semester IV
Core Paper VI:	BIOINFORMATICS
MODEL QUESTION PAPER	TIME: 3 HOURS
MAX. MARK	S: 100
SECTION A	(10  X  1 = 10)
CHOOSE THE BEST ANSWER	
1. Which of the following is a intern	et domain
(a) · com	(b). Org
(c) . net	(d) all of the above
2. Which of the following is a search	n engine?
(a) alta vista	(b) google
(c) hot bot	(d) all of the above
3. Gen bank data base from	
(a) NCBI	(b) EMBL
(c) MIPS	(d) none of the above
4. SWISS – PROT is a	
(a) DNA sequence database (	b) protein sequence database
(c) Both (a) & (b)	(d) none of the above
5. Genome resources	
(a) TIGR	(b) NCBI
(c) ensembl	(d) all of the above
6. DNA markers	
(a) SNP	(b) RFLP
(c) Both (a) & (b)	(d) none of the above
7. Human genome project officially	begun in
(a) 1985	(b) 1990
(c) 1992	(d) 1993
8 is the analysis of the prote	ein complement expressed by a genome (or) a cell
(or) a tissue type	
(a) Genomics	(b) proteomics
(c) transcriptome	(d) none of the above
9. Which of the following is used for	local similarity searches?
(a) FASTA	(b) MMDB
(c) BLAST	(d) All of the above
10 is the technique of method	ically demonstrating a family relation ship between
Species	
(a) Phylogenetic analysis	(b) Blocks
(c) PRINTS	(d) IDENTITY
SECTION B	(5 X 6 = 30)
ANSWER ALL QUESTIONS, CHO	OSING EITHER (a) OR (b)
11. (a) Write short notes on WW	W.
Or	
(b) Give a note on web brows	jes.

12. (a) Comment on biological database protocol. Or (b) Give a note on pathway database. (a) What do you know about EST. 13. Or (b) Give an account on SAGE. 14. (a) Explain protein – protein interaction. Or (b) Describe mass spectrometry. 15. (a) Comment on drug designing. Or (b) Explain chemoinformatics. SECTION C (5 X 12 = 60)ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b) 16. (a) Discuss about electronic mail. Or (b) Write an essay on search engine. 17. (a) Write an essay on sequence analysis tools. Or (b) Write an essay on primary nucleic acid sequence database. (a) Discuss DNA and RNA microarrays. 18 Or (b) Explain in brief the different method used in human genome mapping. (a) Discuss about 2 D gel electrophoresis. 19. Or (b) Comment on pathway analysis . 20. (a) Explain phylogenetic analysis. Or

(b) Write a note on the following.

(1) Pearl (2) E - cell.

#### Semester IV DIPLOMA PAPER II: HUMAN PATHOLOGY

#### TIME: 3 HOURS

MARKS: 100

#### SECTION - A (10 x 1 =10 MARKS)

- 1. Cirrhosis
- 2. Rotor drugs
- 3. Cystic fibrosis
- 4. Ketotic Hyperglycemic
- 5. Creatine kinase

- 6. ----- Hormone helps in detection of pregnancy
- 7. Myocardial infraction refers to-----
- 8. Proteinuria means-----
- 9. Acromegaly results because of-----
- 10. Jaundice is an example of-----

SECTION - B (5 X 6 = 30 MARKS)

- 11. a. Write short note on disorders of kidney Or
  - b. Write about biotechnological approaches to treatment of liver diseases.
- 12. a. Write short note on Disorders of carbohydrate metabolism.
- Or

b. Write about the significance of Diabetes in pregnancy.

13. a. Write briefly about metabolic aspects of cancer

#### Or

b. Write short note on enzymes which serve as tumor markers

#### 14. a. Explain disorders of hypothalamus Or

b. Write about the Prostate cancer markers.

15. a. Explain disorders of plasma proteins

Ōr

b. Write briefly about Abnormal Haemoglobin derivatives.

