Sample Paper ive examination M.Sc. Biotechnology Institute of Molecular Biology and Biotechnology (IMBB)

udent Name:	Time: 60 min	Total Marks: 60	
,011 P.O.,	d Frankly	Hing is not allowed	лткоги т 950 бу.
1717000	Cutting and ove	r-writing is not allowed	
very used	a feach oming acid are determined	9) Viruses exhibit several different inter	active strategies in
1) The unique properties of each amino acid are determined		the host cell. Which of the following does not describe one	
by its particular	is saids b) P group	of those strategies?	
a) Number of bonds to other amino acids b) R group c) Kinds of pentide bonds d) Presence of hydrogen		a) The virus takes over the host cell me	tabolism and
c) Kinds of pept	tide bonds d) Presence of hydrogen	destroys the cell	
	a company that is induced by	b) The virus does not take over the cell	's metabolism and
2) The process of	of uptake of free DNA that is induced by	leaves the cell without killing it	
subjecting bacte	eria to high voltage electric field in the	c) The virus integrates its nucleic acid in	nto that of the host
presence of DN	A is known as	cell and lives harmoniously with the cel	I for a long time
a) Conjugation	b) Electroporation	d) The virus takes over the host cell me	etabolism but does
c) Recombination	on d) None of these	not reproduce itself	. 9 6. 4
	institution a principalism of Denatured by	e) The virus integrates its nucleic acid	into that of the host
3) Bacillus thu	ringiensis is used to control	cell, but eventually lyses the host cell u	pon exiting from it
a) Nematodes	b) Fungal pathogens	cent, but eventually 19565 and 11-1	Arough
c) Bacterial pat	hogens d) Insect pests	10) Biolistics (Gene-gun) is suitable for	NISMIR TO SEE
	factor management and factors are to come as a factor of the	a) Constructing recombinant DNA by j	oining with vectors
4) The rate of n	nigration of DNA within an agarose gel in the	b) Transformation of plant cells	
gel electrophor	esis technique is primarily based on what	c) Disarming pathogen vectors	
factor?		d) DNA finger printing	
a) The negative	e charge of the DNA	a) DNA tinger printing	
b) The number	of DNA fragments	11) Many physicians prefer mother's m	ilk over cow's milk
c) The size of t	the wells of the gel	for infants. This is because mother's m	ilk has the following
d) The size of t	the DNA fragments	for infants. This is because mother's mile	ink has the following
		property not found in cow's milk	
5 You are worl	king in a government laboratory in charge of	a) Antibodies against human disease	
dectroving bio	terrorism materials. An envelope containing	b) Proteins for growing human muscle	
the chare-form	ing bacterium anthrax must be disposed of	c) More essential amino acids	o
cafely What w	would be the best method for ensuring the	d) MHCs for maturing the infant's im-	nune system
anthrax is kille	ed? Ab are an instable for controlling	e) B cells	rather than Children
a) Autocla		12) Detection of GMOs accomplished	Dyblot
b) Dry He		4)1011	outhern blot
c) Disinfe		d) a and b e) a and c f) b	and c
	ing Spray	- 36) The first francismo copy was	
- 1	Soaking	13) A researcher is maintaining a liqui	d culture of bacteria at
e) Bleach	Southing	exponential growth. The pump provid	ing nutrients to the
() The technic	que that utilizes probes to detect specific DNA	culture breaks down over the weekend	l, allowing culture to
6) The technic	nown as what?	run out of nutrients. It is very likely th	at upon return to the
sequences is k	ot b) Northern blot	lab the researcher will find the culture	in which of the
a) Western blo		following stages of growth?	analysis
c) Southern b	of Lastern olov	a) Continuous	o) Log
=: 0 :11' · · ·	n of tissue culture medium is done by	c) Lag	d) Stationary
7) Sterilizatio	If Of tissue culture mediani is done of	many different sequences are possible?	
a) Mixing the	medium with antifungal agents	14) The plant tissues have high rates of	of cell division and
b) Filtering th	ne medium through fine sieve	either concentration or production rec	uired growth regulating
c) Autoclavin	g of medium at 120° for 15 min	substances including	
d) Keeping th	ne medium at –20°C	a) Auxine b) (Cytokinine
**************************************	1 - DNA transcript for a gang	c) Both Auxine and cytokinine d) (Gibralline
8) A scientist	has a processed mRNA transcript for a gene		
he/she wants	to clone into a bacterial vector. What must	15) Differentiation of organs and tiss	ues in a developing
he/she do as	a first step in this process?	organism is associated with	
a) Use PCR t	o create a cDNA molecule	a) Differential expression of genes	
b) Generate p	orimers to the processed mRNA	b) Differential expression of protein	
c) Sequence	the mRNA transcript	c) Both (a) and (b)	None of the above
d) Ligate the	mRNA into the cloning vector	c) Dour (a) and (b)	
			-3

