

This question paper contains 2 printed pages]

PB—27—2024

FACULTY OF SCIENCE AND TECHNOLOGY

B.Sc. (Third Year) (Fifth Semester) EXAMINATION

APRIL/MAY, 2024

(New Course)

BIOTECHNOLOGY

Paper—DSEBT-4E-I

(Advanced Bioinformatics)

(Saturday, 13-4-2024)

Time : 10.00 a.m. to 1.00 p.m.

Time—3 Hours

Maximum Marks—75

N.B. :— (i) Attempt all questions.

(ii) Figures to the right indicate full marks.

(iii) Illustrate your answers with suitable diagram scheme etc.

1. What is bioinformatics ? Give a detailed account of advanced fields of bioinformatics. 15

Or

Write on :

(a) HTML 8

(b) WWW. 7

P.T.O.

2. Describe in detail multiple sequence alignment. 15

Or

Write notes on :

(a) Rasmol 8

(b) Cn3D. 7

3. Describe in brief Protein sequence database. 15

Or

Write notes on :

(a) GenBank 8

(b) DDBJ. 7

4. Describe secondary structure classification of protein. 15

Or

Write notes on :

(a) Motif 8

(b) Domain. 7

5. Write short notes on (any *three*) : 15

(i) WWW

(ii) PyMol

(iii) Pubmed

(iv) Homology modelling.

This question paper contains 2 printed pages]

PB—24—2024

FACULTY OF SCIENCE

B.Sc. (Third Year) (Fifth Semester) EXAMINATION

APRIL/MAY, 2024

(New Pattern)

BIOTECHNOLOGY

(Paper-CCBT-3E)

(Bioprocess Technology)

(Friday, 12-4-2024)

Time : 10.00 a.m. to 1.00 p.m.

Time—3 Hours

Maximum Marks—75

N.B. :— (i) All questions are compulsory.

(ii) All questions carry equal marks.

(iii) Draw a well labelled diagrams wherever necessary.

1. Define Bioreactor. Explain in detail construction, design and operation of Bioreactor. 15

Or

(a) Explain welding, surface treatment components and specification of the fermenters. 8

(b) Explain in detail materials of construction of fermenter. 7

P.T.O.

2. Define sterilization. Explain in detail media sterilization with principle and mechanism. 15

Or

- (a) Define media. Explain in detail constituents of design of media. 8
- (b) Describe design of sterilization cycle using kinetics of thermal death of microbes. 7
3. Define Growth. Explain in detail batch and continuous culture kinetics. 15

Or

- (a) Explain in detail effect of temperature on cell growth. 8
- (b) Give an account on strategies of fermentation control. 7
4. Explain in detail quality control and quality assurance. 15

Or

- (a) Describe scale up in Bioprocesses fermentation. 8
- (b) Give an account on foam and its control. 7
5. Write short notes on (any *three*) : 3×5=15

- (a) Fermenter
- (b) Decimal reduction
- (c) Fed batch culture
- (d) SOP
- (e) Costing of media.

This question paper contains 2 printed pages]

PB—16—2024

FACULTY OF SCIENCE AND TECHNOLOGY

B.Sc. (Third Year) (Fifth Semester) EXAMINATION

APRIL/MAY, 2024

(New Course)

BIOTECHNOLOGY

(Developmental Biology)

(Monday, 08-04-2024)

Time : 10.00 a.m. to 1.00 p.m.

Time—3 Hours

Maximum Marks—75

Note :— (i) All the questions are compulsory.

(ii) Each question carries equal marks.

1. Describe in detail competence, determination and commitment. Explain each with an example. 15

Or

- (a) Explain in brief about fertilization. 8
(b) Describe in detail organogenesis in frog. 7

2. Describe in detail developmental stages of Drosophila. 15

Or

- (a) What is stem cell ? Describe in detail different types of stem cells. 8
(b) What is progenitor cells ? Explain in detail cell lineages in animal.

P.T.O.

7

3. What is seedling development ? Explain floral patterning in Arabidopsis.

15

Or

(a) Write a note on meristem structure.

8

(b) Describe in detail photomorphogenesis.

7

4. What is transgenic technology ? Explain its application in plant and animal.

15

Or

(a) Write a note on concept of test tube baby.

8

(b) Write a note on citrus.

7

5. Write short notes on (any *three*) :

15

(i) Types of Cleavage

(ii) Apoptosis

(iii) Hybrid

(iv) Embryoculture

(v) Patterns of Cleavage.

This question paper contains 2 printed pages]

PB—28—2024

FACULTY OF SCIENCE

B.Sc. (Third Year) (Fifth Semester) EXAMINATION

APRIL/MAY, 2024

(New Pattern)

BIOTECHNOLOGY

(Medical Biotechnology)

(Saturday, 13-4-2024)

Time : 10.00 a.m. to 1.00 p.m.

Time—3 Hours

Maximum Marks—75

N.B. :— (i) All questions are compulsory.

(ii) Draw neat and well labelled diagrams wherever necessary.

1. What is stem cell ? Describe in detail stem cell therapy with examples.15

Or

(a) Write principle and applications of live, killed and attenuated vaccines. 8

(b) Write principle and applications of recombinant DNA and protein based vaccines. 7

2. How to diagnose bacterial, viral and parasitic diseases by using ELISA and Western blot ? 15

Or

(a) Write about production of monoclonal antibodies. 8

(b) Write applications of monoclonal antibodies in diagnostic reagents.7

P.T.O.

3. Describe in detail role of adult and embryonic stem cells and a note on its clinical applications. 15

Or

- (a) Write about properties and potency of stem cells. 8
- (b) Explain the concept of tissue engineering. 7
4. Describe in detail role of oncogenes and give examples. 15

Or

- (a) What are tumor suppressor genes and how do they work ? 8
- (b) Write about symptoms and treatment of SCID. 7
5. Write short notes on (any *three*) : 15
- (i) Secondary immunodeficiency
- (ii) Apoptosis
- (iii) Plant based vaccines
- (iv) Passive immunization
- (v) Therapeutic interventions of uncontrolled cell growth.

This question paper contains 2 printed pages]

PB—10—2024

FACULTY OF SCIENCE

B.Sc. (Third Year) (Fifth Semester) EXAMINATION

APRIL/MAY, 2024

(New Pattern)

BIOTECHNOLOGY

(r-DNA Technology)

(Friday, 05-04-2024)

Time : 10.00 a.m. to 1.00 p.m.

Time—3 Hours

Maximum Marks—75

Note :— (i) All questions are compulsory.

(ii) Each question carries equal marks.

1. What are reporter genes in gene cloning ? Explain in detail types of reporter genes and add a note on their assays in gene cloning. 15

Or

(a) Explain various vectorless gene transfer methods. 8

(b) Explain pBR 322 as a vector. 7

2. Describe in detail principle and mechanism of PCR and write a note on its types and applications. 15

Or

(a) Explain in detail southern hybridization. 8

(b) Describe in detail DNA Microarray. 7

3. What are probes ? Explain their types and explain screening of library using probe based direct and indirect methods. 15

Or

(a) Explain construction of DNA library. 8

(b) Explain the technique of Autoradiography of DNA. 7

4. What is protein engineering ? Explain various strategies to improve properties of proteins and enzymes. 15

Or

(a) Explain in detail production technology of recombinant insulin. 8

(b) Explain the concept of gene therapy. 7

5. Write short notes on any *three* of the following : 3×5=15

(a) Restriction endonucleases

(b) Sanger's method of DNA sequencing

(c) Chemical synthesis of DNA

(d) BT cotton.