Printed Pa		Subject Code:- ABT0403 Roll. No:
	NOIDA INSTITUTE OF ENGINEERING AI (An Autonomous Institute Affi B.Tec	liated to AKTU, Lucknow)
	SEM: IV - THEORY EXAM	
	Subject: Structural and C	,
Time: 3	Hours	Max. Marks: 100
 The que Section Section Section 	A - Question No- 1 is 1 mark each & Question B - Question No-3 is based on external choice C - Questions No. 4-8 are within unit choice questions be left blank. Any written material after the control of the control o	carrying 6 marks each. uestions carrying 10 marks each. er a blank sheet will not be evaluated/checked.
	SECTION A	A 20
1. Attempt	t all parts:-	
1-a.	The process by which DNA makes a copy of	itself during cell division is called as? (CO1)
	(a) Transcription	
	(b) Translation	
	(c) Reverse Transcription	
4.1	(d) Replication	
1-b.	Which is called as start codon? (CO1)	1
	(a) AUG	
	(b) AAC (c) AUU	
	(d) CAU	
1-c.	During centrifugation, separation of analyte is	s based on(CO2)
1 0.	(a) Size	(002)
	(b) Shape	
	(c) Density	
	(d) All	
1-d.		detailed and comprehensive description of the tween all proteins whose structure is known.
	(a) SCOP	
	(b) CATH	
	(c) Both (a) & (b)	
	(d) Niether (a) nor (b)	
1-e.	(CO3)	sy to ionize atoms and disrupt molecular bonds?
	(a) Visible radiation	
	(b) Infrared radiation	
	(c) UV radiation (d) X-rays	
1-f.	•	-nucleic acid interactions is an application of 1
1-1.	(CO3)	-nucleic acid interactions is an application of
	(a) XFEL(b) Circular Dichroism	
	(c) chronia Dichiolom	

	(c) EPR	
	(d) X-ray crystallography	
1-g.	Aptamers are(CO4)	1
	(a) short, ssDNA or ssRNA	
	(b) long, ssDNA or ssRNA	
	(c) short, dsDNA or dsRNA (d) long, dsDNA or dsRNA	
1-h.	Nucleoside contains(CO4)	1
1-11.	(a) Nitrogenous Base	1
	(b) Sugar	
	(c) Phosphate group	
	(d) Only (a) & (b)	
1-i.	The study of genome starts with the analysis of(CO5)	1
	(a) Nucleosides	
	(b) Nucleotides	
	(c) Chromosomes	
	(d) All	
1-j.	Turner syndrome is characterized by (CO5)	1
	(a) X0	
	(b) XX	
	(c) XY	
	(d) XXY	
	apt all parts:-	
2.a.	Explain the major difference between glycoproteins and proteoglycans? (CO1)	2
2.b.	What is difference between soluble and membrane proteins? (CO2)	2
2.c.	What is the full form of FRET and why we use this? (CO3)	2
2.d.	What is SELEX? (CO4)	2
2.e.	Which compound is used for visualization of DNA bands in Gel doc? (CO5)	2
	SECTION B 30	
3. Answ	er any <u>five</u> of the following:-	
3-a.	Discuss the role of chaperons in protein folding? (CO1)	6
3-b.	Explain the G-protein coupled receptors (GPCRs) protein families. (CO1)	6
3-c.	What is NMR? Describe the method to prepare sample for NMR Spectroscopy? (CO2)	6
3-d.	Explain the significance of Ramachandran Plot in 3D structure validation of generated protein model. (CO2)	6
3.e.	What are the various applications of circular dichroism (CD) in structural biology? (CO3)	6
3.f.	Write down various types of RNA and also explain their function? (CO4)	6
3.g.	Write notes on (a) Surface Plasmon Resonance (b) Fluorescence quenching (c) Fluorescence Resonance Energy Transfer. (CO5)	6
	SECTION C 50	
4. Answ	er any <u>one</u> of the following:-	
4-a.	Explain the importance of α -helix, β -pleated sheet and turn in protein structure. (CO1)	10
4-b.	What do you understand by sequence alignment? What type of approaches you will follow to align the sequences? Explain with a sequence alignment problem. (CO1)	10
5. Answ	er any one of the following:-	
5-a.	Explain SCOP and CATH? (CO2)	10

5-b.	Why we perform SDS-PAGE and Western Blot technique. Explain. (CO2)	10
6. Answer	any one of the following:-	
6-a.	Write notes on (a) Circular Dichroism (b) Electron Paramagnetic Resonance spectroscopy (c) Single molecule fluorescence. (CO3)	10
6-b.	Explain Electron Paramagnetic Resonance (EPR) spectroscopy. How EPR works? Explain the different ways by which EPR shows similarity with NMR. (CO3)	10
7. Answer	any one of the following:-	
7-a.	Why aptamers are widely used in nanotechnology? Discuss. (CO4)	10
7-b.	What do you understand by isomer, epimer and anomer? Explain with suitable examples. (CO4)	10
8. Answer	any one of the following:-	
8-a.	Discuss the dynamics of Protein-RNA complexes. (CO5)	10
8-b.	Describe the solution methods and enzymatic approaches to measure the kinetics of RNA-protein interactions. (CO5)	10