Seat	
No.	

## M. Sc. (Bioinformatics) (Semester –II) (New) (CBCS) Examination, 2017 ADVANCED BIOINFORAMTICS

Day & Date: Wednesday, 19-04-2017 Max. Marks: 70

Time: 10.30 AM to 01.00 PM

- N.B.: 1) Part-I, Question 1 is compulsory.
  - 2) Attempt any Four questions from part-II.
  - 3) Figures to the **right** indicate **full** marks.
  - 4) Answers to the Part-I and –II are to be written in same answer booklet only.

#### Section-I

		Section-i	
Q.1	A)	Rewrite the sentence after choosing the correct from the given alternatives.  1) illustrate the relatedness of the leaf nodes without making assumptions about ancestry at all.  a) Super trees	07
		MP in phylogenetic refers to     a) Multiple Parsimony     c) Maximum Parsimony     d) Maximum Phylogeny	
		<ul> <li>3) PubMed and Medline are library databases.</li> <li>a) Visual Library b) Vertebral Library</li> <li>c) Virtual Library d) All of these</li> </ul>	
		4) is a tool in EMBOSS which gives protein statistics a) Showfeat b) Infoseq c) Pepstat d) None of these	
		5) The PAM matrices were introduced by a) Margeret Dayhoff b) Henikoff and Henikoff c) Feng and Doolittle d) None of these	
		<ul> <li>6) PAUP stands for</li> <li>a) Protein Analysis Using Proteomics</li> <li>b) Phylogenetic Analysis Using Parsimony</li> <li>c) Phylip Analysis Using Parsimony</li> <li>d) None of these</li> </ul>	
		7) BLOSUM stands for a) Block Substitute Matrix b) Block Substitution Matrix c) Block Substituent Matrix d) None of these	

	B) Definition  1) ExPASy 2) KEGG 3) PDB 4) PSI BLAST 5) BankIt 6) gap penalty 7) alpha helix	<b>80</b> 07
	Section-II	
Q.2	Answer any four the following Explain EMBOSS and its utilities and add a note on for what purpose EMBOSS is use?	14
Q.3	What is pairwise sequence alignment? Give a detailed description of Smith-Waterman algorithm?	14
Q4	Explain different type's identification of SNPs methods and a details account on SNP database	14
Q5	<ul> <li>Answer any two from the following</li> <li>a) Explain protein arrays, its basic principles and applications.</li> <li>b) Explain MUMmer and suffix tree and add a note on comparative genomics.</li> <li>c) Write a detailed note on Maximum parsimony method.</li> </ul>	14
Q.6	Write short notes on (any two) a) Phylip b) Gene prediction in Prokaryotes c) KEGG	14

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M.Sc	. (Bioinformatics) (Semester – II) (New) (CBCS) Examination, 2017 MICROBIOLOGY AND BIOTECHNOLOGY	
Day & Da	ate: Friday, 21-04-2017 Max. Marks: 70	)
Time: 10	.30 AM to 01.00 PM	
Ins	1) Part-I, Questions-1 is compulsory. 2) Attempt any four question from part- II. 3) Figures to the right indicate full marks. 4) Answer to the Part- I and Part- II are to be written in same answer Booklet only.  Section-1	
Q.1 A)	Rewrite the sentence after choosing the correct answer from the given alternatives:  1) The bacterium grows in salt concentration is called a) Acidophiles b) Alkaliphiles c) Halophiles d) Xerophiles  2) bacterial genome was synthesized artificially. a) Mycoplasma b) E. coli c) Pseudomonas d) H1N1  3) Sanger's method of sequencing makes use of a) Oligonucleotide b) Deoxynucleotide c) Dideoxynucleotide d) All of these  4) Molecular taxonomy is based on a) 28S rRNA b) 16S rRNA c) 30S rRNA d) 23S rRNA  5) The genetic material of virus is a) RNA b) DNA c) RNA or DNA d) Protein  6) Prions are basically made up of a) RNA b) Protein c) DNA d) Virus  7) plasmid contains viral fragment. a) Phagemid b) cosmid c) pBR322 d) pUC19	07
В)	Definitions: 1) Ribosomes 2) Endonuclease 3) SV40	07

- 4) BACs

- 5) Probes6) Domain of life7) Growth kinetics

### Section- II

### Answer Any Four of the following:

Q.2	Explain the general structure of prokaryotic cell with neat labeled diagram.	14
Q.3	Describe the structure and function of cloning vectors-pUC18 and pBR322.	14
Q.4	Explain the gene transfer in plant systems.	14
Q.5	<ul> <li>Answer any two from the following:</li> <li>a) Add a note on application of r-DAN technology in crop improvement.</li> <li>b) Write a note on animal tissue culture media.</li> <li>c) Explain the stages of growth curve.</li> </ul>	14
Q.6	<ul><li>Write short notes on (any two)</li><li>a) Electroporation</li><li>b) Structure of T4 bacteriophage</li><li>c) Bacterial conjugation</li></ul>	14

