Total No. of Questions : 10]	SEAT No. :
PD3198	[Total No. of Pages : 3

[6477]-101 M.Sc. - I BIOCHEMISTRY

CHB - 501 - MJ : Biomolecules (T) (2023 Credit Pattern) (Semester - I)

Time: 3 Hours] [Max. Marks: 70

Instructions to the candidates:

- 1) Q.1 and Q.6 are compulsory and carry 5 marks each.
- 2) Attempt any three questions from question 2 to question 5.
- 3) Attempt any three questions from question No. 7 to question No. 10.
- 4) Answer of the two sections should be written an two separate answer sheets.
- 5) Figures on right hand side indicate marks.

SECTION - I

Q1) Answer the following. [5] a) Define the term Macromolecules with example. Write the Important function of HDL. b) Write various source of vitamin A. c) Write the structure of $\beta(Beta)$ Maltose. d) Define the term weak acid with example. e) Answer the following questions. **Q2**) a) Define Monosacctrides. How they classify? Give examples. i) [3] ii) How vitamins are classified? Elaborate your answer with example.[3] Answer the following question. b) i) Why water is called universal solvent? [2] Write the structure of triglycerides. [2] ii)

Q 3)	Writ	e sho	ort account on the following.	$[5 \times 2 = 10]$
	a)	Stru	cture of water molecule.	
	b)	Gly	cosidic bond.	
	c)	Deo	xy sugars	
	d)	Coe	nzymes	
	e)	Mice	elles	
<i>Q4</i>)	a)	Ans	wer the following.	
		i)	Define the term polysaccharides? With the help of example 1.	mple classify
			polysaccharides.	[3]
		ii)	Why sucrose is called an Invert sugar? Draw the structure	
	b)		wer the following.	
	- /	i)	Write the biochemical function and deficiency of 'Ribe	oflavin' [2]
		ii)	Write short account on 'Rancidity'.	[2]
		11)	write short account on Ranciarty.	[4]
Q 5)	Ans	wer tl	he following any two.	[10]
	a)	Wha	at are fatty acids? How they are classify? Give examples	s. [5]
	b)	Writ	e about osazone formation of glucose with reaction.	[5]
	c)	Wha	at are lipoproteins? How they are classify? Write fund	ction of each
		class	s of lipoprotein.	[5]
			<u>SECTION – II</u>	
Q6)	Ans	wer tl	ne following questions.	[5]
	a)	Defi	ne amino acids.	
	b)	Wha	at are 'Motifs'?	
	c)	Writ	te any two biological significance of protein.	
	d)	Wha	at are nucleosides?	
	e)	Writ	te the structure of phenylalanine.	
Q 7)	a)	Ans	wer the following questions.	
~ .		i)	With the help of diagram explain the primary function of	of protein. [3]
		ii)	Draw the structure of tRNA. Explain its and elaborate	-
		,	of tRNA.	[3]
	b)	Ans	wer the following questions.	
	•	i)	How amino acids are classify on the basis of dietary	need? Give
		,	examples.	[2]
		ii)	Explain the α helix secondary structure of protein.	[2]
[647	7 7]-1	01	2	- -

Q8) Ans	swer the following short notes.	$[5\times2=10]$				
	a)	Biuret Reaction					
	b)	Rare amino acids					
	c)	Nucleotides					
	d)	Acid base properties of amino acid					
	e)	Ninhydrin Reaction of amino acid					
Q9) a)	Answer the following questions.					
		i) Explain Watson and crick model of DNA.	[3]				
		ii) Write about steps involved in protein sequencing.	[3]				
	b)	Answer the following questions.					
		i) Write short account on Ramchandran plot.	[2]				
		ii) What is β pleated sheets? Explain.	[2]				
Q1	<i>0</i>)An	swer the following (any two).	[10]				
	a)	With the help of diagram explain titration curve of Glycine.	[5]				
	b)	Explain structure of Haemoglobin with the help of diagram.	Explain structure of Haemoglobin with the help of diagram. [5]				
	c)	Explain types of DNA using diagram.	[5]				







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[6477]-102 First Year M.Sc. **BIOCHEMISTRY**

CHB-502-MJ: Enzymology and Biophysical Techniques (2023 Credit Pattern) (Semester - I)

Time: 3 Hours [Max. Marks: 70 Instructions to the candidates: Q.1 and Q.6 are compulsory and carry 5 marks. 2) Attempt any three questions from Q.2 to Q.5. 3) Attempt any three questions from 0.7 to 0.10. 4) Answer to the two sections should be written in seperate answer sheets. Figures to the right indicate full marks. **SECTION - I Q1)** Answer any five of the following: [5] What are the predomenant elements that alter the rate of enzymatic reactions? Define the following term. b) i) ii) V_{max}^{m} How is substrate bound to the active site of an enzyme? c) What are affinity lables? d) Write M.M. equation. e) Which enzyme has highest turnover no.? f) **Q2)** Answer the following: What is meant by enzyme inhibition? Discuss in detail competitive a) inhibition. [3] What is the effect of pH and temperature on enzyme activity? [3] ii) How is zymogen activated? [2] b) What is enzyme turnover? Comment on its significance. [2] *Q3*) Write short notes on (Any 5): [10]Binding and active site of enzyme. a) b) Lock and key hypothesis

- c) Isoenzymes
- Presteady state kinetics d)
- Enzyme specificites e)
- Allosteric enzymes f)

