# End Semester Examinations April 2022

Programme: M.Sc. Biochemistry

**Session: 2021-22** 

Semester: I

First

Max. Time: 3 Hours

Course Title: Analytical Biochemistry

Max. Marks: 70

Course Code: SIAS BC 1103 C 3104

### **Instructions:**

1. Question no. 1 has seven parts and students need to answer any four. Each part carries three and half Marks.

2. Question no. 2 to 5 have three parts and student need to answer any two parts of each question. Each part carries seven marks.

Q 1. (4X3.5=14)

- a) Describe the principle of compound microscope? How do you calculate linear and angular magnification?
- b) On what basis does agarose gel electrophoresis separate molecules? Which factors affect the migration of DNA in gel electrophoresis?
- c) Explain the principle of centrifuge. What factors have an influence on centrifugation?
- d) What is the principle of high performance liquid chromatography and its application? Could you tell some disadvantages of high performance liquid chromatography?
- e) Draw Jablonski diagram to illustrate fluorescence. Make sure that you include the name of the various processes involved. What do you understand by "Stokes shift" in fluorescence?
- f) What are the basic principles of NMR spectroscopy? Where is NMR used? What causes chemical shifts in NMR?
- g) What kind of information do we get from flow cytometry? What is the principle behind this technique? What are the three main components of a flow cytometer?

Q 2. (2X7=14)

- a) What are the basic components of "Fluorescence microscope"? Describe in brief its optical system. What is the principle behind confocal microscopy?
- b) What are the two different types of Electron Microscope? How do they differ? How do you prepare samples for them?
- c) What are the basic principles of Atomic-force microscope (AFM)? Elaborate the methods used in AFM to acquire images.

- a) How does polyacrylamide gel electrophoresis (PAGE) work? Elaborate step by step. What is the difference between PAGE and SDS-PAGE? What is the role of SDS in SDS-PAGE?
- b) What is the difference between gel electrophoresis and pulsed field gel electrophoresis? When and why pulse field gel electrophoresis is preferred?
- c) What is density gradient centrifugation? How does it differ from differential centrifugation? Describe in brief the two types of density gradient centrifugation.

Q 4. (2X7=14)

- a) What is the principle of affinity chromatography? How does antibody affinity chromatography work? What are the main differences between ion exchange chromatography and gel filtration chromatography?
- b) What is gas chromatography and how does it work? Which is better between high performance liquid chromatography and gas chromatography? Give reasons to support your answer.
- c) What are the three major types of radiation detectors? What is the advantage of using a scintillation counter rather than a Geiger-Muller counter to detect radiation? How do you handle radioactive sources safely?

Q 5. (2X7=14)

- a) Which spectroscopy will you use to understand secondary structure of proteins? What are the basic principles of this spectroscopy? How do you analyse the data to predict the secondary structure?
- b) How is Beer-Lambert law used in absorption spectroscopy? How do you determine extinction coefficient?
  - A protein has one tryptophan and two tyrosine in its sequence. Assume molar extinction coefficients at 280 nm of tryptophan and tyrosine as 3000 and 1500 M<sup>-1</sup> cm<sup>-1</sup>, respectively. What would be the molar concentration of that protein if its absorption at 280 nm is 0.60 in a 1 cm cuvette?
- c) What is electrospray ionisation mass spectrometry? Does electrospray ionization cause fragmentation? Why is ionisation important in mass spectrometry? Discuss about the mass analysis in this technique.

# **End Semester Examinations April 2022**

Program: M.Sc. Biochemistry

Session: 2021-22

Semester: First

Max. Time: 3 Hours

**Course Title: Fundamentals of Biochemistry** 

Max. Marks: 70

Course Code: SIAS BC 1 1 01 GEC 4004

#### Instructions:

- 1. Question no. 1 has seven parts and students need to answer any four. Each part carries three and half Marks.
- 2. Question no. 2 to 5 have three parts and student need to answer any two parts of each question. Each part carries seven marks.

Q 1.