component isolated f	owing terms is used to describe the rom a plant, for in vitro culturing in the	28) RNA primer attracts a) Ligase	DNAb) Polymerase	er Effects van ber
specific medium? a) Synthetic seeds	b) Embryoid	c) Halicase	d) all	
c) Callus		29) Polymerase chain rea		
17) Two microbes	and found to be	a) Kary Mullis c) Miescher	b) Griffi d) Avery	
very useful in genetic a) E.coli and Agroba	engineering.	30) 4 1: : 1:		
b) Plasmid and Agrob	actrium tumefaciens	30) A dividing and undif		of cells is called
c) E. coli and Aspirgi			b) Embryo	
d) Algea and E.coli	general in DNA can be case to	c) Explant	d) Zygote	
10) 0 1 1	b) Election microscore	31) In plants, DNA is for	ınd in	
18) Cryl endotoxins	obtained from Bacillus Thuringiensis	a) Nucleus	b) Plastic	
are effective against:		c) Mitochondria	d) All of	the above
a) Flies		SAY ON HOUSE HOUSE HER AT HE		
c) Boll worms	d) Nematodes	32) Genetic information		
10) The atmesture of a	en be used to introduce DNA mor		b) protein	
a) Presence of oxygen		c) Chromosome	d) RNA	
c) Presence of carbon		33) Somatic hybridization	n is achieved thro	ugh
d) The polar bonds of	water molecules		b) Protoplast fusion	
		c) Conjugation	d) Recombinant I	ONA technology
20) A method that co	unts only live microbes is the			Unit essential all a
a) Pure culture meth		34) The enzymes required	d to obtain wall-fr	ee/naked
c) Turbidity method	d) cell counting method	protoplasts are		
21) Maria	sit's newpord by rac along hast.	 a) Cellulase and proteinas 		and pectinase
21) Meristem culture	is practiced in horticulture to get	c) Cellulase and amylase	d) Amylase	and pectinase
c) Haploids	us b) Somaclonal variation	the production of the second		
	d) Virus-free plants	35) Why does the Enviro monitor the release of tra	nmental Protectio nsgenic bacteria u	n Agency closely sed for
22) Micropropagation	is a technique for production of	agricultural purposes?		
a) True to type plants		a) They want to monitor to	the destruction of	crops by GMOs.
c) Somatic hybrids	d) Somaclonal plants	b) They want to observe tc) They want to ensure th	the effect the GMO	Os have on crops
23) The protein produ	acts of the following Bt toxin genes cry	environment and pose a t	hreat to humans	Tomerate in the
IA c and cry II Ab are	responsible for controlling	d) They want to ensure th	at people are awa	re that GMOs
a) Moth	b) Fruit fly	may have played a role in	the production of	f a particular food
c) Bollworm	d) Roundworm	product.	EDNA indecutes	with hear as a
24) A mixture of orga	nisms is in a pool of seawater. Which	36) The first transgenie of	CYNER molecular	
organism will have a	growth advantage as the water	a) Pea b) Tobac	cco c) Flax	d) Cotton
evaporates?	S 49 1000	a) 1 ca	c) Hax	d) Cotton
a) neurophiles	b) aerotolerant anaerobes	37) All methods of DNA	fingernrinting der	end on some
c) mesophiles	d) osmotolerant prokaryotes	variation of what strategy		cha on some
	The Hartseini rolls		b) GMOs	cas ?
25) In genetic enginee	ering, a chimera is		d) Microarray ana	lvsis
a) An enzyme that lin		C) THE TEXAMORETE	STAR OF	10.50
b) A plasmid that con		38) A tripeptide contains	glycine, alanine, a	and serine. How
c) Virus that infects b	acteria d) A fungi	many different sequences a) 3 b) 4	are possible?	
26) Problems in obtai	ning large amounts of proteins encoded		c) 5	d) 6
by recombinant genes	can often be overcome by using	39) Transgenic plants are	developed by	
a) BACS	b) Expression vectors	a) Introducing foreign ger	nes	
c) YACS	d) all of these	b) Introducing gene muta	tions	
	The company south to represent the particles of	c) Deleting certain chrom	osomes parts	
	ology, Polyethylene glycol method is	d) Stopping spindle forma		
used for		and boundaries of the period gr	thesis hade up of	
a) Energy production		40) Repressor molecules	bind to the:	
b) Gene transfer with		a) Promoter	o) Enhancer	
c) Biodiesel productio	on d) Seedless fruit production	c) Operator	d) Hormone respon	nse element