04)	Solv	e the	following:	
2 -)	a)	i)		[3]
	,	ii)	Hepatic enzyme gives a V_{max} of 90m mole min ⁻¹ and initial veloc of 30mm min ⁻¹ at (s) of 2.5×10^{-2} M. Assuming enzyme obeys M.	ity
	b)	i)	How proximity and overnation enhances rate of enzyme catalys reaction.	sed [2]
		ii)	What is the relation between enzyme catalyzed reaction and transitistate of reaction?	ion [2]
Q5)	Atte	mpt a	any two of the following:	[0]
	a)	How	enzymes are classified? Elucidate your answer with suitable exampl	es.
	b)	Exp	lain in detail mechanism of action of chymotrypsin enzyme.	
	c)	Disc	cuss the kinetics of two substrate reaction.	
			<u>SECTION - II</u>	
Q6)	Atte	mpt a	any five of the following:	[5]
	a)	Wha	at is cation exchange column?	
	b)	Wha	at are the disadvantages of gel electrophoresis?	
	c)	Whi	ch buffer is used for isoelectric focusing?	
	d)	Wha	at is dialysis in relation to membrane?	
	e)		v do you calculate Rf values of separated amino acids in paromatography?	er
	f)	Wha	at is chromatogrpahy?	
Q7)	Ans	wer tl	ne following:	
	a)	i)	Describe any three factors that have an impact on the biomolecu absorption spectra.	les [3]
		ii)	How will you check the purity of an eznyme?	[3]
	b)	i)	Mention two unique advantages of Gel filtration chromatography	y. [2]
		ii)	Comment on TLC.	[2]

Q8)	Writ	e sho	rt no	otes on (Any 5)	[10]	
	a)	Lyop	hiliz	zation		
	b)	disc	PAC	GE		
	c)	Reve	erse (dialysis		
	d)	Abso	orpti	ion spectrum		
	e)	Aga	rose	gel electrophoresis		
	f)	Mov	ing 1	boundary electrophoresis		
Q9)	Solv	e the	follo	owing:		
	a)	i)		aw the schematic diagram of U.V. visible spectrophotometer blain the instrumentation.	r and [3]	
		ii)		scuss why ammonium sulphate is preferred over other salt cipitating proteins.	s for [3]	
	b)	i)	Giv	ve an exmaple for the following:	[2]	
			1)	A stationary phase used in hydrophobic chromatography	<i>r</i> .	
			2)	A gas use in GLC as mobile phase		
		ii)	Stat	te the principle of GLC.	[2]	
Q10)	Attei	npt a	ny tv	wo of the following:	[10]	
	a)	Expl	Explain the components of HPLC.			
	b)	Disc	uss t	the principle and applications of DNA-Cellulose chromatogra	aphy.	
	c)	Men	tion	the role of the following:		
		i)	Bro	omophenol blue in sample buffer		
		ii)	Am	npholytes in isoelectric focusing.		

- iii) High temperature in GLC
- iv) Photomultiplier tube in U.V. spectrophotometer.
- TEMED in gel electrophoresis. v)



Total No	o. of Que	stions: 5]	SEAT No. :	
PD32	200		[Total No. of Pages :	2
		[6477]	-103	
		First Yea	r M.Sc.	
		BIOCHEN	MISTRY	
		CHB-503-MJ:	Cell Biology	
		(2023 Credit Patter	rn) (Semester - I)	
Time : 2	Hours]		[Max. Marks : 3	35
Instruct	ions to th	he candidates:	_	
1)	_	on 1 is compulsory.	0.5	
2) 3)	_	t any three questions from Q.2 s to the right indicate full mar	_	
,	S	· ·		
Q1) A:		ne following questions (any	5): [5	5]
a)		t is mycellium?		
b)	Expl	ain the function of mitochor	ndria.	
c)	Wha	t are cell adhesion molecule	s?	
d)	Expl	ain the difference between	mitosis and meosis.	
e)	Expl	ain fibronectin structure.		
f)	How	does cyclins and cyclin-de	pendent kinases function?	
Q2) a)	Ansv	wer the following question:	[6	6]
	i)	Explain the structure and fu	unction of xylem and phloem?	
	ii)	Distinguish between interreukaryotic cell.	nal organization in prokaryotic cell an	ıd
b)	Ansv	wer the following question:	[4	4]
	i)	Draw and label structure of	plant cell.	
	ii)	Explain the term sister chro	matid.	

Q3) Write short note (any 5):

- a) Chloroplast
- b) Germ cells
- c) Animal cell
- d) Acrosome reaction
- e) Extracellular matrix
- f) Golgi apparatus

O4)	a)	Answer	the	follow	ing	question	
\mathcal{Q}^{4}	a)	Allow CI	uic	TOHOW	mg	question	٠

[6]

- i) Explain with diagram hemidesmosomes.
- ii) Explain density gradient centrifugation.
- b) Answer the following question:

[4]

- i) How is cell cycle regulated.
- ii) Explain specific cell aggregation in sponges.
- **Q5)** Answer the following question (any 2)

- a) Describe the process of fertilization.
- b) Describe different phases of cell cycle.
- c) Explain with the diagram the process of meosis.



Total No. of Questions : 5]	SEAT No. :
PD3201	[Total No. of Pages : 2

[6477]-104 First Year M.Sc. **BIOCHEMISTRY** CHB-505-MJ: Microbiology

(2023 Credit Pattern) (Semester - I) Time: 2 Hours [Max. Marks: 35 Instructions to the candidates: 1) Q.1 is compulsory. Attempt any three questions from Q.2 to Q.5. 2) Figures to the right indicate full marks. **Q1)** Answer any five of the following: [5] Define enterotoxin and neurotoxin. a) Differentiate between sporocidal and bactericidal. b) Enlist the methods for pure culture preparation. c) Explain pathogenesity of bacteria. d) Define viroids and give an example of disease caused by it. e) f) Define mutant, mutation and mutagen. Attempt the following: [6] **Q2)** a) i) Explain continuous culture of bacteria. ii) Write the principle and applications of phase contrast microscopy. Attempt the following: b) [4] Enlist the groups of chemical antimicrobial agents with suitable i) examples.

> Explain phototrophs and chemotrophs. <u>ii)</u>

<i>Q3</i>)	Write	e a sh	ort note on any five of the following:	[10]
	a)	Aero	obic and anaerobic bacteria.	
	b)	Bact	reriophage	
	c)	Deni	itrification	
	d)	Vaco	eines	
	e)	SEM	1	
	f)	Strea	ak plate technique	
Q4)	a)	Atte	mpt the following:	[6]
		i)	Explain the production of Lysine.	
		ii)	Compare the advantages and disadvantages of various techniof isolation of micro-organisms in pure cultures.	ques
	b)	Atte	mpt the following:	[4]
		i)	Explain the mode of action of tetanus toxin.	
		ii)	Explain isolation and purification of auxotrophic mutants.	
Q5)	Ansv	wer a	ny two of the following:	[10]
	a)	Give	e a detail account on nitrogenase enzyme complex.	
	b)	Expl	lain the mechanism of Lysogeny.	
	c)	Diff	erentiate between prokaryotic and eukaryotic cells.	