(4X3.5=14)

- a) Describe mutarotation phenomenon.
- b) Write a short note on prostaglandins and their functions.
- c) Why monosaccharides are called reducing sugars? Explain with examples.
- d) What are hexose derivatives?
- e) How many categories of amino acids are there? Explain with examples.
- f) Write down name and structure of different kinds of nitrogenous bases found in nucleic acids.
- g) What are bile salts? Mention their functions.

Q 2.

(2X7=14)

- a) Describe with examples the different heteropolysaccharides.
- b) Write down convention of writing name of disaccharide/oligosachharide in biochemistry.
- c) What are glycoconjugates? Give their structural details.

Q3.

(2X7=14)

- a) Draw the structure of amino acids with aromatic side chains.
- b) What are motifs and domains? Describe the forces that control the tertiary structure of proteins.
- c) Describe classification of proteins.

Q 4.

(2X7=14)

- a) Describe the Chargaff's rule, also explain the characteristics of different forms of DNA.
- b) What do you mean by denaturation and renaturation of DNA?
- c) Describe structural organization of eukaryotic DNA.

Q 5.

(2X7=14)

- a) Describe different types of sphingolipids with their structures.
- b) Describe properties and functions of cholesterol and its derivatives.
- c) What are lipids? Write down their classification.

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## **End Semester Examinations April 2022**

Programme: M.Sc Biochemistry

Session: 2021-22

Semester:

First

Max. Time: 3 Hours

Course Title:

**Cell and Molecular Biology** 

Max. Marks: 70

Course Code:

SIAS BC 11 01 C 4004

#### Instructions:

- 1. Question no. 1 has seven parts and students need to answer any four. Each part carries three and half Marks.
- 2. Question no. 2 to 5 have three parts and student need to answer any two parts of each question. Each part carries seven marks.

Q 1. (4X3.5=14)

- a) Draw the structure of a plant and an animal cell mentioning their important organelles
- b) Although the membrane is tightly packed by phospholipids, it is not static. Justify
- c) What are the unique properties of plasma membrane? Draw the structure of phospholipids and lipid bilayer
- d) In fluid mosaic mosaic model, how are the phospholipids and proteins arranged in the membranes of the cells? Explain the model with an illustration.
- e) State three essential functions of the genetic material.
- f) Schematically show the overview of cellular processes from DNA to proteins.
- g) A preliminary RNA sequence has the following nucleotides:

5'-AUGAAAGUUCGCAUUACUAACUACAGUGGUAG-3'. What would have been the template and non-template of genomic DNA sequence of this pre-mRNA?

Q 2. (2X7=14)

- a) What are the constituents of nuclear pore complex? Explain with an illustration.
- b) Explain the molecular trafficking that allows proteins to move to and fro through nuclear pore.
- c) Provide indirect evidences to show that genetic information is stored in DNA. Explain in detail atleast one experimental proof to show DNA mediates transformation with clear illustration.

Q3. (2X7=14)

- a) What is the structure of DNA? Explain with clear structures! What is the form in which the genetic information is stored in it?
- b) What are the alternate forms of DNA double helix and show how they are different from each other? Explain the functional state of E.coli chromosome.

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c) What are the five types of RNA molecules and mention their essential roles. Compare the prokaryotic and eukaryotic gene expression with illustration.

Q 4. (2X7=14)

- a) Explain the chemical composition of eukaryotic chromosomes, Explain in detail three levels
  of DNA packaging in eukaryotes.
- b) Telomeric regions are important for DNA stability. Justify by stating three important functions of telomeres. Explain the replication of telomeric region.
- c) A segment of human DNA has the following sequence of nucleotide pairs:
  - 3'-ATATTTACGTGCTACCTTGCTGATAGGACTGCAATATCTACTTTACCATT-5'
  - 5'-TATAAATGCACGATGGAACGACTATCCTGACGTTATAGATGAAATGGTAA-3' When this is segment of DNA is
  - (a) Replicated what will be the sequence of nucleotides in the daughter DNA strands
  - (b)Transcribed by RNA polymerase, identify the promoter sequence and write the sequence of nucleotides in the RNA transcript?
  - (c) Explain clearly how primosome and replicosome complexes co-operate to replicate two strands of DNA

Q 5. (2X7=14)

- a) How transcription is regulated in prokaryotes?
- b) Compare the DNA polymerases of prokaryotes and Eukaryotes with clear illustrations. How are RNA polymerases different from DNA polymerases of prokaryotes?
- c) Explain how all three steps of transcription process in Prokaryotes with clear illustration.

