

Seat Number

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April 2015

BT 401

**Pharmaceutical Biotechnology and Bioinformatics  
(Old)**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

**Instructions to Candidates :**

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. Draw the neat labelled diagrams wherever necessary.
5. All question are compulsory.
6. All question carry equal marks.

**1. Explain the following briefly any eight.****16**

- i) Pharmacogenetic.
- ii) Pharmacokinetics.
- iii) Gene chips.
- iv) Clean rooms.
- v) Microbial contaminants.
- vi) Vaccines.
- vii) Antisense therapy.
- viii) Growth hormones.
- ix) FASTA.
- x) Drug designing.

**2. Explain the following any two.****16**

- a) Pharmaceuticals & Microbial and plant origin.
- b) Use of recombinant interferons in Cancer Therapy.
- c) Gene therapy in inherited single gene disorders

3. Explain the following **any two**. 16
- a) Role & impact of genomics in drug discovery.
  - b) Method of simulated docking.
  - c) Gene Bank & DDBJ.
4. Attempt **any two** of the following. 16
- a) Theory of scoring matrices and their use in sequence comparison.
  - b) Protein structure prediction using bioinformatics approach.
  - c) Antigen and antisense therapy.
5. Write notes on **any four**. 16
- a) Structural genomics.
  - b) Randomised control studies.
  - c) Detection of endotoxin.
  - d) Validation studies.
  - e) Statistical analysis of BLAST results.

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**BT-201**  
**Molecular Biology**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

**Instructions to Candidates :**

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2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions are compulsory.
5. All questions carry equal marks.
6. Draw well labelled diagrams wherever necessary.

1. Attempt any eight of following. 16
  - i) ORF
  - ii) Replication fork
  - iii) Point mutation
  - iv) Pyrosequencing
  - v) Initiation factor
  - vi) RNAi
  - vii) Rec A
  - viii) Replio somes
  - ix) Kornberg polymerase
  
2. Explain the causes of mutations. 16

**OR**

Describe the Direct DNA repair system.
  
3. Explain the following any two. 16
  - a) Enzymes in Replication.
  - b) Control of transcription.
  - c) Splicing mechanism.

4. Attempt any two.

16

- a) Post translational modification of protein.
- b) Structure of Ribosome.
- c) Regulation of gene expression in Bacteriophage.

5. Write short notes on any four.

16

- a) DNA polymerase.
- b) Fidelity of replication.
- c) Positive regulation.
- d) RNA Enzyme.
- e) ara operon.

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**BT-402**  
**Bioinformatics**  
**(New)**

**P. Pages : 2****Time : Three Hours****Max. Marks : 80****Instructions to Candidates :**

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All question are compulsory.
5. All question carry equal marks.

**1. Attempt any eight.****16**

- |                  |                        |
|------------------|------------------------|
| i) Genes         | ii) Regulatory regions |
| iii) pH gradient | iv) Protein chips      |
| v) MMDB          | vi) EMBL               |
| vii) PAM         | viii) CN3D             |
| ix) Data mining. |                        |

**2. Explain the following : any two****16**

- a) Protein expression analysis.
- b) Comparative genomics.
- c) Electro Blot.

**3. Attempt any two.****16**

- |                            |                 |
|----------------------------|-----------------|
| a) Structure databases     | b) PAM - Matrix |
| c) Phylogeny tree analysis |                 |

4. Explain the following :

16

- a) Multiple sequence alignment
- b) Modeller

OR

- a) Visualization softwares.
- b) Secondary structure prediction.

5. Write short notes on any four.

16

- a) Chimera
- b) Similarity Search
- c) Pubmed
- d) digital Imaging
- e) Oligonucleotide finger printing.

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**BT - 402**  
**Food Biotechnology**  
**(Old)**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

## Instructions to Candidates :

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions are compulsory.
5. Figures to right indicates full marks.

1. Answers the following **any eight**.

16

- i) Corn sweeteners.
- ii) Cellulose.
- iii) Bakers yeast.
- iv)  $\beta$ -Carotene.
- v) ELISA.
- vi) Patent.
- vii) Fermentation biotechnology.
- viii) SCP.
- ix) Strain improvement.
- x) Beverages industry.