PD3202

SEAT No.:		
[Total	No	of Dogge . 2

[6477]-106 First Year M.Sc. **BIOCHEMISTRY**

CHB -508 RM: Research Methodology (2023 Credit Pattern) (Semester - I)

Time: 3 Hours] [Max. Marks: 70]

Instructions to the candidates:

- Attempt all question.
- Figures to the right indicate full marks. *2*)
- **Q1**) Answer the following questions.

 $[5 \times 2 = 10]$

- a) Which is universal solvent? list the other solvents generally used.
- Define trade secretes and give example. b)
- c) What is use of cobalt in Irradiation.
- What is incinerator? write application. d)
- Differentiate between word processing and spreadsheet packages. e)
- **Q2**) Attempt any two of the following.

 $[2 \times 5 = 10]$

- What is use of centrifuge? Write short account or safety of it. a)
- What is PPE? Explain wide range of equipments used in it. b)
- c) What is patent? list rights of patents.

Q3) Attempt the following questions as short notes.

 $[3 \times 5 = 15]$

- a) Use of microwave in disintection.
- b) Importants of patent.
- c) Storage of chemicals.

Q4) Attempt following questions:

 $[2 \times 5 = 10]$

a) Calculate mean of the following using formula.

61, 68, 69, 70, 63, 70, 69, 78.

b) Calculate median of the following using formula.

61, 63, 68, 69, 70, 70, 78

Q5) Answer the following questions.

 $[3 \times 5 = 15]$

- a) Write short account on Basic Rules of Laboratory work.
- b) How will you Import.
 - i) $C \times F$ file
 - ii) A structure into drawing editor.
- c) List the types of Intellectual property with importants.

Q6) Answer the following questions.

 $[2\times5=10]$

- a) Give graphical representation of patent grading procedure.
- b) What is Biomedical waste? Write short account on disposal of Biomedical waste.



Total No. of Questions : 10]	SEAT No. :
PD3203	[Total No. of Pages : 3

[6477]-201 **M.Sc. - I**

BIO-CHEMISTRY

CHB-551-MJ: Bioenergetics and Metabolism

(2023 Credit Pattern) (Semester -II) Time: 3 Hours] [Max. Marks: 70] Instructions to the candidates: Q. number 1 and Q. number 6 are compulsory and carry 5 marks. *2*) Attempt any three questions from Q. number 2 to Q. Number 5. *3*) Attempt any three questions from Q. number 7 to Q. number 10. Use separate answer paper sheets for both sections. *4*) Figure on right hand side indicate marks. 5) **SECTION-I Q1**) Attempt the following questions. [5] Define the term Enthalpy? a) Define the term oxidative phosphorylation? b) List the energy rich compounds in the body. c) Draw struture of ATP. d) e) Write the names of types of fatty acid oxidation. Attempt the following questions. [6+4=10]**Q2**) a) With the help of example explain substrate level phosphorylation. i) [3] ii) Explain carnitine shuttle. [3] Attempt the following questions. b) i) Write the energetics TCA [2] Explain the reaction of conversion of pyruvate to acetyl coenzyme

Α.

[2]

Q 3)	Writ	e sho	ort notes on the following. Any 5.	[10]
	a)	Pay-	-off phase of glycolysis	
	b)	Thre	ee bypass reactions of gluconeogenesis	
	c)	Glyd	oxalate cycle	
	d)	Trig	lyceride synthesis	
	e)	Fate	of pyruvate	
	f)	Reg	ulation of glycolysis	
Q4)	a)	Atte	empt the following questions.	6+4=10]
		i)	Draw a neat, labelled diagram of Electron Transport Chair	(ETC). [3]
		ii)	Explain how many ATP are produce when palmitoyl Co A beta oxidation.	enters in [3]
	b)	Atte	empt the following questions.	
		i)	Write the flow chart of the synthesis of fatty acid?	[2]
		ii)	How is TCA cycle regulated?	[2]
Q5)	a)	Atte	empt the following questions.	6+4=10]
		i)	Explain the entry of glycogen in glycolysis with reactions.	[3]
		ii)	Elaborate the Q cycle in Electron Transport chain.	[3]
	b)	Atte	empt the following questions.	
		i)	What are ketone bodies?	[2]
		ii)	Define the term Inborn error of carbohydrate metabolic example.	ism with [2]
			SECTION-II	
Q6)	Atte	mpt t	he following questions.	[5]
	a)	Wha	at do you mean by salvage pathway?	
	b)	Wha	at is phenylketonuria?	
	c)	Defi	ine the term proteolysis with significance.	
	d)	Give	e the examples of nucleotide coenzymes.	
	e)	Wha	at is Tyrosinemia?	

Q 7)	a)	Atte	empt the following questions.	[6+4=10]
		i)	Explain the Urea cycle with diagram.	[3]
		ii)	Explain the mechanism of Ribonucleotide reductase enzy	yme. [3]
	b)	Atte	empt the following questions.	
		i)	Define the term proteolysis.	[2]
		ii)	Write the conversion Histidine \rightarrow Histamine	[2]
Q8)	Writ	e sho	ort notes on the following. Any 5.	[10]
	a)	Tran	nsamination of amino acids	
	b)	dUN	$MP \rightarrow dTMP$ conversion	
	c)	Sign	nificances of glutathione biosynthesis	
	d)	Con	version of phenyl alanine \rightarrow Tyrosine	
	e)	PRP	PP synthesis from Ribose -5-phosphate	
	f)	IMP	$P \rightarrow GMP$ conversion	
Q9)	a)	Atte	empt the following questions.	[6+4=10]
		i)	Explain the conversion of Serine to Glycine with role of te folate.	etra-hydro-
		ii)	Explain the salvage pathway of purine nucleotide.	[3]
	b)	Atte	empt the following questions.	
		i)	How is Pyrimidine biosynthesis regulated?	[2]
		ii)	What are decarboxylation reactions of amino acids? Give	e example. [2]
Q10)a)	Atte	empt the following questions.	[6+4=10]
		i)	Explain the regulation of urea cycle.	[3]
		ii)	How Isoleucine, leucine enters in degradation pathway?	[3]
	b)	Atte	empt the following questions.	
		i)	Write the following conversation of chorishmate → Peny	l-alanine
				[2]