SECTION- C  $(5 \times 12 = 60 \text{ Marks})$ 

- a. Write elaborately on inherited abnormalities of bilirubin metabolism or
   b. Write an essay on Thyroid related deficiencies
- 17. a. Write about metabolic complication of diabetes

r

- b. Write elaborately on diagnosis and management of diabetes.
- 18. a. Add a significant note on disorders of adrenal medulla

or

b. Explain the disorders on Adrenal Cortex.

19. a. Write about biotechnological approaches to liver diseases especially in development of vaccines and drugs.

Or

b. Write an essay on the relationship of liver and alcohol

**Core Paper VII** 

20. a. Write in detail about the various growth hormone related deficiencies.

Or

b. Write significantly on hyper prolactinaemia and cushing's disease

#### Semester V

**IMMUNOLOGY** 

Duration 3 hrs SECTION – A Marks 100 10 x 1 = 10

Answer all

- 1. Variolation
- 2. Passive immunity
- 3. IgM
- 4. Hematopoiesis
- 5. Avidity
- 6. RA
- 7. Alveolar macrophages
- 8. Cardinal signs of inflammation
- 9. Alexin
- 10. Allergic reactions

SECTI	ON – B	Answer all	5 x 6 = 30
11.	a) Briefly note on cell	mediated immune response (or)	
	b) Give short notes on	Mechanisms of defense	
12.	a) Explain thymus wit	h a neat diagram (or)	
	b) Briefly explain (i) H	Haptens (ii) Adjuvants	
13.	a) Write in short the m	nechanism of resistance to tumors (or)	
	b) Give a brief account	t on (i) RIA and (ii) ELISA	
14.	a) Explain briefly the	mechanism of phagocytosis (or)	
	b) Clonal selection the	eory – Comment on it	
15.	a) Give a brief accoun	t on (i) Th cell and (ii) Tc cell. (or)	

b) Explain type II hypersensitivity reaction.

SECT	ION – C Answer all	5 X 12 = 60
16.	a) Give a detailed note on history of immunology	
	b) Explain innate and acquired immunity with example	
17.	a) Write notes on types of antibodies and their properties (or)	
	b) With neat schematic diagram, explain secondary lymphoid organ	ns.
18.	a) Explain in detail autoimmunity (or)	
	b) Discuss about primary and secondary immunodeficiency in deta	ail
19.	a) Describe the mechanism of response of B cell to antigens (or)	
	b) Give a detailed account on (i) Interferons (ii) Interleukins	
20.	a) Explain in detail Cell mediated immunity to intracellular bacter (or)	ia and viruses
	b) Give a detailed account on complement synthesis	

# Semester VCore Paper VIIIPLANT BIOTECHNOLOGY

**DURATION: 3 HRS** 

#### SECTION A

- 1. Mass selection.
- 2. Mutation
- 3. callus
- 4. Cybrid
- 5. Vir gene
- 6. R plasmid
- 7. RFLP
- 8. cry protein
- 9. Plant stress response
- 10. Ethylene

# SECTION B

(5X6 = 30)

11a. Write a brief account on the pedigree method of selection:-

MAX MARKS: 100

(10X1 = 10)

b. Explain Mutation

12a. How will you produce virus free plants?

Or

b. Give a brief account on somaclonal variation:-

13a. What do you mean by gene family? Add a brief note on it:-

or

b. Explain briefly how the proteins are targeted to the chloroplast?

14a. Give a brief note on the nif genes:-

or

b. Write a brief account on the opines and its importance:-

15a. how plantibodies are produced?

Or

b. Write brief notes on selectable markers

SECTION C

(5X12 =60)

16a. Write an essay about polyploidy in crop improvement:-

0

b. Write a detailed account on the backcrossing technique and its application in crop improvement

17a. Give a detailed account on the somatic hybridization and its application in crop improvement:-

or b. Write an essay on the various methods of sterilization in plant tissue culture:-

18a. Cytoplasmic Male sterility & its application – Discuss.

