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Enrolment No:			
<div>UPES</div> <div>End Semester Examination, May 2025</div> <div><div>Course: Biophysics</div><div>Program: B.Tech – Biotechnology &amp; B.Tech - Biomedical</div><div>Course Code: PHYS2030</div></div> <div><div>Semester: IV<sup>th</sup></div><div>Duration: 3 hours</div><div>Max. Marks: 100</div></div>			
Instructions: Carefully read and attempt all the questions.			
S. No.	Section A  Short answer questions/ MCQ/T&F (20Qx1.5M= 30 Marks)	Marks	COs
Q1.	<b>Recall the postulate of quantum theory.</b> A. Energy is continuous B. Electrons follow circular orbits only C. Energy is quantized and exchanged in discrete units called quanta D. Atoms do not emit radiation	1.5	CO1
Q2.	<b>Hydrogen bonding occurs when hydrogen is bonded to:</b> A. Carbon, sulfur, or silicon B. Oxygen, nitrogen, or fluorine C. Phosphorus, arsenic, or bromine D. Any metal ion	1.5	CO2
Q3.	<b>Electrostatic interactions are strongest when:</b> A. The dielectric constant is high B. The charges are far apart C. The dielectric constant is low and charges are close D. In aqueous environments	1.5	CO2
Q4.	<b>One of the unusual properties of water is:</b> A. Low surface tension B. Maximum density at 0°C C. High heat capacity and heat of vaporization D. It behaves as a non-polar solvent	1.5	CO1
Q5.	<b>In an enzyme-catalyzed reaction, the rate increases with substrate concentration until:</b> A. The enzyme is saturated B. All substrate is converted to product C. Temperature reaches 0°C D. The reaction becomes zero-order	1.5	CO2
Q6.	<b>The enthalpy change (ΔH) of a reaction reflects:</b> A. The amount of disorder in the system B. The heat absorbed or released C. Activation energy D. The reaction rate	1.5	CO2
Q7.	<b>Recall the most abundant lipids in biological membranes.</b> A. Sphingomyelin B. Phosphatidylcholine C. Cholesterol D. Cardiolipin	1.5	CO1
Q8.	<b>Identify a secondary active transporter.</b> A. Na <sup>+</sup> /K <sup>+</sup> -ATPase B. Aquaporin C. Glucose-Na <sup>+</sup> symporter D. Voltage-gated K <sup>+</sup> channel	1.5	CO1

<b>Q9.</b>	<b>During an action potential, depolarization is caused by:</b> A. Cl <sup>-</sup> influx B. K <sup>+</sup> efflux C. Na <sup>+</sup> influx D. Ca <sup>2+</sup> efflux	<b>1.5</b>	<b>CO2</b>
<b>Q10.</b>	<b>The patch clamp technique is used to measure:</b> A. Entire neuron activity B. Single ion channel currents C. Metabolic rate D. Membrane elasticity	<b>1.5</b>	<b>CO2</b>
<b>Q11.</b>	<b>Recall the compounds used in Anfinsen's renaturation experiments?</b> A. SDS and ATP B. Urea and β-mercaptoethanol C. Heat and ethanol D. Acetone and NaCl	<b>1.5</b>	<b>CO1</b>
<b>Q12.</b>	<b>Levinthal's paradox suggests:</b> A. Protein folding is a random and slow process B. Proteins cannot fold spontaneously C. Proteins fold via specific pathways, not by random search D. All proteins fold the same way	<b>1.5</b>	<b>CO1</b>
<b>Q13.</b>	<b>The energy landscape theory of protein folding illustrates:</b> A. Folding requires enzymes B. Proteins fold through a linear pathway C. Proteins fold via multiple paths down an energy funnel D. All folding happens in one step	<b>1.5</b>	<b>CO2</b>
<b>Q14.</b>	<b>A folding funnel represents:</b> A. Increase in entropy B. Progressive decrease in free energy toward native state C. Increase in protein size D. Increase in temperature	<b>1.5</b>	<b>CO1</b>
<b>Q15.</b>	<b>Molecular chaperones assist protein folding by:</b> A. Directly folding proteins B. Preventing aggregation and providing folding environment C. Breaking misfolded structures D. Hydrolyzing misfolded proteins	<b>1.5</b>	<b>CO2</b>
<b>Q16.</b>	<b>The delay between absorption and fluorescence emission is typically:</b> A. Microseconds B. Nanoseconds C. Seconds D. Hours	<b>1.5</b>	<b>CO2</b>
<b>Q17.</b>	<b>UV absorbance at 280 nm is primarily due to:</b> A. Peptide bonds B. Aromatic amino acids C. Aliphatic residues D. Sulfur-containing residues	<b>1.5</b>	<b>CO1</b>
<b>Q18.</b>	<b>The melting temperature (T<sub>m</sub>) of a protein is defined as:</b> A. The temperature at which all proteins degrade B. The temperature at which 50% of the protein is unfolded C. The temperature at which proteins are phosphorylated D. The point of maximum absorbance	<b>1.5</b>	<b>CO1</b>
<b>Q19.</b>	<b>In ITC, the binding isotherm provides information about:</b> A. Molecular geometry B. Stoichiometry, enthalpy, and binding constant C. Molecular diffusion D. pH dependence only	<b>1.5</b>	<b>CO1</b>
<b>Q20.</b>	<b>..... is not a limitation of ITC?</b> A. High sample consumption B. No need for labeling C. Low throughput D. Sensitivity to heat artifacts	<b>1.5</b>	<b>CO2</b>

<b>Section B</b> <b>(4Qx5M=20 Marks)</b>			
<b>Q1.</b>	Explain the concept of secondary active transport, classify its types, and provide an example for each type.	<b>5</b>	<b>CO3</b>
<b>Q2.</b>	Discuss the role and significance of hydrogen bonding and van der Waals interactions in biological systems.	<b>5</b>	<b>CO2</b>
<b>Q3.</b>	Describe the patch clamp technique and summarize its importance in studying the properties of biological membranes.	<b>5</b>	<b>CO2</b>
<b>Q4.</b>	Describe binding energy and analyze its contribution to enzyme catalysis.	<b>5</b>	<b>CO3</b>
<b>Section C</b> <b>(2Qx15M=30 Marks)</b>			
<b>Q1.</b>	Design an experiment to evaluate the suitability of two different buffers in maintaining protein stability. (7) Describe the experimental method and its detailed steps. (8)	<b>15</b>	<b>CO4</b>
<b>Q2.</b>	Explain the principle behind isothermal titration calorimetry (ITC). (5) Describe the working mechanism of ITC with the help of a labeled diagram. (10)	<b>15</b>	<b>CO4</b>
<b>Section D</b> <b>(2Qx10M=20 Marks)</b>			
<b>Q1.</b>	Identify the conditions necessary for conformational changes in proteins. (3) Explain how diphosphoglycerate (2,3-BPG) influences the binding affinity of oxygen to hemoglobin. (7)	<b>10</b>	<b>CO3</b>
<b>Q2.</b>	Analyze how protein folding processes comply with the second law of thermodynamics. (4) Illustrate and explain the energy-entropy funnel model with a labeled diagram. (6)	<b>10</b>	<b>CO4</b>