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PD3204	[Total No. of Pages : 3

[6477]-202 M.Sc. - I

BIOCHEMISTRY

CHB-552-MJ: Membrane Biochemistry and Genetics (2023 Credit Pattern) (Semester - II) Time: 3 Hours] [*Max. Marks* : 70 Instructions to the candidates: Q.1 and Q.6 are compulsory and carry 5 marks each. Attempt any Three questions from Q.2 to Q.5. *2*) 3) Attempt any Three questions from O. No. 7 to O. No. 10. Answers the two sections should be written in separate answer books. *4*) Figures to the right indicate full marks. *5*) **SECTION - I** (Membrane Biochemistry) **Q1**) Answer the following. [5] What is the difference between simple and facilitated diffusion? a) Define Exocytosis b) In what way are membrane proteins asymmetrical? c) Name any two ion translocating antibiotics d) Justify the statement: Phosphatidylcholine and Sphingomyelin are generally e) present outside the leaflet of cell membrane Q2) Attempt the following questions. How does Na⁺K⁺ ATpase pump work? [3] a) How does a gap junction facilitate intercellular communication? [3] b) Discuss facilitated diffusion and its type. [4] c) Q3) Answer the following questions. What happens to the blood cells when suspended in isotonic, hypotonic a) and hypertonic solution? [3] How is calcium transported out of the cell? [4] b) Comment on factors affecting physical properties of membrane. [3] c) P.T.O.

Q 4)	Exp	lain the following.	
	a)	Nuclear Pore	[2]
	b)	Phosphotransferase system	[3]
	c)	Acetylcholine receptor	[2]
	d)	Uniporter, Symporter and Antiporter	[3]
Q 5)	Atte	mpt the following questions.	
	a)	Describe the process of receptor mediated endocytosis with suita example.	ble [5]
	b)	What is the mechanism of voltage gated ion channel? What triggers voltage gated ion channels.	age [5]
		SECTION - II	
		(Genetics)	
Q6)	Ans	wer the following.	[5]
	a)	Describe Mendel's law of dominance.	
	b)	What are Prototrophs? Explain with an example.	
	c)	What are cosmids? Give example.	
	d)	What is complementation tests?	
	e)	What do you mean by colinearity in genetics?	
Q 7)	Ans	wer the following.	
	a)	Explain Hershey and Chase experiment to prove DNA as genetic material	.[3]
	b)	Describe in detail Mendel's law of independent assortment with suita examples for crossing of genes.	ible [3]
	c)	What is fertility factor? Explain the process of conjugation in bacte with a well labelled diagram.	eria [4]

[4]

Q8) Explain the following.

- a) Denaturation of DNA
 b) Concept of gene by Benzer
 c) Plasmid as vector
 [3]
- **Q9**) Attempt the following questions.
 - a) Define genotype and phenotype with examples. [2]
 b) What are conditional mutants? Give examples. [2]
 c) Describe any two genetic disorders of chromosomal origin. [3]
 d) How are mutants isolated and selected? [3]
- Q10) Answer the following.
 - a) Write a short note on human teratogenesis. [5]b) Define transduction and explain in detail the mechanism of transduction
 - with a neat labelled diagram. [5]



Total No	o. of Questions : 5]	SEAT No. :
PD32	205	[Total No. of Pages : 2
	[6477]-203	_
	M.ScI	
	BIOCHEMISTI	RY
CH	B -553-MJ : Techniques in Charact	erization of Biomolecules
	(2023 Credit Pattern) (So	emester -II)
<i>Time</i> : 2	-	[Max. Marks : 35
	ions to the candidates:	
1)	Q.1 is compulsory and carry 5 marks.	
2)	Attempt any three questions from Q.2 to Q.5.	
3)	Answer to both sections should be written on	separate answer sheets.
4)	Figures to the right indicates full marks.	
Q1) A1	nswer any five of the following.	[5]
a)	Why does viscocity change with temper	erature.
b)	What is sedimentation coefficient and i	n what unit it is measured?
c)	Why are Ramchandran plots important	t?
d)	How does light affect fluorescence?	

What is the main use of MALDI TOF in protein studies?

What is X-ray diffraction of biomolecule?

Which radiation is used in autoradiography.

Discuss the principle of IR spectroscopy in the molecular structure

[3]

[3]

[2]

[2]

P.T.O.

Why does radiation occur?

elucidation.

Explain the principle of CD.

e)

f)

a)

b)

Q2) Answer the following.

i)

ii)

i)

ii)

Q3)	Writ	e sho	ort notes on. (Any five)	[10]		
	a)	Intri	nsic viscosity			
	b)	Fluc	prescence			
	c)	Ultra	a centrifugation			
	d)	l) Autoradiography				
	e)	X-ra	y Diffraction			
	f)	MA	LDI-TOF-MS			
0 4)	Solv	e the	following.			
(7)	a)	i)	Write the factors affecting viscosity.	[3]		
	u)	ii)	Explain the difference between diffusion and equilibrium			
	b)	i)	What is zonal centrifugation	[2]		
		ii)	What is the use of chromophose in spectroscopy?	[2]		
Q 5)	Atte	mpt a	any two of the following.	[10]		
	a)	Writ	te the application of preparative centrifugation.			
	b)	Writ	te a note on NMR.			
	c)	Exp	lain the ionization methods used in LCMS and GCM	S.		

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Total No. of Questions: 5	[[
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SEAT No. :	
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PD3206

[Total No. of Pages: 2

[6477]-204 M.Sc. - I

BIO-CHEMISTRY

CHB-555/560-MJ: Bioinformatics (2023 Credit Pattern) (Semester - II)

Time: 2 Hours] [Max. Marks: 35

Instructions to the candidates:

- 1) Q.1 is compulsory.
- 2) Attempt any 3 questions from Q.2 to Q.5.
- 3) Figures to the right indicate full marks.
- Q1) Answer the following questions (Any 5)

[5]

- a) What is Scopus, and how does it help researchers?
- b) What is h-index?
- c) What is UniRef, and why is it important?
- d) What is phylogenetic analysis used for?
- e) Explain protein data bank.
- f) How does NCBI help in science?
- **Q2**) a) Answer the following questions.

[6]

- i) What does phylogenetic analysis help scientists understand?
- ii) How can you evaluate the quality of a scientific journal using impact factors?
- b) Answer the following questions.

[4]

- i) What kind of tools and databases are available through the EBI?
- ii) What is the role of PubMed, and Google Scholar in conducting a literature search?

Q3) Write a Short note (Any 5)

[10]

- a) CLUSTALW
- b) GenPept
- c) dbEST
- d) BLAST
- e) Multiple sequence alignment
- f) Local alignment

Q4) a) Answer the following questions.

[6]

- i) What kind of information can be found in GenBank?
- ii) Explain how pairwise alignment works and why it is useful in comparing sequences.
- b) Answer the following questions.

[4]

- i) What is the main use of GenBank, and who uses it?
- ii) How does UniProtKB help in understanding proteins?

Q5) Answer the following questions (Any Two).

- a) How do BLAST and CLUSTALW help in sequence analysis?
- b) How do you conduct a scientific literature search using PubMed, Scopus, and Google Scholar?
- c) How do GenPept and UniProtKB assist in protein sequence analysis?







Total N	No. of	Questions	:	7]
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SEAT No.: PD3207 [Total No. of Pages : 2

> [6477]-301 S.Y. M.Sc.

BIOCHEMISTRY CHB - 601 - MJ : Molecular Biology

(2023 Credit Pattern) (Semester - III)

Time: 3 Hours] IMax. Marks: 70

Instructions to the candidates:

- All questions are compulsory.
- Figures to the right indicate full marks. *2*)
- **Q1**) Answer in short (any 5):

[10]

- What are transposable elements? a)
- What is translation? Give the role of mRNA in translation. b)
- What is nonsense codon? Give its functions. c)
- d) What is the role of origin of replication locus in DNA replication.
- What is pyrimidine dimer? e)
- Why is RNA splicing important? f)
- Give the significance of signal sequence in protein targetting. g)
- Answer any 2 of the following. **Q2**) a)

[6]

- Describe in detail initiation of translation in prokaryotes. i)
- ii) How is protein targetted across ER membrane?
- Describe in detail mechanism of bacterial defense against viruses. iii)
- Describe the process of transcription in prokaryotes. b)

[4]

Q 3)	a)	Ans	wer any 2 of the following.	[6]
		i)	Write a short note on the enzyme RNA polymerase.	
		ii)	Write a short note on Apoptosis.	
		iii)	Explain in detail epigenetic modification.	
	b)	Desc	cribe in detail the mechanism of base excision Repair.	[4]
Q4)	a)	Ans	wer any 2 of the following.	[6]
		i)	Describe the consequences of defective DNA repair system and disease they cause.	nd the
		ii)	Describe the phenomenon of self splicing and its significance	· •
		iii)	What are transcription factors, and how do they regulate expression?	gene
	b)	Expl	lain the mechanism of homologous genetic recombination.	[4]
Q5)	a)	Ans	wer any 2 of the following.	[6]
		i)	Differentiate between DNA polymerase I, II and III.	
		ii)	How is protein synthesis terminated in Bacteria?	
		iii)	Explain in detail pest translational modification.	
	b)	Desc	cribe ubiquitin dependent pathway for protein degradation.	[4]
Q6)	a)	Ans	wer any 2 of the following.	[6]
		i)	Describe any 3 inhibitors of transcription.	
		ii)	Explain alternative splicing.	
		iii)	How does DNA replication in Eukaryotes differ from prokary	otes?
	b)	With	n diagram explain the discontineous synthesis of okazaki fragmer	ıts.[4]
Q 7)	Ans	wer a	ny 4 of the following.	[10]
	a)	Writ	te a short note on 3 polytailing	
	b)	Defi	ne DNA strands at the replication fork	
	c)	Writ	te the properties of group I intron.	
	d)	Wha	at is the role of ligase enzyme?	
	e)	Give	e the significance of chromatin remodeling in transcription.	

Total No. of Questions : 10]	SEAT No. :
PD3208	[Total No. of Pages : 3

[6477]-302 M.Sc. - II

BIOCHEMISTRY

CHB-602-MJ: Medical Biochemistry and Immunology (2023 Credit Pattern) (Semester - III)

Time: 3 Hours] [Max. Marks: 70 Instructions to the candidates: Q.1 and Q.6 are compulsory. 2) Solve any three questions from Q.2 to Q.5. 3) Solve any three questions from Q.7 to Q.10. 4) Use separate answer sheets for both sections. Figures to the right indicate full marks. **SECTION - I** (Medical Biochemistry) **Q1)** Answer the following: [5] What is 'autophagy'? a) List the enzymes studied in coronary heart diseases. b) Define the term apoptosis. c) Define the term cancer. d) What kind of drugs are macrolides? e) **Q2)** a) Answer the following questions: How 'azoles' works as antifungal drug. [3] How protooncogens are converted to oncogenes? [3] Attempt the following questions. b) i) Define the term 'Angina'? [2] List the names of enzymes found in lysosomes. [2] Q3) Attempt the following short notes. [10]Phagocytosis a) Cancer causing agents b) Hallucinogens and their effects c) Lysosomal storage diseases (LSD). d) Mode of action of sulfonamides/sulfa drugs. e)

Q 4)	a)	Answer the following questions:	
		i) Discuss with the help of diagram mode of action of antibiotics	that
		inhibits SOS Ribosomal submit action.	[3]
		ii) Write the short account on mode of action of analgesics.	[3]
	b)	Attempt the following questions:	
		i) List types of Tualassemia.	[2]
		ii) What is Necrosis.	[2]
Q5)	Atte	mpt the following questions: (Any two)	[10]
	a)	Discuss extrinsic pathway of apoptosis?	[5]
	b)	How microorganisms develop resistance to tetracyclins?	[5]
	c)	Write the mode of action of penicillin with diagram and explanation.	[5]
		SECTION - II	
		(Immunology)	
Q6)	Ansv	wer the following questions:	[5]
	a)	What are interferons?	
	b)	Write the principle of 'RIA' technique.	
	c)	What do you mean by 'opsonization'.	
	d)	Define the term 'Humoral Immune Response'.	
	e)	What are blood group antigens?	
Q 7)	a)	Answer the following questions:	
		i) With the help of diagram explain structure of class MHC-II.	[3]
		ii) What are vaccines? Explain various types of it.	[3]
	b)	Answer the following questions.	
		i) Draw and labelled the diagram of antibody.	[2]
		ii) How 'monoclonal antibodies' are produced in the laboratory?	[2]
Q8)	Writ	e short notes on the following:	[10]
	a)	Bone marrow	
	b)	Type I hypersensitivity	
	c)	Applications of immuno electrophoresis.	
	d)	Types of immuno diffusion.	
	e)	Haematopoiesis.	

Q9) a)	Answer	the follow	ing question	s:
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i)	Explain classical pathway of complement system.	[3]

ii) Discuss antigen presentation & processing by 'cytosolic pathway'.

[3]

[4]

- b) Answer the following questions:
 - i) What is active and passive immunity?
 - ii) Write principles of 'Western Blot' technique.

Q10)Answer the following questions:

- a) With the help of diagram explain cell mediated immune response. [5]
 - [5]
- b) What is ELISA? Explain various types using diagrams.



Total No. of Questions : 5]		SEAT No. :
PD3209		[Total No. of Pages : 2
	[6477]-303	

S.Y. M.Sc. **BIOCHEMISTRY**

CHB-603-MJ: Biochemistry of Specialized Tissue (2023 Credit Pattern) (Semester - III)

Time: 2 Hours] [Max. Marks: 35 Instructions to the candidates: 1) Q.1 is compulsory and carries 5 marks. 2) Attempt any three questions from Q.2 to Q.5. Figures to the right indicate full marks. 3) **Q1)** Answer the following: (Any 5) [5] How do bacterial flagella move? Which organelle provides ATP energy for muscle contractions? b) How does sarin affect the cholinergic synapse? c) What role do the ossicles play in the transmission of sound? d) e) Explain how taste perception is initiated at the molecular level. What are mechanoreceptors of the skin f) **Q2)** Answer the following: What is the difference between thick and thin filaments? i) [3] a) Describe the structure of the skin's epidermis and its relevance to touch sensation. [3] What are the energy sources for muscle contraction? b) i) [2] Define chemotaxis and explain its importance in cell motility. <u>ii)</u> [2] Q3) Write short notes on (any five): [10] Types of cones in the retina? a) b) Outer segment of photoreceptor cells. Stereocilia c) Auditory transduction d) Sensation of itch e) G-proteins in olfactory signaling. f)

Q4) Solve the following:

- a) i) How does the organization of myofibrils contribute to muscle contraction? [3]
 - ii) How do potassium ions move through potassium channels in axons?
 - [0]
- b) i) Describe the molecular structure of actin. [2]
 - ii) How does acetylcholine transmit signal from nerve to muscle? [2]

Q5) Attempt any two of the following:

- a) Discuss contractile proteins in eukaryotic cells other than muscle filaments, highlighting their roles in cell motility and structure.
- b) How does the visual cycle regenerate rhodopsin?
- c) Discuss the creation and propagation of a nerve impulse, detailing the phases of depolarization and repolarization.



Total No. of Questions : 5]	SEAT No. :
PD3210	[Total No. of Pages : 2

[6477]-304 S.Y.M.Sc. BIOCHEMISTRY

CHB - 610 MJ: Toxicology (2023 Credit Pattern) (Semester - III) Time: 2 Hours] [*Max. Marks* : 35 Instructions to the candidates: *1*) Q.1 is compulsory. Solve any three questions from Q.No. 2 to Q.No.5. *2*) *3*) Figures to the right indicate full marks. **Q1**) Answer the following questions. [5] Define the term Toxicology. Define Antidote. b) c) What are hallucinogens. Define the term biotransformation. d) List the various areas of toxicology study. **Q2**) a) Answer the following questions. **[6]** What is Dose Response? List the types of it. i) What measures to be taken during poisoining treatment. ii) Answer the following. [4] b) Define the term tolerance & give its types. i) ii) Write few applications of forensic toxicology. **Q3**) Write short note on the following. $[5 \times 2 = 10]$ Selective Toxicity. a) b) Nacrotics. c) Systemic toxicity Cardiac poisons d) Classification of antidote. e)

Q4) a) Answer the following question	Q4) a)	Answer the follow	ving questions
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[6]

- i) Write short account on phase I reactions of Biotransformation.
- ii) What are asphyxiant poisons?
- b) Answer the following questions.

[4]

- i) What are inorganic poisons?
- ii) List the sites of exposure of toxic substances.

Q5) Attempt the following questions.

- a) With the help of example explain phase II Biotransformation reactions.
- b) What are miscellaneous poisons? Give their types with examples.



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[5]
[6]
idney, [3]
[3]

- b) Answer the following question. **[4]**
 - How carbohydrates are digested? i)
 - Define the term 'Metabolic audosis'. ii)
- *Q3*) Write short notes on the following.

