CHANDGAD - III (C.B.C.S.) (2012 Course): SUMMER - 2017

Misc Sem-III

Subject: Environmental Biotechnology

Day: Wednesday Time: 02.00 PM TO 05.00 PM Date: 05/04/2017 Max Marks: 60 Total Pages: 1 N.B. : 1) Q.No.1 and Q.No.5 are COMPULSORY. Attempt ANY TWO remaining questions from Section-I and Section-II each. 2) 3) Answers to both the sections should be written in the SEPARATE answer books. Figures to the right indicate FULL marks. 4) SECTION-I Q.1 Answer ANY FIVE of the following question in brief: (10)Types of environmental pollution. b) Effect of noise pollution. c) Significance of environmental issues Primary treatment of sewage and waste water. Standards of water Detection of microorganisms in environmental fresh water. Q.2 Answer the following questions: (10)a) Describe the degradation of xenobiotic compound in environment. Role of Bioreactors in waste water treatments. Q.3 Explain the following: (10)Soil pollution sources and its control. a) b) Types and application of biosensors Write short notes on ANY TWO of the following: (10)0.4 Control of air pollution through Biotechnology. b) Water pollution monitoring. Biofilms in treatment of waste water. SECTION-II (10)Q.5 Answer the following: Which are the global environmental issues? b) What is bioremediation? Explain its role in environmental pollution control. (10)0.6 Answer ANY TWO of the following: Role of vermicomposting in solid waste management. b) Describe desalination technique. c) How the Biotechnology is useful in healthy environment. (10)Q.7 Write short notes on the following: a) Carbon credit b) Kinetics of biodegradation of waste Describe the aerobic and anaerobic treatments for waste water. (10)0.8 Describe the importance of genetically modified organisms for improving (10)the environment.

Subject : Plant Biotechnology

Day: Friday Time: 02.00 PM TO 05.00 PM Date: 07/04/2017 Max Marks: 60 Total Pages: 1 N.B: All questions are COMPULSORY. 1) 2) Write section I and II on separate answer sheets. Draw well labeled diagrams WHEREVER necessary. 3) All questions carry EQUAL marks. 4) SECTION-I 0.1 Answer ANY FIVE of the following: (10)What is hotspots of Biodiversity? a) Describe the objectives of modern plant breeding. b) What is pure line selection? c) d) What are the principles of marker assisted breeding. What is bioprospecting? e) Explain genetic basis of cross pollinated crops. f) (10)Answer the following questions. 0.2 Describe in detail the conservation strategies of Biodiversity. a) b) Describe systems and methods of breeding in self pollinated crops. (10)Attempt the following questions: 0.3 Explain the characterization of Biodiversity. Differentiate between breeding in self pollinated and cross pollinated plants. b) (10)Write short notes (ANY TWO): 0.4 Molecular methods for characterization of Biodiversity a) Mass selection b) Backcross breeding methods c) SECTION-II (10)Answer the following questions. Q.5 Describe the technique of double haploid plant production. a) Explain manipulation of gene expression in plants. b) Explain ANY TWO of the following: 0.6 Embryo culture technology. a) Production of transgenic plants resistant to insects. b) Utilization of microorganisms for strawberry. c) (10)Write short notes. Q.7 Differentiate between Ti and Ri plasmid based vectors. a) Seed and micropropagation industries in India. b) (10)Explain diagrammatically. Q.8 Micropropagation of Ginger via axillary shoot proliferation. a) Technique of protoplast isolation and culture. b)

CHANDGAD - III (C.B.C.S.) (2012 Course): SUMMER - 2017

Subject: Animal Tissue Culture

Day: Saturday Time: 02.00 PM TO 05.00 PM Max Marks: 60 Total Pages: 2 Date: 08/04/2017 N.B: Question ONE & FIVE are COMPULSORY. 1) Answer any TWO from questions 2, 3 & 4 and TWO from questions 6, 7 & 8. 2) Both the sections should be written in SEPERATE answer book. 3) Draw well labeled diagrams WHEREVER necessary. 4) SECTION-I (10)Answer ANY FIVE of the following questions in brief: Q.1 Name the instrument used for sterilization of plastic ware and reagents. Give its principle. What is cross contamination? **b**) Name any two cell lines and state their use. What are HEPA filters? For what purpose it is used? d) How tissue culture medium is sterilized? e) Define the role of mitomycin- C in animal tissue culture. (10)Answer the following questions: 0.2What are the common contaminants encountered in tissue culture? How contamination in tissue culture is avoided? b) Why it is important to incubate cell cultures in 5% carbon-dioxide atmosphere? (10)Write notes on ANY TWO of the following: Q.3 Suspension culture a) Balanced salt solution b) Immortal cell lines c) Describe various methods used for disaggregation of cells. Add a note on (10) Q.4

OR

choice of enzymes made for this purpose.

What are advantages and disadvantages of including serum in tissue culture medium?

P.T.O

SECTION-II

| Q.5 | | Answer the following questions: | (10) |
|-----|----------|--|--------------|
| | a) | What is generation number? How and why is it important to keep track of generation number during maintenance of cell lines? | |
| | b) | Describe the methods for viable counting. What do they determine? | |
| | | | |
| Q.6 | a) b) | Define stem cells? State the different types and applications of stem cells. What is MTT assay? For what purpose it is used? | (05) (05) |
| Q.7 | | Write short notes on ANY TWO of the following: | (10) |
| | a) | Microcarriers | |
| | b) | Bioreactor | |
| | c) | Therapeutic proteins | |
| Q.8 | | Define anchorage dependent and anchorage independent cells. Describe any one method for scale up of each type of cells. | (10) |
| | | OP | |

of biotechnology.

Give an account of applications of animal tissue culture in different disciplines

Subject: Human Genetics

Day: Tuesday Time: 02.00 PM TO 05.00 PM Max Marks: 60 Total Pages: 1 Date: 11/04/2017 N.B.: Q.No.1 and Q.No.5 are COMPULSORY. Out of remaining questions attempt 1) **ANY TWO** questions from each section. Answers to both the sections should be written in the SEPARATE answer books. 2) 3) Figures to the right indicate FULL marks. **SECTION-I** 0.1 Answer ANY FIVE of the following in brief: [10] Define autosomal recessive trait. a) b) What is F 2 generation? Why 'O' group is a universal donor? c) d) What is epistasis? State the genetic abnormality in Jacob's syndrome. e) Give an example of genetic disorder caused due to point mutation. 0.2 a) Explain the Mendel's law of segregation giving example. 1061 b) What is Rh factor? Explain its significance in blood transfusion. [04] 0.3 a) What are mitochondrial genetic defects? How they are caused? [05] b) Describe the structure of nucleosome. [05]Describe the probable factors responsible for induction of mutations in the [10] Q.4 genome. What are their consequences? What is aneuploidy? How is it caused? Enlist various structural chromosomal abnormalities giving one example of each. **SECTION - II** Write the principle of fluorescent in situ hybridization (FISH) technique. [05] 0.5 a) Explain its advantages and limitations in detecting genetic abnormalities. b) Describe the genetic abnormality and sympotms of Klinefelter's syndrome. [05] Explain and compare maternal and Mendelian patterns of inheritance giving Q.6 a) suitable examples. b) What is X-Chromosome inactivation? Explain its mechanism. [05] [10] Write short notes on ANY TWO of the following: Q.7 a) Down's syndrome b) Hardy - Weinberg law Amniocentesis Describe the method of karyotype analysis. Explain its significance in [10] Q.8 diagnosis of genetic abnormalities. Describe codominance, incomplete dominance and lethal allele combinations giving suitable examples.