41) Somatic gene therapy could potentially correct a genetic 51) which of these genetic markers is most likely to be highly polymorphic (have many different alleles)? a) Affected individual and his or her off spring only a) An RFLP b) A microsatellite b) Affected individual and all his or her descent c) An SNP d) All of these c) Parents of affected child d) Affected individual only 52) Plasmid has almost 20 different reorganization sites for-42) Agriculture by using only biofertilizers is called a) Restriction endonucleases b) restriction exonucleases a) Manuring b) Composting c) a and b d) None of these c) Inorganic farming d) organic farming 53) Which vehicles are often used for gene therapy to carry a 43) Nucleotide arrangement in DNA can be seen by healthy gene? a) Light microscope b) Electron microscope a) BAC c) X-Ray crystallography h) YAC d) Ultracentrifuge c) Bacterial Vectors d) Viral Vectors 44) Agrobacterium tumefaciens is -----54) Why might use of microsatellites in genetic mapping a) A disease in humans that causes loss of sight studies be an advantage over RFLPs? b) A bacterium that can be used to introduce DNA into a) Microsatellites are easier to detect. plants b) Microsatellites are more abundant than RFLPs. c) A fungi that is used to produce antibiotics in large c) Microsatellites have more potential alleles than RFLPs. amounts d) All of these. d) Disease in humans that causes loss of weight 55) The addition of a phosphoryl group to a OH group of a 45) Good cloning vectors must possess all but which of the sugar by a kinase is a form of: following qualities? a) Dehydrogenation a) They should possess their own origin of replication b) Reduction c) Oxidation d) Esterification b) They should be readily accepted by the cloning host c) They should be easily manipulated 56) In gene therapy, in order to be successful, the healthy d) They should be capable of carrying a significant piece of gene inserted into a target cell must donor DNA a) Take over and kill the defective gene e) They should be resistant to restriction endonucleases b) Be inserted manually into the cell's mitochondria c) Become attached to the cell's mRNA molecules 46) A molecular marker which is amplified by PCR and is d) Be able to make the correct amount of the protein needed polymorphic by length is a(n): a) Restriction fragment length polymorphism. 57) Plasmids are suitable vectors for gene cloning because b) Variable number of tandem repeats site. a) These can shuttle between prokaryotic and eukaryotic c) Amplified fragment length polymorphism. cells d) Single nucleotide polymorphism. b) These are small circular DNA molecules with their own replication origin site 47) A nucleoside is a combination of c) These are small circular DNA molecules, which can a) Nitrogen base+sugar b) Nitrogen base+phosphate integrate with host chromosomal DNA c) sugar+phosphate d) none d) These often carry antibiotic resistance genes 48) Restriction endonucleases are most widely used in 58) Linkage mapping can determine the distance between recombinant DNA technology. They are obtained from which of the following pairs of DNA sequences? a) Bacteriophages b) Bacterial cells a) AFLPs and RFLPs b) Two AFLPs c) Plasmids d) All prokaryotic cells c) Two known genes d) All of these 49) PCR proceeds in three distinct steps governed by 59) Which one is a true statement regarding DNA temperature, they are in order of polymerase used in PCR? a) Annealing, Synthesis, Denaturation a) It is isolated from a virus b) Synthesis, Annealing, Denaturation b) It remains active at high temperature c) Denaturation, Annealing, Synthesis c) It is used to ligate introduced DNA in recipient cells d) Denaturation, Synthesis, Annealing d) It serves as a selectable marker 50) Nucleic acid segment tagged with a radioactive molecule 60) For transformation, micro-particles coated with DNA to is called be bombarded with gene gun are made up of a) Plasmid b) Probe a) Silicon or Platinum c) Clone b) Gold or Tungsten d) Vector c) Silver or Platinum d) Platinum or Zinc

