CHANDGAD - III (C.B.C.S.) (2012 Course): WINTER - 2015

Subject: Environmental Biotechnology

Day: Saturday Time: 02.00 PM TO 05.00 PM Date: 10/10/2015 Max Marks: 60 Total Pages: 1. 25583 N. B. : Q. No. 1 and Q. No.5 are COMPULSORY. 1) 2) Attempt ANY TWO remaining questions from Section – I and Section – II each. Answers to both the sections should be written in the SEPARATE answer books. 3) 4) Figures to the RIGHT indicate full marks. SECTION-I Q. 1 Answer ANY FIVE of the following questions in brief: (10)Mention any four methods of water pollution monitoring. b) Differentiate primary and secondary air pollutants. c) What are the effects of noise pollution? d) Write any four sources of soil pollution. e) Enlist various types of reactors used in wastewater treatment. How can we detect the micro organisms in the environment? 0.2 Answer the following questions: (10)a) Describe the overview drinking water standards in relation to public health. b) What are biosensors? Explain their types and applications. 0.3 Explain the following: (10)a) Principles of industrial water treatment. b) Various gases responsible for global warming. Write short notes on ANY TWO of the following: 0.4 (10)a) Vermicomposting b) Bioposticides and their merits Microbes in wastewater treatment SECTION - II Q. 5 Answer the following: (10)What is hazardous waste? Discuss their impact on human health. Discuss the significance of techno-economic feasibility of conversion of waste into energy. Q. 6 Answer ANY TWO of the following: (10)a) Define acid rain. Write about the causes and effects of acid rain. b) What is meant by carbon credit? Discuss various methods of carbon crediting. c) Discuss the merits and demerits of bioremediation. Q. 7 Write short notes on ANY TWO of the following: (10)Ozone depletion b) Bioaugmentation 0.8 Answer the following: (10)a) Define desalination. Mention various techniques used in this process. Add a note on byproducts and industrial applications of desalination. What are the global environmental challenges? Mention any four of them

with respect to their impacts.

Subject: Plant Biotechnology

Day: Monday Time: 02.00 PM TO 05.00 PM Date: 12/10/2015 Max Marks: 60 Total Pages: 1 25584 N.B.: 1) O. No. 1 and O. No. 5 are COMPULSORY. 2) Answer any TWO from Questions 2, 3 and 4 and from 6, 7, & 8. 3) Figures to the right indicate FULL marks. Answers to both the sections should be written in SEPARATE answer book. 4) SECTION-I (10)0.1 Explain any **FIVE** of the following questions in brief. Plant Diversity a) Threatened and extinct species c) Biodiversity hot spots of a India Techniques for the production of hybrid varieties d) Characterization of biodiversity e) Research areas in plant biotechnology Q.2 Answer the following: (10)What are the objectives of modern plant breeding? Briefly explain methods of plant breeding in self pollinated plants. Explain the following: 0.3 (10)Write a note on selection procedure following hybridization. What is marker assisted plant breeding? Discuss its applications. Write short notes on the following 0.4 (10)a) Bioprospecting of plant diversity for product development b) Conservation strategies of plant diversity SECTION-II 0.5 Answer the following questions (10)a) Explain in vitro approaches for plant genetic improvement. b) What is present status of plant genetic Engineering? Q.6 Answer any TWO of the following questions: (10)What are secondary metabolites? Explain their applications. b) Enlist seed industries and plant tissue culture industries in India. c) Describe the applications and advantages of micropropagation. Write short notes: Q.7 (10)Ti and Ri plasmids Molecular markers and their applications 0.8 Answer any TWO of the following: (10)a) Briefly describe Biopestisides and Biofertilizers. b) Give diagrammatic representation of transgenic plant production resistant to

Explain the techniques of micropropagation via somatic embryogenesis.

Subject : Animal Tissue Culture

Day: Wednesday Time: 02.00 PM TO 05.00 PM Max Marks: 60 Total Pages: 1 Date: 14/10/2015 25585 N.B.; 1) Q. No. 1 & Q. No. 5 are COMPULSORY. Out of the remaining attempt any TWO questions from Section - I and any TWO questions from Section - II. 2) Answers to both the sections should be written in **SEPARATE** answer books. 3) Draw well labeled diagrams WHEREVER necessary. SECTION-I 0.1 Answer the following: (10)What are HEPA filters? What is its role? State the role of phenol red in tissue culture medium. b) What is cross contamination? c) What is senescence of a cell line? d) Define density dependent inhibition of mitosis Describe the method of sterilization of: Q.2 a) (05)i) Heat resistant reagents, ii) Heat sensitive reagents What are the common microbial contaminations encountered in tissue (05) b) culture? How are they detected? State the significance of CO₂ incubator. Why animal cell cultures are Q.3 (05)incubated in it? Why tissue culture medium is supplemented with serum? (05)b) 0.4 Write short notes on Any TWO of the following: (10)Suspension cultures Balanced salt solution b) Organ culture c) SECTION-II Attempt any TWO of the following: (10)0.5 Define primary culture. Outline the steps involved in preparation of primary a) culture using enzymatic disaggregation. Compare the characteristics of normal diploid cell line with continuous cell b) Define anchorage dependent cells. Describe any one method of their scale c) up. Describe the method and significance of viable counting (05)Q.6What is microtitration (MTT) assay? For what purpose it is used? (05)b) State true or false Giving reasons 0.7 Animal tissue culture does not find any application in biotechnology industry (05)a) Stem cells can be used to replace diseased cells in the body (05)b) Write short notes on any TWO of the following (10)Q.8 Nunc cell factory a) Fluidized bed reactor b)

Perfused monolayer culture

Subject: Human Genetics

Day: Friday Time: 02.00 PM TO 05.00 PM Max Marks: 60 Total Pages: 1 Date: 16/10/2015 N.B.: 1) Q.No.1 and Q.No.5 are COMPULSORY. Out of remaining questions attempt ANY TWO questions from each section. Answers to both the sections should be written in the **SEPARATE** answer books. 2) Draw neat and labeled diagrams WHEREVER necessary. 3) Figures to the right indicate FULL marks. 4) SECTION - I 0.1 Attempt the following: [10] a) What is a dihybrid cross? b) Define epistasis. c) Define autosomal dominance. d) What is triploidy? e) Define monosomy giving one example. 0.2 Attempt the following: [10] a) Explain Mendel's law of segregation. b) Describe 'ABO blood group' system. Q.3 Attempt the following: [10] Describe the structure and role of Telomere. b) Explain the role of Y chromosome in sex determination. Write short notes on ANY TWO of the following: 0.4 [10] a) Dosage compensation b) Cystic fibrosis c) Chromosomal banding SECTION - II Q.5 Attempt the following: [10] Describe numerical chromosomal abnormalities. b) Explain the cause and clinical symptoms of Klienfelter's syndrome. Attempt the following: 0.6 [10] a) Describe amniocentesis and state its applications. b) Describe mitochondrial genetic defects. Write short notes on ANY TWO of the following: 0.7 [10]a) Down's syndrome b) FISH c) Haemophilia Describe the various types of mutations with illustrations. [10] 0.8

Explain various inborn errors of amino acid metabolism.