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## M.Sc. (Bioinformatics)(Semester – II) (New) (CBCS) Examination, 2017 BASIC BIOCHEMISTRY & IMMUNOLOGY

		BAGIO BIOGITEMIOTICI A IMMONOLOGI	
Day &	Date	e: Monday, 24-04-2017 Max. Marks	: 70
Time: 1	10.3	0 AM to 01.00 PM	
		<ul> <li>Instruction: 1) Part-I, Questions 1 is compulsory.</li> <li>2) Attempt any four question from part-II</li> <li>3) Figures to the right indicate full marks.</li> <li>4) Answer to the Part-I and Part-II are to be written same answer Booklet only.</li> </ul>	in
Q.1 /	A)	Rewrite the sentence after choosing the correct answer from the given alternatives:  1) is the main source of free energy in biological system.  a) Glucose b) Fatty acids c) ATP d) All	07
		The basic components of lipids are     a) Amino acids b) Vitamins c) Fatty acids d) glucose	
		Amino acids possessing both the charges are called     a) Divalent ions b) Zwiter ions c) dipole ions d) None	
		4) is structural polysaccharide. a) Starch b) Cellulose c) Glycogen d) Sucrose	
		5) B cells are derived from lineage. a) Erythroid b) Myeloid c) Osteoid d) Leucoid	
		6) is a secondary lymphoid organ. a) Bursa of Fabricious b) Lymph node c) Thymus d) None	
		7) The surface maker present on T cell subsets are a) CD4 & CD8 b) CD12 & CD32 c) CD18 & CD24 d) All	
E	В)	Definitions:  1. Fatty acid 2. Polysaccharide 3. Vitamin D 4. IgA 5. Phagocyte 6. Cytokine 7. CMI	07

### Section-II

### Answer any four of the following

Q.2	Explain different types of amino acids.	14
Q.3	Define Antibody. Explain its structure with a neat diagram.	14
Q.4	Write a detailed note on adaptive immunity.	14
Q.5	<ul> <li>Answer any two from the following:</li> <li>a) Add a note on types of proteins.</li> <li>b) Write a note on functions of carbohydrates.</li> <li>c) Explain different types of nucleic acids.</li> </ul>	14
Q.6	Write short notes on (any two) a) Classification of lipids b) Primary lymphoid organs c) Autoimmunity	14

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## M.Sc. (Bioinformatics) (Semester-II) (New) (CBCS) Examination, 2017 Industrial and Environmental Biotechnology

Day & Date: Monday, 24-04-2017 Max. Marks: 70

Time: 10.30 AM to 01.00 PM

#### Instructions:

- 1) Part-1, Question 1 is compulsory
- 2) Attempt any four questions from Part-II
- 3) Figures to the right indicate full marks.
- 4) Answer to the Part-I and Part-II are to be written in same answer Booklet only.

#### PART - I

### Q.1 Rewrite the sentence after choosing the correct answer from the given alternatives:

07

- 1) Stationary phase is described as
  - A) No further increase in the cell population after a maximum value
  - B) Deceleration of growth and division rate after the growth rate reaches a maximum
  - C) Acceleration of growth and division rate after the growth rate reaches a maximum
  - D) Deceleration of growth and division rate after the growth rate reaches a maximum
- 2) The number of baffles in a standard stirred tank bioreactor is
  - A) 8

B) 6

C) 4

D) 2

- 3) The specific growth rate is affected by
  - A) Substrate concentration
  - B) Product concentration
  - C) Oxygen supply
  - D) All of these
- 4) Bioreactors are used for
  - A) Large scale production of desired substances by using cells/microbes
  - B) Kill bacteria
  - C) To Store viruses
  - D) To get Chemicals

energy sources.  A) Petroleum oil B) Sunlight C) Coal D) Natural gas  6) Dendrotheraml energy is included in type of energy source. A) Conventional B) Renewable C) Non-renewable D) Both A & C  7) The forest conservation act was passed in by Indian parliament A) 1099	
<ul> <li>6) Dendrotheraml energy is included in type of energy source. <ul> <li>A) Conventional</li> <li>B) Renewable</li> <li>C) Non-renewable</li> <li>D) Both A &amp; C</li> </ul> </li> <li>7) The forest conservation act was passed in by Indian parliament</li> </ul>	
source. A) Conventional B) Renewable C) Non-renewable D) Both A & C  7) The forest conservation act was passed in by Indian parliament	
C) Non-renewable D) Both A & C  7) The forest conservation act was passed in by Indian parliament	
7) The forest conservation act was passed in by Indian parliament	
parliament	
A) 1000 D) 1000	
A) 1988 B) 1980	
C) 1981 D) 1972	
<ul> <li>B) Definitions</li> <li>1) Secondary metabolites</li> <li>2) Stationary Phase</li> <li>3) Immobilization</li> <li>4) Spargers</li> <li>5) Environmental ethics</li> <li>6) Biosensor</li> <li>7) Bio Sorption</li> </ul>	07
PART - II Answer any four of the following	
Q2 Explain about isolation of Microorganism and note on Microbial growth phases	14
Q3 Define Biosensor? Note with examples and its application	14
Q4 Describe about the environment protection and conservation	14
Q5 Answer any two of the following A) Methods of Preservation of microorganism B) Continuous fermentation C) Streptomycin production	14