Sample Paper

Comprehensive examination M.Sc. Biotechnology Institute of Molecular Biology and Biotechnology (IMBB)

Student Name:		Signature	
Roll No.:		Time: 60 min	Total Marks: 40 。
	Biotechia Mol Blobgy (1 17) Biotechia Cell and Tiesbe Cusu	rie, skh	
Q No:1	What is tissue culture? Discustissue culture?	ss the types of cultures and t	their importance used in
Q No: 2	Define Genetic Engineering. I constructs?	Discuss steps of genetic eng	ineering to design genetic
Q No:3	Write a note on: a) Food preservation	b) Advantages of GN	A foods
Q No:4	Describe the essential biosafe		110005
Q No:5	Define replication? What are replication?	the basic differences in prok	caryotic and eukaryotic
Q No:6	What are the composting orga functioning?	nism? What are the require	ments for their proper

Institute of Bio Technology

Bahauddin Zakariya University Multan

Programme Details

M.Sc. Molecular Biology & Biotechnology (Morning)

Core Courses for Comprehensive Exam Sessions 2014-16

S/No	Course No.	CourseTitle	Cr.Hrs
	0	elongation and resonation. Transcr	ripilon in Eukaryoles. Post-transcessi
emes	iter 01		
emes 1	Biotech-3	Mol.Biology II (F)	
emes 1 2		Mol.Biology II (F) Cell and Tissue Culture. (M)	3

Semester 02

4	Biotech-9	Agriculture Biotechnology (M)	ninchen and on the 3 medians. Anderse
5	Biotech-10	Food Biotechnology (M)	3

Semester 03

6	Biotech-13	Recombinant DNA Technology (M)	3
7	Biotech-15	Metabolomics, Proteomics and Genomics (M)	2

Semester 04

8	Biotech-20	Environment Biotechnology (M)	3
9	Biotech-22	Bio-safety & Bioethics (M)	1



MOLECULAR BIOLOGY-II COURSE OBJECTIVES:

(2 + 1)

To acquaint the student with the chemistry and biology of macromolecules.

COURSE CONTENTS:

Mutations, Types of Mutations, Biochemical basis of Mutagenesis, Base- Analogue Mutagens, chemical Mutagens, Intercalating Agents, Reversion, Transcription, Basic Features of RNA synthesis, Enzymology of RNA synthesis, RNA chain synthesis Initiation, elongation and Termination, Transcription in Eukaryotes, Post-transcriptional Modifications, Translation, Genetic code, codons, Decoding system, Role of mRNA, Role of tRNA, chemical composition of Ribosomes, initiation of protein synthesis, Elongation of polypeptide chain, Termination of polypeptide chain, Difference between protein synthesis in prokaryotes and Eukaryotes, Post-translational Modifications, Regulation of Genes and gene products in prokaryotes, Regulation in Eukaryotes, Protein sorting and transport. DNA repair mechanisms.

Practicals:

Restriction digestion of DNA and preparation of restriction maps, Gel Electrophoresis, PCR, Blotting Techniques, RNA isolation and RT-PCR.

- 1. Molecular Biology by Daved Freifelder. Jones & Barlet Publisher, Boston
- 2. Molecular Biology of the cell by B. Alberts. D. Brag, J. Lewis. M. Raff, K. Roberts and J. D. Watson, Garland Publishers. Jones & Barlet Publisher, Boston



CELL AND TISSUE CULTURE

(2+1)

COURSE OBJECTIVES:

To acquaint the student with the techniques to produce disease free plant material.

COURSE CONTENTS:

Cell and Plant Tissue culture, Introduction, history and importance, Methods of cell and tissue culture, callus culture, organogenesis, somatic embryogenesis, protoplast isolation and fusion, anther and pollen culture Micropropagation, improvement of Plants via Plant cell culture, production of variant plants form selected cells, selection for stress tolerance, production of disease resistant plant material.