Or b. Explain in detail about the Chloroplast genome organization:-

19a. How will you produce transgenic plants?

Or

b. Explain the various physical methods of gene transfer in plants

20a. Describe in detail on the herbicide resistance transgenic plants:-

or

b. write a detailed account on the pest resistance transgenic plants

**Core Paper IX** 

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#### Semester V : ANIMAL BIOTECHNOLOGY

MODEL QUESTION PAPER TIME: 3 HOURS

#### MAX. MARKS: 100

SECTION A

(10 X 1 = 10)

# CHOOSE THE BEST ANSWER

1 refers to the culture of di	spersed cells obtained from original tissue or from a cell
Line	
(a) Cell culture	(b) cell line
(c) Both (a) & (b)	(d) none of the above
2 refers to the irrerversible	e loss of specialized properties of cells when they are
cultured invitro	
(a) Dedifferentiation	(b) redifferentiation
(c) Differentiation	(d) none of the above
3. The end of the proliferate life spa	n of cells is refers to as
(a) Senescence	(b) apoptosis
(c) Cell line	(d) none of the above
4. The process of programmed cell	death is refers to as
(a) Senescence	(b) apoptosis
(c) Both (a) & (b)	(d) none of the above
5. Vaccine are	
(a) Dead bacteria	(b) attenuated bacteria
(c) Viral fragments	(d) all of the above
6. Treatment of disease by use of ge	ene or DNA sequences
(a) Gene therapy	(b) genetic immunization
(c) Gene silencing	(d) none of the above
7. Cryopreservation broadly means	the storage of germ plasm
(a) Over solid CO <sub>2</sub> (at -79 $^{\circ}$	C)(b) low temperature deep freezers at ( at -80 $^{\circ}$ C)
(c) In liquid nitrogen ( at -19	$6^{\circ}$ C) (d) all of the above
8. Application of embryo culture	
(a) Prevention embryo abort	ion (b) overcoming seed darmency
(c) Production of haploids	(d) All of the above
9. The target gene responsible for th	e development of transgenic organisms.
(a) Transgene	(b) transgenic
(c) Both (a) & (b)	(d) none of the above
10. The first animals used for transg	genic was a
(a) Mouse	(b) goat
(c) Dog	(d) horse
	(5 M ( 20))
SECTION B	$(3 \land 0 = 30)$

ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

11. (a) Write short notes on characterization of cultured cells.

Or

- (b) Comment on animal cell culture media.
- 12. (a) Give a note on primary cell culture.

Or

- (b) Write an account on cell growth kinetics.
- 13. (a) Explain electrophoration.

Or

- (b) Explain in brief on gene therapy.
- (a) Explain the molecular events during fertilization.. 14.

Or

- (b) Write in brief about cytokines.
- 15. (a) Write in detail about embryo culture.

Or

(b) Give a note on blood clotting factors.

SECTION C (5 X 12 = 60)

#### ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

16. (a) What are the facilities required to set up animal cell culture laboratory.

Or

- (b) Comment on cell synchronization and senescence.
- (a) Write an essay on stem cell culture. 17.

Or

- (b) Write an essay on crypreservation.
- 18 (a) comment on human genome project. Add a note on major highlights of human Genome.

Or

- (b) Write an essay on vaccine.
- (a) explain the genetic regulation during embryonic development.

Or

- (b) Briefly explain collection and preservation of embryo.
- 20. (a) write an essay on transgenics and their applications. Or

(b) Write short note on the following.

(1) Hormone

19.