2. Answer the following **any two**.

16

- i) Explain metabolic engineering of bakers yeast.

ii) Why glucan sucrose is importance in synthesis of oligosaccharide.

iii) Surface plasma resonance.

3. a) How cold enzymes play important role in food preparation. 8

OR

Application and production of bacterial cellulose.

b) Genetic engineering of Baker's yeast. 8

OR

What are the approaches of probiotics in fermented food.

4. a) Attempt any two of the following. 10

i) How solid state fermentation is important for fermented food products.

ii) International aspects of quality and safely assessment of food.

iii) Explain application and uses of Dunaliella as food ingredient production.

b) Enzyme technology in dairy products. 6

OR

How you will improve quality of food by using Biotechnology.

5. Write short note on any four. 16

i) Microbial biotechnology of fermented food.

ii) Biotransformation with example.

iii) Prebiotics.

iv) Food born pathogens.

v) Mycotoxin.

vi) Corn Sweeteners.

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**BT-102****Biomolecules and Molecular Enzymology****P. Pages : 2****Time : Three Hours****Max. Marks : 80****Instructions to Candidates :**

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions are compulsory.
5. Draw neat and labeled Diagram(s) wherever necessary.

**1. Answer the following Any Four.****8**

- 1) Anomer.
- 2) Saponification number.
- 3) Antimetabolites.
- 4) Isoenzymes.
- 5) Writing number.
- 6) Feed-back inhibition.

**2. Explain any four.****8**

- 1) Mutarotation.
- 2) Physiologically important glycosides.
- 3) Cyclic fatty acids.
- 4) Xerophthalmia.
- 5) Competitive inhibition.
- 6) Matrix.

3. Attempt any two. 16
- 1) Classification of vitamins.
  - 2) Protein stabilizing bonds.
  - 3)  $\beta$ -oxidation of fatty acid.
4. Attempt any two. 16
- 1) Glycolysis.
  - 2) Protein classification.
  - 3) Structure and functions of coenzyme A.
4. a) Briefly answer Oxidative phosphorylation. 10
- OR
- Mechanism of enzyme action.
- b) Describe secondary structure of protein. 6
- OR
- Significances of  $V_{max}$  and  $k_m$ .
5. Write short notes on any four. 16
- 1) C value paradox.
  - 2) Unisubstrate enzyme kinetics.
  - 3) Enzyme inhibition.
  - 4) Homopolysaccharides.
  - 5) Hemoglobin.
  - 6) Industrial applications of Immobilized enzymes.

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BT-302

**Plant Technology**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

**Instructions to Candidates :**

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions carry equal marks.
5. Draw neat and labelled diagrams whenever necessary.

1. Explain any eight. 16

i) Cybrid	ii) Cytoplasmic male sterility
iii) Gametoclonal variation	iv) parthenogenetic embryo
v) Feeder layer	vi) Molecular marker
vii) Taxol	viii) Chloroplast engineering
ix) Viability of plant cells	x) STS.
2. Describe any two. 16
  - a) Plant tissue culture media.
  - b) Production of virus free plants.
  - c) Haploid identification.
3. Explain any two. 16
  - a) Protoplast culture technology.
  - b) Promoters used in plant vectors.
  - c) Secondary metabolites of commercial importances.

4. Explain any two.

16

- a) Mitochondrial genome.
- b) Selection of fused protoplast.
- c) Herbicide resistances transgenic plants.

5. Write short notes any four.

16

- a) Binary vectors.
- b) RAPD.
- c) Sub protoplast isolation.
- d) Gynogenesis.
- e) Clonal propagation.

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BT-202

**Bioinstrumentation and Biostatistics**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

## Instructions to Candidates :

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions are compulsory.
5. All questions carry equal marks.
6. Draw neat and labelled diagram wherever necessary.

1. Define any eight.

16

- |                                 |                       |
|---------------------------------|-----------------------|
| i) Coefficient of variation     | ii) Fluorescence      |
| iii) Relative centrifugal force | iv) Range             |
| v) Population                   | vi) Sample            |
| vii) Radioactivity              | viii) Resolving power |
| ix) Sedimentation coefficient   | x) Frequency.         |

2. a) Calculate mean, median mode of given data.