Practicals:

Preparation of stock and working solutions, preparation of culture media (liquid) semiliquid and semi-solid) Explants preparation, callus culture and organogenesis. Preparation and fusion of protoplasts.

- 1. Plant Tissue culture manual. K. Lindeseg Kluwer. Academic Publisher, The Netherlands
- 2. Principles of Tissue Engineering, R.P. Lanza, R. Langer and J. Vacantuy. Academic Press, California, USA.

COURSE OBJECTIVES:

To acquaint the student with the importance and basic concepts of biotechnology.

COURSE CONTENTS:

The cell its organelles and their functions, growth requirements, Characteristics and Industrial importance of algae, fungi and bacteria, Glycolytic pathway and enzymes, fermentative ethanol production, High- energy compounds and coenzymes, TCA cycle and its metabolic and industrial importance, Protein structure, synthesis and function, Nucleic acids and microbial strain selection techniques, Cell growth parameters, fermentor assembly and systems, cultivation conditions, sterilization and inoculation procedures, Biomass and Single-Cell Protein production. Aerobic and anaerobic yeast fermentations: products & processes.

RECOMMENDED TEXT BOOKS:

- 1. Principles of Plant Biotechnology an introduction to Genetic Engineering in plants. S. H. Mantel, J. A. Mathews, R. A. Mecee, Blackwell Scientific Publication Oxford, London, Boston.
- 2. Biotechnology in 21st century, Ayyana, C. Mcgraw Hill.
- 3. Shuler, M.L.and F. Kargi. 1992. Bioprocess Engineering, Prentice-Hall, Englewood Cliffs, NJ.
- 4. Bailey, J.E. and D.F. Ollis. 1986. Biochemical Engineering Fundamentals, 2nd ed. McGraw Hill, New York.

Siglechnology. The Genetic Manipulation of Plants, Oxford University

AGRICULTURAL BIOTECHNOLOGY

(2+1)

COURSE OBJECTIVES:

To acquaint the students with the techniques to develop skills to produce Transgenic Crops.

COURSE CONTENTS:

The concepts of Plant Molecular Markers, Historical Back ground of Tissue Culture, Requirements for in-vitro cultures, Role of Phyto-hormones in somatic embryogenesis, Types of Cultures: Tissue culture and regeneration, Cell culture, Haploid Culture, Protoplast culture. Somaclonal variations as breeding tool, Somatic Hybridization, Commercial application and Issues related to tissue culture, Plant Tansformation; Gene Gun Method of Transformation, Agrobacterium- Mediated transformation, Chloroplast Transformation, PEG mediated transformation etc, Field Evaluation and Commercialization, Transgenic crops for Herbicide, Biotic and Abiotic stress resistance, Introduction to Biofertilizers. Biosafety Concerns and Bioethics on GM crops.

Practical:

Selection of ex-plant, Medium Preparation and Callus Induction, Culturing Agrobacterium and Infection to plant callus, Selection of Transformants, Regeneration of Plantlets and acclimatization, Plant DNA extraction and PCR for Trans gene.

- 1. Jitendra Prakash, R.L.M. Pierik, 1993. Plant Biotechnology: Commercial Prospects and Problems, Intercept
- 2. Peter M. Gresshoff, 1992. Plant Biotechnology and Development, CRC Press
- 3. Adrian Slater, Nigel W. Scott, Mark R. Fowler, 2008. Plant Biotechnology: The Genetic Manipulation of Plants, Oxford University Press.
- 4. Sheela Srivastava, Alka Narula, S. S. Bhojwani, Inc NetLibrary, 2004 "Plant
- 5. Biotechnology and Molecular Markers" publishers Springer link
- 6. H.S. Chawla, 2002 "Introduction to Plant Biotechnology" Second Edition, ISBN 978-1-57808-228-5; 562 Pages, Science publishers, USA
- 7. S Harding, 2007. Biotechnology and Genetic Engineering Reviews: V. 24, Nottingham University Press
- 8. Jane K Setlow, 2003. Genetic Engineering: Principles and Methods, Springer
- 9. National Biosafety Guidelines Biosafety rules 2005
- 10. Clarice Swisher, Carol Wekesser, 1996. Genetic Engineering, Lucent Books

FOOD BIOTECHNOLOGY

(2+1)

COURSE OBJECTIVES:

To acquaint the student with the role of microorganisms in food and food industry, and also with the principles of enzymology, and food engineering.