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### BIOINFORMATICS (Semester - IV) (New) (CBCS) Examination, 2017

			BIOLOGICAL SIMULATION AND MODELING	
Day a	& Da	te: \	Wednesday,19-04-2017 Max. Mark	(s: 70
Time	: 02.	30 F	PM to 05.00 PM	
			N.B.: 1) Section-I, Question 1 is Compulsory.  2) Figures to the right indicate full marks.  3) Attempt Any Four questions from Section –II  4) Answers to the Section-I and Section –II are to be written in same answer Booklet only.  Section -1	oe
Q.1	A)		elect appropriate word:  Python is types of language.  a) Dynamic b) Semi-dynamic  c) Static d) None of these	07
		2)	The function creates a Python file object. a) Fopen() b) Open() c) Fileopen() d) None of these	
		3)	x = 4.5 y = 2  print x//y? what will be the output a) 2.0 b) 10.0 c) 5.0 d) 1.0	
		4)	The first in simulation is a) Calculation b)Processing c) Model building d)All	
		5)	is a base of any simulation. a) Statistics b) Mathematics c) Physics d) Chemistry	
		6)	MD in simulation stands for a) Microbial Dynamics b) Macroscopic Dynamics c) Molecular Dynamics d) None	
		7)	The first protein simulated was a) Insulin b) Trypsin inhibitor c) Polymerase d) Protease	
	1)   2)   3)   4)   5)   6)	Pyth Dyna Stati Clas	namic tic ss tem	07

		SLK-KC - 101
Q.2	Section-II  Answer any four of the following  Explain string functions in python with example.	14
Q.3	Explain working with python	14
Q.4	Write a note on principle applications of simulations	14
Q.5	Answer any two form the following  1) Write a note simulation Softwares 2) Explain python editor in details 3) Add a note on Molecular mechanics	14
Q.6	<ul><li>Write short notes on (Any two)</li><li>1) Biological simulations</li><li>2) Functions in python</li><li>3) Examples of molecular dynamics</li></ul>	14

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# M. Sc. Bioinformatics (Semester – IV) (New) (CBCS) Examination, 2017 CLINICAL BIOINFORMATICS

Day & Date: Friday, 21-04-2017 Max. Marks: 70				
Time: 02.3	Time: 02.30 PM to 05.00 PM			
	N.B.: 1) Section-I, Question 1 is con 2) Attempt <u>any four</u> from Sec 3) Figures to the right indicate 4) Answer to the Section-I and in same answer Booklet or <u>Section - I</u>	ction-II e full marks. d Section-II are to be written		
Q.1 A)	Rewrite the sentence after choosing the control the given alternatives.  1) is also called whole transciptome slap a) RNA Seq b) Chlp c) FAST A	hotgun sequencing.		
	2) Peptic ulcers is caused by  a) E.coli b) Helicobacter pylori c) H	IIV d) Fungus		
	<ul><li>3)to prepares a standardized bill for patient.</li><li>a) Payer</li><li>b) Provider</li><li>c) Occur</li></ul>	-		
	A fast quality control toolkit Illumin     a) Pyroclenar b) FAST X c) PrinSe			
	5) Cancer is caused by a) Hepatitis b) HI c) Human papilloma virus d) Al	V I of these		
	6)is a genome browser for vertebrat a) Google b) Ensembl c) Pu			
	, ,	 dministration ssessment		
В)	Definitions.  1) Cystic fibrosis  2) ADR  3) Mapviewer.  4) NGS library  5) SNPdb  6) BAM	07		

### 7) HTQC

	Section - II	
Q.2	Answer any four of the following.  Explain in detail about types of Cancer.	14
Q.3	Explain quality control tool for Next generation sequencing data in details and add a note on challenges for bioinformatics in NGS details.	14
Q.4	Genome data visualization using Ensemble and Map viewer.	14
Q.5	<ul><li>Answer any two from the following.</li><li>1) Explain Basic NGS chemistry in details.</li><li>2) Describe pathogen genomes.</li><li>3) Explain transcriptomics in details.</li></ul>	14
Q.6	Write short notes on (any two)  1) System Biology  2) