# VA-1001-2024

#### FACULTY OF ALL FACULTIES

# All (Third Year) (Fifth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(CBCS/New Pattern)

ENVIRONMENTAL STUDIES (Compulsory)

पर्यावरण अभ्यास (अनिवार्य)

Paper-V

(Wednesday, 27-11-2024)

Time: 10.00 a.m. to 12.00 noon

Time—2 Hours

Maximum Marks—40

- N.B. := (i) Attempt all questions.
  - (ii) All questions carry equal marks.
  - (iii) Draw neat and well labelled diagram wherever necessary.
    - (i) **सर्व** प्रश्न सोडवा.
  - (ii) **सर्व** प्रश्नांना समान गुण आहेत.
  - (iii) आवश्यक तेथे सुबक आकृती काढून नावे द्या.
- 1. Write in detail the effects of modern agriculture.

15

आधुनिक शेतीमुळे होणारे दुष्परिणाम सविस्तर माहिती लिहा.

Or

## (किंवा)

(a) Describe the importance of Environmental Study.

8

पर्यावरण अभ्यासाचे महत्त्व विशद करा.

Describe grassland ecosystem.

7

'गवताळ परिसंस्था' विशद करा.

WT		(2)	VA—1001—2024
2.	Write	biogeographical classification of India.	15
	भारताती	ल सजीवांचे भौगोलिक परिस्थितीनुसार वर्गीकरण करा.	
		Or	
		(किंवा)	
	(a)	Describe alternative energy source.	8
		पर्यायी ऊर्जा स्रोत वर्णन करा.	
	<i>(b)</i>	Discuss the role of an individual in pollution and	abatement. 7
		प्रदूषण व त्याच्या नियंत्रणात मानवाचा वैयक्तिक वाटाः	
3.	Write	short notes any two:	10
	(i)	Desertification	
	(ii)	Food web	
	(iii)	Noise pollution	
	(iv)	Environmental awareness.	
	खालील	पैकी कोणत्याही <b>दोन</b> वर थोडक्यात टिपा लिहा :	
	(i)	वाळवंटीकरण	
	(ii)	अन्न जाळे	
	(iii)	ध्वनी प्रदूषण	
	(iv)	पर्यावरण जागृती.	
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## VB—10—2024

#### FACULTY OF SCIENCE

# B.Sc. (Third Year) (Fifth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(New Course)

#### BIOTECHNOLOGY

(r-DNA Technology)

 (Friday, 29-11-2024)
 Time: 10.00 p.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 labelled diagrams wherever necessary.
- What is gene cloning? Explain in brief various gene cloning strategies used in r-DNA technology.

Or

- (a) Explain various reporter genes used in gene cloning. 8
- (b) Explain construction of M13 vector and add a note on its applications. 7
- 2. Describe in detail the technique of DNA micro array and explain its applications.

15

Or

- (a) Explain in detail Maxam and Gilbert's technique of DNA sequencing. 8
- (b) Explain in detail Northern blotting.

7

WT		( 2 ) VB-	<b>—</b> 10 <b>—</b> 2024		
3.	Describe	in detail the steps involved in construction of c-DNA	library. 15		
		Or			
	(a) E	xplain in detail chemical synthesis of DNA.	8		
	( <i>b</i> ) D	escribe the technique of Autoradiography of DNA.	7		
4.	What is protein Engineering? Explain various strategies to improve properties				
	of protei	ns and enzymes.	15		
		Or			
	(a) E	xplain the concept of Gene therapy.	8		
	( <i>b</i> ) D	escribe in detail production of recombinant insulin.	7		
5.	Write sl	nort notes on any three of the following:	$3 \times 5 = 15$		
	(a) R	estriction enzymes			
	( <i>b</i> ) A	garose gel electrophoresis			
	(c) N	ucleic acid probe			
	( <i>d</i> ) B	t cotton.			

# VB-16-2024

# FACULTY OF SCIENCE

# B.Sc. (Third Year) (Fifth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(New Pattern)

**BIOTECHNOLOGY** 

Paper-CCBT-2E

(Developmental Biology)

(Monday, 2-12-2024) Time: 10.00 a.m. to	1.00 p.m.
Time—3 Hours Maximum M	!arks—75
N.B. := (i) All questions are compulsory.	
(ii) All questions carry equal marks.	
(iii) Draw a well labelled diagram wherever necessary.	
1. Explain types and patterns of cleavage in detail.	15
Or	
(a) Describe developmental stages in chick in detail.	8
(b) Give an account on Gametogenesis and Fertilization.	7
2. Explain in detail concept of stem cells and stem cell technology with ap	plications.
	15

WT-16-2024 OrGive an account on ageing and apoptosis. (*a*) (*b*) Explain in detail abnormal development. 3. 15 Describe in detail development in Arabidopsis. Describe in detail photomorphogenesis. (*b*) Explain meristem structure and its activity. Explain in detail transgenic technology and its applications in plants and 15 animals. Or8 Describe cloning of mammals in detail. (a)(*b*) Describe embryo culture and preservation. 7 Write short notes on (any three):  $3\times5=15$ Commitment (a)(b) Concept of test tube baby (c) **GMOs** Differentiations (d)Competence.