 $[5 \times 2 = 10]$

- a) Blood Brain Barrier (Barrier)
- Fibrinolysis b)
- Transport of O₂ (Oxygen) c)
- Transport of CO₂ (Carbon dioxide) d)
- **Bohrs Effect** e)

<i>Q4</i>)	a)	Ans	wer the following question.	[6]
		i)	Explain Intrinsic pathway of blood cogulation.	[3]
		ii)	How proteins are digested and absorbed?	[3]
	b)	Ans	wer the following question.	[4]
		i)	What is Respiratory audosis?	[2]
		ii)	Write about any four functions of kidney.	[2]
Q 5)	Ans	wer tl	ne following questions.	[10]
	a)		te about platelet aggregation and Extrinsic pathway llation.	of blood [5]
	h)	Defi	ne the term ultamins. Write about its absorption	[5]

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Total No. of Questions: 5]	SEAT No. :
PD3212	[Total No. of Pages : 2
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[6477]-306 S.Y. M.Sc. **BIOCHEMISTRY**

	CHB-612-MJ : Applied Plant Biochemistry (2023 Credit Pattern) (Semester - III)					
		Q.1 is Attem	the candidates: compulsory and carry 5 marks. pt any three questions from Q.2 to Q.5. es to the right indicate full marks.	[Max. Marks: 35		
Q1)	An	swer t	he following: (Any 5)	[5]		
	a)	Giv	e role of Mn in plants.			
	b)	Def	ine photophosphorylation.			
	c)	Wha	at is ligin?			
d) Give any two applications of rubber?						
e) What are macromolecules in plant?						
	f)	Nan	ne the alkaloid obtained from papaver somniferum.			
Q2)	An	swer t	he following:			
	a)	i)	What is Green house technology?	[3]		
		ii)	Write a note on terpenoids?	[3]		
	b)	i)	Give role of ethylene in plant growth.	[2]		
		ii)	Draw the structure of chloroplast.	[2]		
Q3)	Wri	ite sho	rt notes on (any five):	[10]		
	a)	Gib	berellins			
	b)	Pho	tosystems			
	c)	Part	henocarpy			
	d)	Acc	essory pigments in photosynthesis			
	e)	Leg	hemoglobin			
	f) Photorespiration					

Q4) Solve the following:

- a) i) What are advantages of Phytoremediation? [3]
 ii) Discuss hormones in senescence. [3]
 b) i) Give names of any two plants used in phytoremediation. [2]
 - ii) Give deficiency symptoms of Mg in plants. [2]
- **Q5)** Attempt any two of the following:

- a) Discuss Hydroponics as a advanced farming technology.
- b) Describe calvin cycle.
- c) Explain biological and abiological Nitrogen fixation.



Total No. of	f Questions	:	6]
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SEAT No.:	

[Total No. of Pages: 2

PD3213

[6477]-401 M.Sc. -II BIOCHEMISTRY

CHB-651-MJ: Genetic Engineering (2023 Credit Pattern) (Semester - IV)

Time: 3 Hours [Max. Marks: 70

Instructions to the candidates:

- 1) All questions are compulsory.
- 2) Figures to the right indicates full marks.
- **Q1**) Attempt the following.

 $[5 \times 2 = 10]$

- a) Write the applications of genetic engineering.
- b) Define transformation and transfection.
- c) Draw and explain the structure of pUC18.
- d) Write the role of restriction endonuclease in genetic engineering.
- e) What are the applications of RT-PCR?
- **Q2**) Attempt the following questions (any 2).

 $[2 \times 5 = 10]$

- a) Explain blue plaques and clear plaques formation while selection of recombinants.
- b) What are the advantages of DNA vaccine over conventional vaccines?
- c) Explain principle and procedure of southern blotting.
- **Q3**) Write short note on the following (any 3).

 $[3 \times 5 = 15]$

- a) RFLP.
- b) Recombinant vaccine.
- c) Ti plasmid.
- d) Plaque hybridization.

P.T.O.

Q4) Answer the following:

[10]

- Why are sticky ends important in genetic engineering experiments? What are the methods of putting sticky ends to blunt ended DNA molecules?
- b) Explain yeast vectors.

Q5) Answer the following:

 $[3 \times 5 = 15]$

- What are the steps to prepare edible vaccine? What are the applications a) of edible vaccine?
- Explain the principle and applications of hot-start PCT. b)
- What do you understand by artificial chromosome? Explain structure of c) yeast artificial chromosome with its advantages.

Q6) Answer the following:

- Plant viruses exhibit natural tendency of gene transfer to plants and thus have the potential for use as gene transfer and expression vectors. Explain the statement. Give a brief account on DNA and RNA viruses, which are used as expression vectors for plant transformation.
- Describe the principle and procedure of RNA labeling by in vitro b) transcription.







Total No. of Questions: 10]	SEAT No. :
PD3214	[Total No. of Pages : 3

[6477]-402 M.Sc. - II

BIOCHEMISTRY CHB-652-MJ: Fermentation Technology and Tissue Culture (2023 Credit Pattern) (Semester - IV) Time: 3 Hours 1 [*Max. Marks* : 70 Instructions to the candidates: Q1 and Q6 are compulsory and carry 5 marks each. *2*) Attempt any Three questions from Q.2 to Q.5. Attempt any Three (3) questions from Q. No. 7 to Q. No. 10. *3*) Answer the two sections should be written on separate answer sheets. *4*) 5) Figure on right hand side indicate marks. **SECTION - I** (Fermentation Technology) **Q1**) Answer any five of the following. [5] Define COD a) How dilution rate is useful in continuous culture? b) What is meant by enrichment culture? c) What is meant by antifoams? d) Define feedback repression? e) What is media formulation? f) [6] **Q2**) a) Attempt the followings: What is the importance of antifoaming agents in fermentation? i) What is meant by inoculum development? Discuss with example Attempt the following: b) [4] How inhibitors will be useful in fermentation process? i) ii) Which are the nitrogen sources used for fermentation? [6] **Q3**) a) Attempt the followings: What are the things that are filter sterilized? i) Discuss the importance of aeration for the fermentation ii) Attempt the following: [4] b) i) What are the different types of impellers used for agitation?

What is the fate of pyruvate under anaerobic condition?

ii)

P.T.O.