12

Value	6	8	10	12	14	16	18	20	22	24
Frequency	20	30	40	40	55	60	55	20	15	25

OR

Calculate standard deviation and coefficient of variation of given data.

Grain yields	15-17	18-20	21-23	24-26	27-29	30-32	33-35
No. of Plants	2	2	4	5	7	9	6
	36-38	39-41	42-44				
	4	3	2				

b) Explain in brief ion exchange chromatography.

4

OR

Describe in brief measures of dispersion.

3. Attempt any two of the following.

16

- i) Describe principle working and applications of SEM.
- ii) Explain principle and applications of cytophotometry and flow cytometry.
- iii) Explain autoradiography and its applications.

4. Describe any two.

16

- i) Write types, principle and applications of density gradient centrifugation.
- ii) Measurement of radioactivity by ionization chamber and proportional counter.
- iii) Affinity chromatography.

5. Write short note on any four.

16

- i) Standard deviation.
- ii) Chi square test.
- iii) ANOVA.
- iv) Types of centrifuge.
- v) Fluorescence microscope.

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BT-403

## Industrial and Business Biotechnology (New)

P. Pages : 2

Time : Three Hours

Max. Marks : 80

### Instructions to Candidates :

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions are compulsory and carry equal marks.
5. Draw neat and labelled diagram wherever necessary.

1. Define / Explain any eight.

16

- |                          |                                  |
|--------------------------|----------------------------------|
| a) Hydrolytic reactions. | b) Glucose isomerase             |
| c) Dextrans              | d) Biosafety                     |
| e) ISO 9000              | f) Polyketide antibiotics        |
| g) Cobalamin             | h) $\beta$ - lactam antibiotics. |
| i) Management concept.   |                                  |

2. Write short notes on any four.

16

- a) Time event - time study.
- b) Microorganisms for citric acid production.
- c) Principles of Management.
- d) Regulation of biosynthesis of streptomycin
- e) PHB.

3. Describe methods of fermentative production & Recovery of following any two. 16
- Tetracycline
  - Erythromycin
  - Lactic acid.
4. Write in detail any two. 16
- Write production and applications of Glucose isomerase.
  - Transformation of prostaglandins.
  - Discuss Biopol - a biodegradable plastic.
5. a) Explain in brief microbial production of cellulases. 8

OR

Write in detail Microorganism involved, production and Recovery of Penicillin - G.

- b) What are polysaccharides? Write in detail production and Recovery of Dextran. 8

OR

Write in detail different methods for industrial amino acid production.

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**BT-403**  
**Industrial Biotechnology**  
**(Old)**

**P. Pages : 2****Time : Three Hours****Max. Marks : 80****Instructions to Candidates :**

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. Draw a neat labelled diagram wherever necessary.
5. All questions are compulsory and carry equal marks.

**1. Attempt any eight of the following.****16**

- a) Xanthan
- b) Erythromycin
- c) Riboflavin
- d) Gluconic acid
- e) Polyhydroxybutyrate (PHB)
- f) Penicillin.
- g) L - lysine
- h) Lactic acid
- i) Dextran
- j) Cephalosporin

**2. Attempt any two of the following.****16**

- a) Describe production, recovery and application of citric acid.

- b) Comment on production and application of vitamin - B12.
- c) Describe importance of microbial polysaccharides.

3. Attempt any two of the following. 16

- a) Write a chemistry, applications and production of xanthan.
- b) Explain biosynthesis and commercial production of Riboflavin.
- c) Write microorganisms used production & recovery of tetracycline production.

4. Attempt any two of the following. 16

- a) Write a note on lactic acid production.
- b) Describe chemistry and properties of polyhydroxy butyrate.
- c) Write an account on production, chemistry and recovery of penicillin.

5. Attempt any four of the following. 16

- a) Alginate
- b) t-tryptophan
- c) Production of alcohol by fermentation.
- d) Recovery and applications of dextran.
- e) Erythromycin.

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**BT-103**  
**Immunology**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

## Instructions to Candidates :

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. Draw a well labelled diagram.
5. All questions are compulsory.
6. All question carry equal marks.