(2) Plasminogen

# Semester V

# (AOS-A)- Paper I

# **MOLECULAR GENETICS**

(10X1 =10)

**DURATION: 3 HRS** 

# SECTION A

MAX MARKS: 75

1. DNA lielicase	
2. One complete set of DNA is called as	
a) haploid b)diploid c) geneome	d) none
3. Explain genetic code	,
4. Explain operon	
5. Transit peptide	
6. Translation	
7. The resultant organism from mutation is	called
a) Mutant b) mutation c) mutagen	d) all of these
8. What do you mean by Linkage?	,
9. The material which induces mutation is	termed as
a) Mutant b) mutation c) mutagen	d) all of these
10. Explain gene knockout	
SECTION B Answer all	(5X5 =25)
11a. Comment on the semi-conservative type	pe of DNA replication:-
	or
b. Describe the structure of tRNA with il	lustration:
12a. Elaborate the rho dependent termination	on of transcription:-
12a. Elaborate the rho dependent termination	on of transcription:- or
<ul><li>12a. Elaborate the rho dependent termination</li><li>b.: Write a brief account on 'Genetic cod</li></ul>	on of transcription:- or e':-
<ul> <li>12a. Elaborate the rho dependent termination</li> <li>b.: Write a brief account on 'Genetic cod</li> <li>13a. Protein export – Discuss</li> </ul>	on of transcription:- or e':-
<ul><li>12a. Elaborate the rho dependent termination</li><li>b.: Write a brief account on 'Genetic cod</li><li>13a. Protein export – Discuss</li></ul>	on of transcription:- or e':- or
<ul> <li>12a. Elaborate the rho dependent termination</li> <li>b.: Write a brief account on 'Genetic cod</li> <li>13a. Protein export – Discuss</li> <li>b. Discuss RNA Processing</li> </ul>	on of transcription:- or le':- or
<ul> <li>12a. Elaborate the rho dependent termination</li> <li>b.: Write a brief account on 'Genetic cod 13a. Protein export – Discuss</li> <li>b. Discuss RNA Processing</li> <li>14 a. Explain briefly photoreactivation:-</li> </ul>	on of transcription:- or e':- or
<ul> <li>12a. Elaborate the rho dependent termination</li> <li>b.: Write a brief account on 'Genetic code</li> <li>13a. Protein export – Discuss</li> <li>b. Discuss RNA Processing</li> <li>14 a. Explain briefly photoreactivation:-</li> </ul>	on of transcription:- or e':- or or
<ul> <li>12a. Elaborate the rho dependent termination</li> <li>b.: Write a brief account on 'Genetic code</li> <li>13a. Protein export – Discuss</li> <li>b. Discuss RNA Processing</li> <li>14 a. Explain briefly photoreactivation:-</li> <li>b. Give a brief account on 'Induced mutanged account on 'Induced m</li></ul>	on of transcription:- or le':- or or tion' :-
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<ul> <li>12a. Elaborate the rho dependent termination</li> <li>b.: Write a brief account on 'Genetic code</li> <li>13a. Protein export – Discuss</li> <li>b. Discuss RNA Processing</li> <li>14 a. Explain briefly photoreactivation:-</li> <li>b. Give a brief account on 'Induced mutae</li> <li>15a. Explain briefly the transformation:-</li> <li>b. Write a brief account on gene targeting</li> </ul>	on of transcription:- or e':- or or tion' :- or ;:-
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17a .Give a detailed account on lac operon:-

b. Write a detailed account on the ;Elucidation of genetic code':-

18a. Explain protein synthesis in detail :-

or b. Give a detailed account t on the post translational modifications:-

19a. Elaborate the Ames test :-

or

or

b. Comment on the spontaneous mutations:-

20a. Write a detailed account on conjugation:-

(AOS\_A) Paper II

or

b. Elucidate the various steps involved in chromosome mapping:-

# Semester V BIOLOGY OF CLONING VECTORS

DURATION : 3 HRS

MAX MARKS: 75

#### SECTION A

1.