Q 4)	a)	Atte	empt the followings:	[6]
		i)	What is the effect of time of sterilization on the yield of the prod	luct?
		ii)	What are the different types of sensors used during fermentation	on?
	b) Attempt the following:			
		i)	What do you understand from the term fed batch culture?	
		ii)	How will you isolate auxotrophic mutant?	
Q5)	Ans	wer a	any two of the following:	[10]
	a)	Disc	cuss with suitable diagram the construction of a fermenter	
	b)	Exp	lain the steps involved in downstream processing?	
	c)	Wha	at are the different methods used for effluent treatment?	
			SECTION - II	
			(Tissue Culture)	
Q6)	Atte	mpt t	the following.	[5]
	a)	Wha	at is the difference between soft and compact callus tissues?	
	b)	Defi	ine the terms: embryoid and cybrid.	
	c)	Wha	at are the advantages of natural media of animal tissue culture?	
	d)	Wri	te the importance of suspension culture.	
	e)	Defi	ine contact inhibition	
Q 7)	a)	Wri	te short note on the following.	[6]
		i)	Monolayer culture.	
		ii)	Embryo culture.	
	b)	Atte	empt the following.	[4]
		i)	Differentiate between primary cell line and established cell line.	
		ii)	Explain the importance of root culture.	
		•	- · ·	

08)	Write	short no	te on i	the f	follow	ing (์ anv	5) .
$\mathbf{v}_{\mathbf{v}}$	* * 1110		to on	uic i		1115	uii y	J	,.

[10]

- a) Cryopreservation
- b) Applications of tissue culture
- c) Hybridoma technology
- d) Conditioning of tissue cultured plants
- e) Organ culture.
- f) Arrangement of tissue culture laboratory.

Q9) a) Answer the following.

[6]

- i) Describe principle and procedure of preparation of aseptic explant for plant tissue culturing.
- ii) What are the applications of cytotoxic assay.
- b) Attempt the following.

[4]

- i) Describe media preparation and sterilization technique in tissue culture.
- ii) Explain the tests to check viability of cultured cells.

Q10) Attempt any two of the following.

- a) Describe principle, procedure and applications of protoplast culture.
- b) What do you mean by contamination and cross contamination? How to identify it? What care must be taken to avoid it?
- c) Explain the role of following in tissue culture laboratory: Laminar Air flow hood, CO₂ incubator, Culture vessels, Autoclave, Refrigerator



Tota	l No	of Questions: 51
Total No. of Questions: 5] SEAT No.:		
PD	321	[Total No. of Pages : 2
		S.Y.M.Sc.
		BIOCHEMISTRY
		CHB-660-MJ: Endocrinology
		(2023 Credit Pattern) (Semester-IV)
Time	e: 2 H	ours] [Max. Marks : 35
Instr	uctio	s to the candidates:
	<i>1)</i>	Question 1 is compulsory and carry 5 marks.
	<i>2)</i>	Solve any three questions from Q.2 to Q.5.
	<i>3)</i>	Figures to the right indicate full marks.
Q1)	Ans	wer the following questions. [5]
	a)	List the function of cortisol. [1]
	b)	What is Thyroglobulin? [1]
	c)	Define the term Target cell. [1]
	d)	List the major difference between type I diabetes mellitus and type II
	,	diabetes mellitus. [2]
Q2)	a)	Attempt the following questions. [6]
		i) Elaborate a role of IP ₃ and DAf as a secondary messenger with diagram. [3]
		ii) How hormone secreted by thyrotrops from anterior pitutary gland

- b) Answer the following questions.
 - i) What are receptors? How are they classified? [2]

regulates, controls the secretion of harmone from thyroid gland.[3]

- ii) Elaborate the classification of harmone with example. [2]
- Q3) Write short account on the following.

 $[5 \times 2 = 10]$

[4]

- a) Parathyroid gland
- b) Calcium calmodulin system
- c) Insulin and mode of action
- d) Catecholamines
- e) Hypothalamus

Q4) a)	Answer the	following	questions.
~ /		\mathcal{C}	1

[6]

- i) Explain the mechanism of action of hormone, secreted by posterior pitutary gland. [3]
- ii) With the help of diagram explain the synthesis of steroid harmone. [3]
- b) Answer the following questions.

[4]

[2]

- i) Write the physiological role of growth hormones.
- ii) Write in details about any two gastrointestinal hormone secretion and mode of action. [2]

Q5) Answer the following questions.

 $[2 \times 5 = 10]$

- a) Why cyclic-amp is termed as secondary messenger system? Elaborate your answer with diagram and mechanism.
- b) What is target cell insensitivity? Explain with the help of examples.



Total	l No.	of Questions : 5] SEAT No .
		SEAT NO.
PD	321	6 [Total No. of Pages : 2
		M.Sc II
		BIOCHEMISTRY
		CHB-661-MJ: Food Technology
		(2023 Credit Pattern) (Semester-IV)
Time	: 2 H	[Max. Marks : 3:
Instr	uction	ns to the candidates:
	1)	Question 1 is compulsory and carry 5 marks.
	2)	Solve any three questions from Q.2 to Q.5.
	3)	Figures to the right indicate full marks.
Q1)	Ans	wer the following questions. [5]
	a)	What is Spiruline?
	b)	List the purpose of food analysis in food technology
	c)	List the types of Genetically modified food (GMO).
	d)	Which parameters are included in proximate composition analysis o food.
	e)	Define the term food Thickner with example.
Q2)	a)	Answer the following questions. [6]
		Define the terms simple cell markets. I let a decorate a februarie and

Define the term single cell protein. List advantages of single cell 1) protein. [3]

> Explain the effect of germination as the carbohydrate contents of ii) food. [3]

Answer the following questions. b) [4]

Define the term food additives. i) [2]

ii) List anticaking agents used in food Technology. [2]

Q 3)	Write	e sho	rt notes on the following.	$[5\times2=10]$
	a)	Natu	ral sweetners	
	b)	Food	d preservatives	
	c)	Flav	or enhancer	
	d)	Bene	efits of meat Tenderization	
	e)	Enzy	ymes used in fruit juice processing	
Q4)	a)	Ansv	wer the following questions.	[6]
		i)	Discuss the main purpose of food Irradiation.	[3]
		ii)	Write about the process of starch production.	[3]
	b)	Ansv	wer the following questions.	[4]
		i)	Define the term food spoilage.	[2]
		ii)	What are meat tenderizer enzymes? Give examples.	[2]
Q5)	a)	Ansv	wer the following questions.	[6]
		i)	Give source and function of natural colours used in food to	echnology.
		ii)	Discuss the potential advantages of genetically modified	food. [3]
	b)	Ansv	wer the following questions.	[4]
		i)	List the factors responsible to spoil the food.	[2]



Give examples of unconventional source of protein.

[2]

ii)