COURSE CONTENTS:

Food composition, proximate analysis. Probiotics. Fermented foods, Food enzymes, colors and additives, Microbial Food spoilage and food born disease, Food preservation methods, Food engineering principles, Modified atmospheric packaging, Food marketing principles. Mathematical Modeling in food technology .Microbial biotechnology of food flavors production, oil and fats, dairy products, meat and cereal foods, Food Safety and quality control.

Practical:

Estimation of moisture, ash, carbohydrates, protein, fat, crude fiber in food, Detection of proteases, amylases in milk. Determination of iodine number of fats. Determination of specific gravity of sugar. Separation of gluten from wheat flour. Detection of E.coli in drinking water. Detection of yeast and mould in dairy and bakery products. Production of cheese.

- 1. Food Biotechnology by Ulf Stahl, Ute E.B. Donalies and Elke Nevogit, 2008.
- 2. Food colors, flavors and additives technology by NIIR, National Institue of industrial research, Dehli, India 2007.
- 3. Biotechnology and food processing by Meenakshi Paul, 2007
- 4. Food Biotechnology, edited by K. Shetty et al., 2nd edition, 2006.
- 5. Fundamentals of food Biotechnology, Dyong H.Lee, 1996Food Chemistry, lab. Manual by Dennis D. Miller, Willey Inter science., 1998
- 6. Food analysis Manual by Javid Aziz Awan. 2000.
- 7. Biotechnology and our Food by Joan Nordquist, Mass Market paper back, 2000

RECOMBINANT DNA TECHNIQUES (2+1) COURSE OBJECTIVES:

To acquaint the students with Basic techniques and tools used in gene manipulation and its practical uses.

COURSE CONTENTS:

Introduction and History of Recombinant DNA technology, Basic techniques, gel electrophoresis, Blotting techniques, restriction endonucleases, restriction mapping, vectors and their types, cloning vectors, transformations, Polymerase Chain reaction, Cloning strategies, Site-directed mutagenesis. Sequencing strategies, Application of recombinant DNA Technology (agriculture, health, industry, environment and basic research).

Practicals:

DNA and plasmid isolation and agarose gel electrophoresis, conjugation, transformation, role of mutagenic agents in mutation, Blotting techniques.

- 1. Robert F.W., 2005. Molecular Biology. McGraw-Hill.
- 2. Dale, J.W. and von Schantz, M. 2002. From Genes to Genomes: Concepts and Applications of DNA Technology. John-Wiley and Son Limited.
- 3. Meyers, R.A., 2006. Genomics and Genetics. John-Wiley and Son Limited.
- 4. Primrose, S.B., and Twyman, R.M. 2006. Gene Manipulation and Genomics 6 edition. Blackwell Publishing.
- 5. Watson, J.M., Caudy, A.A., Meyers, R.A., and Witkowski, J.A., 2007. Recombinant DNA. Gene and genomes. 3rd Edition. W.h. Freeman and Company, New York.

METABOLOMICS, PROTEOMICS AND GENOMICS (2+0)

COURSE OBJECTIVES:

To acquaint the students with structural and functional genomics, proteomics and metabolomics

COURSE CONTENTS:

Structural genomics, Organization and Structure of the Genomes, Genetic Mapping, Transcript Mapping, Structural Variation in the Genomes, Genomics and proteomics, Molecular Biology of Proteins, Posttranslational modifications, Molecular mechanisms of cellular communication/signaling pathways, Protein-Protein Interactions, receptor identification and characterization, Integral Membrane Proteins and Ion Channels, Advance techniques used in proteomics (MS, LCMS/MS, ICAT, iTRAQ). Introduction to Metabolomics, detection, profiling, analysis and engineering. Micrarray and RNA interference.

- 1. "Handbook of comparative genomics: principles and methodology" by Cecilia Saccone, Graziano Pesole (2003). Published by Wiley-Liss,
- 2. Functional genomics by Chris Town (2002). Published by Springer.
- 3. Human Molecular Genetics-3 by T. Strachan, Andrew P. Read Published by Garland Science, 2004
- 4. Genes IX by Benjamin Lewin . Published by Jones and Bartlett Publishers, 2007
- 5 Systems Biology by Mohamed Al-Rubeai (2006), Martin Fussenegger Published by Springer

ENVIRONMENTAL BIOTECHNOLOGY

(2+1)

To acquaint the students with conservation and reclamation of environment through biotechnology.