- episome
- 2. vector
- 3. cosmid
- 4. concactamers
- 5. Yip
- 6. Lysate
- 7. probe
- 8. FISH
- 9. cDNA Library
- 10. Magic Box

# SECTION B

# (5X5 = 25)

(10X1 = 10)

11a. Describe the structure of plasmid pUC 18:-

or

b.Discuss Construction of PBR 322

- 12a. Explain briefly on M13 vector:-
- or

b. Explain cosmid and its imoptance in rDNA technology

13a. Explain Yep:-

or

b. Write a brief account on the cloning in Bacillus:-

14a. How will you produce a cDNA? or b. Explain chromosome walking:-15a. How will you check the safety of a rDNA laboratory? b.Discuss Clical applications of rDNA technology SECTION C (5X8 = 40)16a. Write a detailed account on the pBR 322 and its importance in rDNA technology:or b. Expression vectors in prokaryotes – Explain 17a. Write a detailed account on SV40 viral vector:or b. Give a detailed account on Ti plasmids:-18a. Explain YAC or b. Cloning in streptomyces – Discuss 19a. How will you construct a genomic library? or b. Comment on the site directed mutagenesis:-20a. Give a detailed account on Human Genome project:or b. Write a detailed account on the gene therapy:-

#### Semester V Diploma paper III: Diagnostic tools

SECTION – A 1. Serum

- 2. Coagulation
- 3. Easinophil

Duration 3 hrs

- 4. Nephron
- 5. Dialysis
- 6. Aciduria
- 7. threshold level
- 8. graft
- 9. PPLO
- 10. Radio immuno assay

Section B

Answer all 5 \* 6 = 30

Marks 100

Answer all

 $10 \ge 1 = 10$ 

11 a) what do	you mean by packed cell volume?	
b) Discuss	Anemia	
12 a) Discuss	differential counting	
b) Discuss	the functions of Basophiles	
13 a) Discuss	the physical analysis of urine or	
b) Discuss	absorption and re-absorption in Kidney	
14 a) Discuss	the functions of blood	
b) Discuss	the types of grafts	
15 a) Write a	brief note on Lymphocytes	
b) Discuss	the alignment of DCD	
0) 2150455	the chinical applications of PCK	
Section C:	Answer all	5 * 12 = 60
Section C: 16 a) Explain	Answer all the biochemistry of blood coagulation or	5 * 12 = 60
Section C: 16 a) Explain b) Discuss	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams.	5 * 12 = 60
Section C: 16 a) Explain b) Discuss 17 a) Explain	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or	5 * 12 = 60
Section C: 16 a) Explain b) Discuss 17 a) Explain b) Explain	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or the extraction and analysis of CSF	5 * 12 = 60
<ul> <li>Section C:</li> <li>16 a) Explain</li> <li>b) Discuss</li> <li>17 a) Explain</li> <li>b) Explain</li> <li>18 a) Discuss</li> </ul>	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or the extraction and analysis of CSF Haemoglobinopathies or	5 * 12 = 60
Section C: 16 a) Explain b) Discuss 17 a) Explain b) Explain 18 a) Discuss b) Discuss	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or the extraction and analysis of CSF Haemoglobinopathies or ELISA	5 * 12 = 60
Section C: 16 a) Explain b) Discuss 17 a) Explain b) Explain 18 a) Discuss b) Discuss 19 a) Discuss	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or the extraction and analysis of CSF Haemoglobinopathies or ELISA the methods in Blood banking or	5 * 12 = 60
Section C: 16 a) Explain b) Discuss 17 a) Explain b) Explain 18 a) Discuss b) Discuss 19 a) Discuss b) Discuss	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or the extraction and analysis of CSF Haemoglobinopathies or ELISA the methods in Blood banking or the mechanism of Graft rejection	5 * 12 = 60
Section C: 16 a) Explain b) Discuss 17 a) Explain b) Explain 18 a) Discuss b) Discuss 19 a) Discuss b) Discuss 20 a) Discuss	Answer all the biochemistry of blood coagulation or the types of Blood cells with diagrams. the importance of urine analysis during infection. or the extraction and analysis of CSF Haemoglobinopathies or ELISA the methods in Blood banking or the mechanism of Graft rejection the examination of AIDS and its importance.	5 * 12 = 60

b) How the computers can be useful in a clinical laboratory on

# Semester VICore Paper XMICROBIAL BIOTECHNOLOGY

Duration 3 hrs

SECTION – A

Marks 100

 $10 \ge 1 = 10$ 

Answer all

- 1. Microbe
- 2. Organic acid
- 3. Vaccine
- 4. Lipase
- 5. Fermentation
- 6. Lactic acid bacteria
- 7. Glutamine
- 8. Tetracycline
- 9. Xenobiotic
- 10. Biofertiliser