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# VB-27-2024

## FACULTY OF SCIENCE AND TECHNOLOGY

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

## NOVEMBER/DECEMBER, 2024

(New Pattern)

**BIOTECHNOLOGY** 

Paper-DSEBT-4E I

(Advanced Bioinformatics)

(Thursday, 5-12-2024)

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? Describe in detail the applications in bioinformatics. 15

Or

Write notes on:

(a) HTML.

8

(b) URLs.

7

WT	( 2 ) VB-	-272024
2.	Describe in detail the Local alignment and Global alignment.	15
	Or	
	Write notes on:	
	(a) Cn3D.	8
	(b) PyMol.	7
3.	Describe in brief Primary databases.	15
	Or	
	Write notes on:	
	(a) PDB.	8
	(b) PubChem.	7
4.	Describe Protein secondary structure prediction methods.	15
	Or	
	Write notes on:	
	(a) Homology modeling.	8
	(b) Domain.	7
5.	Write short notes on (any three):	15
	(i) Role of internet	
	(ii) Rasmol	
	(iii) Pubmed	
	(iv) Motif.	
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# VB-28-2024

# FACULTY OF SCIENCE & TECHNOLOGY

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

# **NOVEMBER/DECEMBER, 2024**

(New Course))

## BIOTECHNOLOGY

(Medical Biotechnology)

(Thursday, 05-12-2024)	Time: 10.00 a.m. to 1.00 p.m.
Time—3 Hours	Maximum Marks—75
N.B. := (i) All questions are compulsory.	
(iii) Draw well labelled diagram wh	erever necessary.
1. Describe in detail protein based vaccines.	. 15
Or	
(a) Write a brief note on plant based	vaccines. 8
(b) Prepare a draft on stem cell thera	py. 7
2. Explain in detail the production of mono	clonal antibodies. 15
Or	
(a) Give the role of ELISA in the dia	gnosis of bacterial disease. 8
(b) Write a note on western blot technology	nique. 7
	P.T.O.

3.	Define	e stem cell. Explain in detail properties and potency of stem cells	. 15
		Or	
	(a)	Elaborate the concept of tissue engineering.	8
	( <i>b</i> )	Give the clinical applications of embryonic stem cells.	7
4.	What	are oncogenes? Describe in detail the cell cycle with respect to ca	ncer.
			15
		Or	
	(a)	Describe the defects in complement system.	8
	( <i>b</i> )	Write a brief not on secondary immunodeficiency with an exan	nple.
			7
5.	Write	short notes on the following (any three):	15
	(a)	Role of scaffolds	
	( <i>b</i> )	Cell based vaccine	
	(c)	Tumor supressor genes	
	(d)	SCID	
	(e)	Conjugate vaccine	

WT

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## VB-07-2024

## FACULTY OF SCIENCE

# B.Sc. (Third Year) (Sixth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(New Pattern)

#### BIOTECHNOLOGY

Paper-CCBT-2F

(Industrial Biotechnology)

(Thursday, 28-11-2024)

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 a well labelled diagram wherever necessary.
  - (iii) All questions carry equal marks.
- Explain in detail isolation and selection of mutants producing improved level
   of primary metabolites with suitable example.

Or

- (a) Describe in detail isolation of mutants which do not produce feedback inhibitors or repressor?
- (b) Explain isolation of mutants which do not recognize presence of inhibitors or repressors.
- 2. Explain in detail physical and chemical method of cell disruption. 15

WT		( 2 ) VB—07—2	024
		Or	
	(a)	Explain in detail filtration for removal of cell mass.	8
	( <i>b</i> )	Give an account on ultrafiltration and reverse osmosis.	7
3.	Expla	in in detail production of citric acid with their applications. $Or$	15
	(a)	Explain in detail production of penicillin.	8
	( <i>b</i> )	Explain in detail production of protease.	7
4.	Expla	in in detail sterility, pyrogen, toxicity and carcinogenicity testing	. 15
		Or	
	(a)	Describe concept of QC and QA.	8
	(b)	Give an account on fermentation economics.	7
5.	Write	short notes on (any three):	15
	(a)	Modification of permeability	
	( <i>b</i> )	HPLC	
	(c)	Vitamin B <sub>12</sub>	
	(d)	GMP	
	(e)	Solvent recovery.	