## **End Semester Examinations April 2022**

**Programme: M.Sc Biochemistry** 

**Session: 2021-22** 

Semester:

First

Max. Time: 3 Hours

Course Title: Protein Biochemistry

Max. Marks: 70

Course Code: SIAS BC 11 04 C 3104

#### Instructions:

- 1. Question no. 1 has seven parts and students need to answer any four. Each part carries 3.5 Marks.
- 2. Question no. 2 to 5 have three parts and student need to answer any two parts of each question. Each part carries 7 marks.

Q 1. (4X3.5=14)

- a) What are hydrophobic and hydrophilic amino acids? Give list of amino acids of these classes.
- b) What is dipole moment? Write in detail about the dipole moment of water.
- c) Explain the energy coupling in biological reactions.
- d) What are good and bad helix formers and why?
- e) Describe Anfinsen's experiment to demonstrate protein refolding.
- f) Explain the varieties of lipid motions which create disorder in the fluid lipid bilayer.
- g) What are the major classes of membrane proteins? How do they interact with membrane lipids?

Q 2. (2X7=14)

- a) Explain Ramachandran plot with torsion angles related to alpha helix and beta sheets.
- b) Explain in detail the non-covalent interactions between molecules in the protein.
- c) Explain the physicochemical properties of the intracellular environment in detail.

Q3. (2X7=14)

- a) Explain the secondary structure of proteins.
- b) Write in detail the protein folding patterns of the motifs and domains.
- c) What are the forces and interactions responsible for the tertiary and quaternary structure of proteins?

Q4. (2X7=14)

- a) Write in detail the chaperone and chaperonin assisted protein folding.
- b) Explain the thermodynamics of protein folding.
- c) What is protein melting curve and how can we determine the fraction of unfolded protein using these curves?

Q5. (2X7=14)

- a) Explain the mechanism of the development of amyloidoses in detail.
- b) How does Mass spectrometry work and explain the methods of ionization?
- c) Write about different methods of protein engineering with examples.

## End Semester Examinations April 2022

Programme: M.Sc. Biochemistry Session: 2021-22

Semester: First Max. Time: 3 Hours

Course Title: Biomolecules and Metabolism Max. Marks: 70

Course Code: SIAS BC 1102 C 4004

## **Instructions:**

1. Question no. 1 has seven parts and students need to answer any four. Each part carries three and half Marks.

2. Question no. 2 to 5 have three parts and student need to answer any two parts of each question. Each part carries seven marks.

Q 1. (4X3.5=14)

- a) Differentiate between B DNA and Z DNA.
- b) What are coupled reactions?
- c) Differentiate between amylose and amylopectin
- d) Explain importance of omega fatty acids in the diet.
- e) What are sphingolipids?
- f) Comment on structural aspects of a glycoprotein.
- g) Differentiate between de novo and salvage pathways of nucleic acids.

Q 2. (2X7=14)

- a) Differentiate between gluconeogenesis and glycolysis.
- b) Describe energy transducing steps of citric acid cycle.
- c) Describe the structure, properties and importance of cellulose and chitin.

Q3. (2X7=14)

- a) Explain oxidation of odd number of fatty acids.
- b) Discuss synthesis of fatty acids at Fatty acid synthase complex.
- c) Briefly classify membrane phospholipids and cite their structural features.

Q 4. (2X7=14)

- a) Discuss tertiary structure of proteins and forces responsible for maintaining it.
- b) Explain features of alpha helix and beta pleated sheet structures.
- c) Give schematic representation of urea cycle steps and mention the enzymes involved at various steps.

Q 5. (2X7=14)

- a) Give structure of nucleotides.
- b) Briefly explain purine nucleotide biosynthesis pathway.
- c) Explain the double-helical structure of DNA.