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SECTION – B
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 $5 \ge 6 = 30$ 

#### Answer all

11.	a) Briefly note on the horizons of microbial biotechnology	
	b) Give an account on the applications of microbial biotechnology	
12.	a) Briefly note on synthetic vaccines (or)	
13.	<ul><li>b) Give short notes on production of proteins in yeast</li><li>a) Explain the production of ethanol from feed stock</li></ul>	
	b) Give a protocol for the production of lactic acid	
14.	a) Briefly note on antifungal agents (or)	
	b) Give a short note on antitumour antibodies	
15.	a) Describe the role of microbes in removal of heavy metals from efflue (or)	ents
	b) Give a brief note on nitrogen fixing bacteria	
SECTI	ON - C 5 X 12 = 60	
	Answer all	

16. a) Scope of microbial biotechnology – Comment on it

(or)

b) Give a detailed note on the public concerns of microbial biotechnology

17.	a) Microbes are living factories of macromolecules - Explain (or)
	b) Explain with an example Microbial Insecticides
18.	a) Acetic acid production (or)
	b) Describe about the industrial production of various diary products
19.	a) Describe in detail the production of aminoacids (or)
	b) Discuss the large scale production of antibiotics
20.	a) Explain the role of microbes in xenobiotics degradation (or)
	b) Discuss about the production of SCP with an example

# Semester VICore Paper XIENVIRONMENTAL BIOTECHNOLOGY

TIME: 3 HOURS

MARKS: 100

SECTION - A  $(10 \times 1 = 10 \text{ MARKS})$ 

- 1. Global Warming
- 2. Chipko movement
- 3. Gene Bank
- 4. Antagonism
- 5. Food web
- 6. Acid rains refers to-----
- 7. Cryopreservation is-----
- 8. Trophic level refers to-----
- 9. FAO is -----
- 10. The word Pollution refers to------

# SECTION- B $(5 \times 6 = 30 \text{ MARKS})$

11. a. Write about the biotic factors that affect ecology

or

- b. Write briefly about Symbiosis
- 12. a. Write about the pond ecosystem.

#### Or

b. Explain the pyramid of biomass

13. a. Write about the role of domestic live stock

or

b. Add a note on ethical and aesthetic values of biodiversity.

or

14. a. Write about Air pollution

or

- b. Add significantly on water pollution
- 15. a. Explain briefly about National park
  - b. Explain briefly about Chipko movement

SECTION - C  $(5 \times 12 = 60 \text{ Marks})$ 

16. a. Add a significant note on parasitism and predation

or

- b. Explain the role of Abiotic factors affecting ecology.
- 17. a. Explain the Biogeochemical cycle of N & P.

or

b. Write short note on a. Food web

#### b. Food chain

#### c. Energy Flow

18. a. Explain ozone and its importance

or

b. Write an essay on Environmental Assessment and monitoring systems.

19. a. Explain the plant Bioresources.

Or

- b. Write an essay on intellectual property Rights.
- 20. a. Write in detail about In-situ and Ex-situ conservation Strategies.
- Or
- b. Write an essay on wild life Sanctuaries