COURSE CONTENTS:

Introduction to Environmental biotechnology, Fundamentals of Biological Intervention, Genetic manipulation strategies in environmental biotechnology, Pollution indicators, Pollution control strategies, Biology of Waste water and its treatment, Sludge treatment, Contaminated land and bioremediation, Aerobes and Effluents, Phytotechnology (Terrestrial Phyto-systems, Metal Phytoremediation, Rhizofilteration etc) Hyper accumulation, Solid Waste treatments, Concept of integrated Environmental Detoxification Biotransformation, Products of environmental biotechnology. biotechnology,

Biodegradation of environmental pollutants by microorganisms, Bacteriology of Practical: Drinking water, Microscopic studies of water specimens collected from various locations, Field survey of polluted areas, Field study for pollution indicators (Plants, Microorganisms).

- Christon Hurst, Ronald Crawford, Jay Garland, David Lipson, Aaron Mills and Linda Stetzenbach (2007) Manual of Environmental Microbiology. 3rd Ed.
- Christopher F. Faster and D. A John Wase, John Wase (2004) Environmental Biotechnology. John Willey & Sons.
- Derek R. Lovley (2000) Environmental Microb-Metal Interactions. ASM Press. 3.
- Environmental Biotechnology by Bhattacharyya and Rintu Banerjee (Paperback -4. Mar 1, 2007) OUP India.
- Environmental Biotechnology by Bruce E. Rittmann and Perry L. McCarty (Paperback - Jan 1, 2001). McGraw-Hill Publishing Co. 5.
- Environmental Biotechnology Theory and application, 2003. Gareth M Evans and 6.
- Environmental Biotechnology, 2008. T, Srinivas New Age international Publishers. 7.
- Environmental Microbiology: Methods and Protocols (Methods in Biotechnology) by John F. T. Spencer and Alicia L. Ragout de Spencer (Hardcover - Jul 15, 2004)
- Laqrence P. Wackett (2001) Biocatalysis and Biodegradations: Microbial Transformation of Organic Compounds. ASM Press

BIOSAFETY AND BIOETHICS

(1+0)

COURSE OBJECTIVES:

To acquaint students with principles of biosafety and ethical perspectives of Biotechnological systems.

COURSE CONTENTS:

Introduction to Biosafety (Definition, Concept, Uses and abuses of genetic information, Biohazards), Good Laboratory Practices, Risks Related to GMOs, International Rules & regulations for Biosafety & GMOs. Introduction to Bioethics, Ethical issues regarding GMOs, Euthanasia, Issues related to Reproductive & Cloning technologies, Issues to transplants and Eugenics, Patenting, Commercialization and Benefits Sharing, role of National Bioethic committees.

RECOMMENDED TEXT BOOKS:

1. WHO. 2006. Laboratory Biosafety manual third edition. AITBS publishers and distributors, India. (Available online free of cost).

2. Lewis RJ. Sax dangerous properties of Industrial materials, 10nth edition. Toronto, John Wiley and sons, 1999.

3. Lenga RE. 1988. The Sigma-Aldrich Library of chemical safety data, 2nd ed. Milwaukee, WI, Aldrich chemical company.

 Furr AK. 2000. CRC handbook of laboratory safety 5th edition. Boca Raton, FL, CRC press.

5. United states Department Health and Human services. 1999. Biosaftey in Microbiological and biomedical laboratories. 4th edition. Centers for disease control and prevention/National institutes of Health, Washington DC.

6. Biosafety in the Laboratory: Prudent Practices for Handling and Disposal of Infectious Materials Committee on Hazardous Biological Substances in the Laboratory, National Research Council ISBN: 0309-55920-0, (1989) available from the National Academies Press at: http://www.nap.edu/catalog/1197.html

7. Genes Technology and Policy. Jose Maria A. Ochave 2003 http://www.apdip.net/publications/iespprimers/eprimer-genes.pdf

either concentration or production require-

8. Bioethics & Biosafety in Biotechnology by V Sree Krishna.