1. Explain the following any eight. 16

i) Hematopoiesis	ii) IgD
iii) Allergy	iv) Immunization
v) Attenuated vaccine	vi) TNF
vii) C <sub>9</sub>	viii) Memory cells
ix) CD <sub>8</sub>	x) Crystal lattice
  
2. Attempt any two of following. 16
  - a) Complex mediated hypersensitivity.
  - b) Structures & functions of cytokines.
  - c) Organ specific autoimmunity.
  
3. Attempt any two of following. 16
  - a) Activation of  $\beta$ - cells.
  - b) Adaptive Immunity.
  - c) Variants of Immunoglobulins.

4. Explain the following.

16

- a) Whole organism vaccines.
- b) Agglutination.

OR

- a) Explain the principle of Immunoelectrophoresis.
- b) Mast cell degranulation.

5. Write short notes on any four.

16

- a) Spleen
- b) BCR - CD19
- c) Selection
- d) HIV genome
- e) ELISA

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BT-203

## Bioprocess Engineering and Technology

P. Pages : 2

Time : Three Hours

Max. Marks : 80

### Instructions to Candidates :

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. All questions are compulsory.
5. All questions carry equal marks.
6. Draw neat and labelled diagram wherever necessary.

1. Answer briefly any eight.

16

- a) Describe feedback inhibition.
- b) Write note on strain improvement for secondary metabolite.
- c) Define Z value.
- d) Define Reynold number.
- e) Write short note on agitator.
- f) Briefly explain process of filtration.
- g) Draw a neat labelled sketch of bioreactor.
- h) Write a note on crystallization.
- i) Explain in short factors affecting sterilization.
- j) Define Bioreactor.

2. a) What is bioreactor? Explain any one bioreactor in detail

12

OR

Explain the continuous sterilization and scale up of the batch sterilization.

b) Write a note on foam separator.

4

OR

Inoculum development.

3. Explain any two of the following.

16

a) Explain strain improvement of industrially important microorganisms.

b) Describe measurement and control of Bio-process.

c) Explain aeration & agitation system for bio-reactor.

4. Explain any two of the following.

16

a) Explain in detail centrifugation technique.

b) Write note on preservation of industrial strain.

c) Write a note on removal of microbial cell and solid matter in down stream processing.

5. Explain any four of the following.

16

a) Exhaust gas.

b) Gas liquid mass transfer.

c) Liquid- liquid extraction.

d) Mutagenesis.

e) Secondary metabolites.

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BT - 404

**Business Biotechnology  
(Old)**

P. Pages : 2

Time : Three Hours

Max. Marks : 80

**Instructions to Candidates :**

1. Do not write anything on question paper except Seat No.
2. Graph or diagram should be drawn with the black ink pen being used for writing paper or black HB pencil.
3. Students should note, no supplement will be provided.
4. Draw neat and labelled diagrams wherever necessary.
5. All questions are compulsory.
6. All questions carry equal marks.

1. Briefly answer the following any **eight**.

16

- a) What are public limited companies ?
- b) Define the 'Bottom-up' approach in nano fabrication.
- c) Mention the need of registering geographical indications.
- d) List the important properties of nanotubes.
- e) What is product planning ?
- f) Give the examples of franchising.
- g) What is 'PLC' ?
- h) List any four agencies supporting entrepreneurship in India.
- i) Give any four examples of bioweapon.
- j) Define the term, nanolithography.

2. Answer the following :

16

- a) Which are the ways to make the organization more effective ?

b) What are the principles of sales management.

OR

What is 'entrepreneurship' and 'self employment' ? Discuss its need in Indian context.

3. Write short notes on any four.

16

- a) SWOT analysis.
- b) Market research.
- c) Nanotubes and Buckyballs
- d) Nanotechnology for drug delivery (method)
- e) Sui Generis system.
- f) patents.

4. Attempt any two of following.

16

- a) Illustrate the working principle and applications of atomic force microscopy.
- b) What are good manufacturing practices ?
- c) Elaborate the applications of nanotechnology in drug delivery.

5. Describe any two of following.

16

- a) ISO 9000 quality system standards.
- b) Guidelines for DNA research activity.
- c) Ethical implications of biotechnological products.

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