# Semester VI (AOS B) Paper I: RECOMBINANT DNA TECHNOLOGY AND BIO ETHICS

#### MODEL QUESTION PAPER TIME: 3 HOURS

MAX. MARKS: 75

SECTION A (10 X 1 = 10) CHOOSE THE BEST ANSWER

1.	ECO R1 restriction site is	
	(a) GAATTC	(b) GG CC
	(c) CCGG	(d) GGCCGG
2.	The main sources of DNA ligase	is
	(a) Ø x 174	(b) E.coli
	(c) T 4 phage	(d) neurospora
3.	Protein engineering is carried out	by
	(a) site directed mutagenesis	(b)Laser beam
	(c) gamma radiation	(d) dimmer formation
4.	Which of the following is used as	probe
	(a) single stranded DNA	(b) Double stranded DNA
	(c) both (a) & (b)	(d) none of the above
5.	Transduction is mediated by	
	(a) phages	(b) bacteria
	(c) both (a) & (b)	(d) none of the above
6.	Which of the following technique	is used for direct DNA transfer
	(a) particle gun	(b) transformation
	(c) translation	(d) none of the above
7.	Antibody is used as probe in	
	(a) northern blotting	(b) southern blotting
	(c) western blotting	(d) none of the above
8.	Southern blotting technique is dev	reloped by
	(a) southern	(b) korn berg
	(c) craig venter	(d) All of the above
9.	discovered polymerase cha	in reaction.
	(a) kary mullis	(b) francis collins
	(c) southern	(d) korn berg
10	DBT is located at	
	(a) new delhi	(b) bombay
	(c) culcutta	(d) hydrabad
SE	ECTION B	$(5 \times 5 = 25)$
A	NSWER ALL QUESTIONS, CHC	OSING EITHER (a) OR (b)
11	. (a) Write short notes on poly	nucleotide kinase.
	(b) Write a brief note on the	nomenclature of restriction enzymes.
12	. (a) Comment on genomic lib	rary.
	Or	, see the second se
	(b) Write an account on prob	e construction.
13	. (a) Explain transformation.	
	Or	
	(b) Explain in brief particle g	un bombardment.

(a) Give a note on 'antisence technology'.. Or 14.

- (b) Describe in brief about terminator gene technology.
- 15. (a) Write in detail about biosafety management.

Or

(b) Write in detail about issues in biotechnology.

# SECTION C (5 X 8 = 40)

#### ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

16. (a) Write an essay on DNA manipulative enzymes.

Or

- (b) Write an essay on restriction and modification systems of bacteria.
- 17. (a) Explain the construction of DNA libraries.

#### Or

- (b) Write an essay on protein engineering.
- 18 (a) Write a note on the following.
  - (1) Electroporation
  - (2) Liposome mediated transfer.

Or

- (b) Write an essay on co cultivation.
- 19. (a) Discuss about southern transfer technique.

Or

- (b) Write about DNA sequencing methods.
- 20. (a) Write an essay on social and ethical issues of biotechnology.

Or

(b) Give your view about biotechnology and hunger.

# Semester VI(AOS \_ B) Paper IIBIOPHYSICS AND BIOINSTRUMENTATION

Answer all

Duration 3hrs SECTION – A Marks 75 10 x 1 = 10

1. Nucleoside

- 2. Secondary structure of protein
- 3. Covalent bond
- 4. Ionic interaction
- 5. Beer Lambert's law
- 6. X ray diffraction
- 7. Ninhydrin
- 8. HPLC
- 9. Sedimentation
- 10. Agarose

SECTI	ION – B	5 x 5 = 25
	Answer all	
11.	a) Briefly describe the primary structure of proteins (or)	
	b) Give the structure and bond formation of (i) Ade	nine and (ii) Thymine
12.	a) Give a short note on Van der Waal's interaction (or)	
	b) Write briefly about Ionic interactions	
13.	a) Explain Beer – Lambert's law (or)	
	b) Comment of X- ray scattering	
14.	a) Gel Filtration chromatography – Comment on it (or)	
	b) Give notes on thin layer chromatography	
15.	a) Write notes on agarose gel electrophoresis (or)	
	b) Give short notes on Ultracentrifugation	
SECTI	ION – C	5 X 8 = 40
	Answer all	
16.	a) Give a detailed note on the protein structure (or)	
	b) Describe in detail the structure of nucleic acid	