## VB-24-2024

#### FACULTY OF SCIENCE

# B.Sc. (Third Year) (Fifth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(New Pattern)

**BIOTECHNOLOGY** 

Paper-CCBT-3E

(Bioprocess Technology)

(Wednesday, 4-12-2024)

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 diagram wherever necessary.
- 1. Define fermenter. Explain in detail construction, design and operation of fermenter.

Or

- (a) Define Bioprocess Engineering. Explain in detail materials of construction of fermenter.
- (b) Explain in detail specification of the fermenter.
- 2. Define media. Explain in detail Design of media and their optimization. 15

P.T.O.

7

Or(*a*) Explain in detail principles, mechanism of capture of particles in air. 8 (b) Give difference between Depth and Screen filters. Explain in detail effect of pH and temperature on cell growth. 3. 15  $(\alpha)$ Explain in detail fed-batch culture kinetics with application. 8 (b) Define Bioproduct. Describe in detail classification of bioproducts. 7 Describe in detail scale up in bioprocess fermentations and factors used in 15 scale up. OrGive an account on standard operating procedures and GMP. 8 (b) Explain computer control fermentations in detail. 7 Write short notes on (any three)  $3\times5=15$ Aeration and agitation (a)(b) Media sterilization Measurement of O<sub>2</sub>/CO<sub>2</sub> (c) **OUR** (*d*) Viscosity and its control.

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WT

## VB-07-2024

#### FACULTY OF SCIENCE

# B.Sc. (Third Year) (Sixth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(New Pattern)

BIOTECHNOLOGY

Paper-CCBT-2F

(Industrial Biotechnology)

(Thursday, 28-11-2024)

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 a well labelled diagram wherever necessary.
  - (iii) All questions carry equal marks.
- 1. Explain in detail isolation and selection of mutants producing improved level of primary metabolites with suitable example.

Or

- (a) Describe in detail isolation of mutants which do not produce feedback inhibitors or repressor?
- (b) Explain isolation of mutants which do not recognize presence of inhibitors or repressors.7
- 2. Explain in detail physical and chemical method of cell disruption. 15

WT		( 2 ) VB—07—2	024
		Or	
	(a)	Explain in detail filtration for removal of cell mass.	8
	( <i>b</i> )	Give an account on ultrafiltration and reverse osmosis.	7
3.	Expla	in in detail production of citric acid with their applications. $Or$	15
	(a)	Explain in detail production of penicillin.	8
	( <i>b</i> )	Explain in detail production of protease.	7
4.	Expla	in in detail sterility, pyrogen, toxicity and carcinogenicity testing	. 15
		Or	
	(a)	Describe concept of QC and QA.	8
	(b)	Give an account on fermentation economics.	7
5.	Write	short notes on (any three):	15
	(a)	Modification of permeability	
	( <i>b</i> )	HPLC	
	(c)	Vitamin B <sub>12</sub>	
	(d)	GMP	
	(e)	Solvent recovery.	

# VB—13—2024

## FACULTY OF SCIENCE

# B.Sc. (Third Year) (Sixth Semester) EXAMINATION NOVEMBER/DECEMBER, 2024

(New Pattern)

## **BIOTECHNOLOGY**

(Environmental Biotechnology)

Time	-3 Hours Maximum Mark	ks—75
N.B.	— (i) All questions are compulsory.	
	(ii) Each question carries equal marks.	
	(iii) Draw a well labelled diagram wherever necessary.	
1.	Describe Anaerobic Biological treatments in detail.	15
	Or	
	(a) Write a note on waste water treatment.	8
	(b) Write a note on packed bed reactors.	7
2.	Describe anaerobic degradation pathways.	15
	Or	
	(a) Write a note on biodegradation of Hydrocarbon.	8
	(b) Write a note on concept of Municipal solid waste managemen	t. 7
	E. The West, Toy,	P.T.O.

VV I		( 2 ) VB—13—20	024
3.	What	is Bioremediation? Describe methods of Bioremediation.	15
		Or	
	(a)	Write a note on Bioremediation of soil.	8
	( <i>b</i> )	Describe phytoremediation with its advantages and disadvantage	s.7
4.	What	is xenobiotics? Describe pesticide degradation with example.	15
		Or	
	(a)	Herbicide degradation pathway.	8
	( <i>b</i> )	Cytochrome P450 system.	7
5.	Write	short notes on (any three):	15
	(a)	Rotating biological contactors	
	(b)	Aerobic degradation pathway	
	(c)	Biodegradation	
	(d)	Phase-II	
	(e)	Important microorganisms in waste water treatment.	

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