- 17. a) Discuss about the forces that stabilize the protein structure
  - (or) b) Describe the forces that stabilize nucleic acid structure
- a) Explain the principle and working of NMR spectroscopy (or)
  b) Explain the principle and working of Mass spectroscopy
- a) Write notes on (i) HPLC (ii) GC (or)
  b) Give a detailed on affinity chromatography
- 20. a) Give a detailed note on SDS- PAGE (or)
  b) Explain the methods of (i) Diffusion (ii) Sedimentation

# Semester VI Diploma Paper IV: PHARMACOLOGY

# MODEL QUESTION PAPER TIME: 3 HOURS

MAX. MARKS: 100

SECTION A	(10  X  1 = 10)
CHOOSE THE BEST ANSWER	

1. The study of mechanism of action	n and action of drug is called	
(a) pharmaco kinetics	(b) pharmaco vigilance	
(c) pharmaco economics	(d) pharmaco dynamics	
2. Receptor serves as		
(a) Recognition molecule	(b) signal transmission	
(c) both (a) & (b)	(d) none of the above	
3. Which of the following is favouri	ing maximum absorption?	
(a) Digoxin	(b) Digitoxin	
(c) Both (a) & (b)	(d) none of the above	
4. Which of the following are involved	ved as primary drug target?	
(a) Enzymes	(b) receptors	
(c) Ion channels	(d) all of the above	
5. The uses of bioassay		
(a) to measure drug toxicity a	and unwanted effects	
(b) to investigate the function	n of endogenous mediators	
(c) both (a) & (b)	(d) none of the above	
6. The main roots of drug administr	ation	
(a) Oral	(b) sublingual	
(c) Rectal	(d) all of the above	
7. Penicillin G act by		
(a) Inhibit protein synthesis	(b) remove intracellular calcium	
(c) Inhibit DNA synthesis	(d) inhibit cell wall synthesis	
8. Hormone insulin is secreted from		
(a) $\beta$ - islet cells of pancreas	(b) $\alpha$ - islet cells of pancreas	
(c) $\gamma$ - islet cells of pancreas	(d) $\delta$ - islet cells of pancreas	
9. Sulfonamides act by	.,	
(a) Inhibiting cell wall synthe	esis (b) inhibiting protein synthesis	
(c) Inhibiting folate synthesis	s (d) inhibiting DNA gyrase	
10. Drug which induce cancer are ca	lled	
(a) Teratogenic agents	(b) hypersensitive drugs	
(c) carcinogenic drugs	(d) ototoxic drugs	
$\sim$		

#### SECTION B $(5 \times 6 = 30)$

#### ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

11. (a) Explain phase I and phase II reactions of drug metabolism in detail.

Or

(b) Discuss the design of pro drug.

12. (a) Discuss the role of human hepatic cytochrome P 450 enzyme system in drug metabolism

Or

(b) Discuss various novel drug delivery systems.

13. (a) Discuss the theories of drug receptor interaction .

#### Or

(b) Describe the various methods of bioassay.

14. (a) Describe the structure of G protein coupled receptor and discuss the role protein Or

(b) Write short on tachyphylaxis.

15. (a) Write an essay on the suffonamide .

Or

(b) Write an essay on drug dependence.

### SECTION C (5 X 12 = 60)

#### ANSWER ALL QUESTIONS, CHOOSING EITHER (a) OR (b)

- 16. (a)Discuss the various process of drug absorption and factors affecting the absorption. Or
  - (b) Discuss the factors modifying the effect of drugs
- 17. (a) Discuss the factors which affect the drug action .

#### Or

- (b) Explain in detail about the excretion of drugs.
- 18 (a) Discuss the factors which affect the drug action.

Or

- (b) Explain in detail about the excretion of drugs.
- 19. (a) discuss the mechanism and uses of various penicillin. Or

(b) Discuss the drug treatment of diabetes mellitus.

20.

(a) write an essay on cancer chemotherapy .

#### Or

(b) Comment on anti fertility and ovulation inducing drugs.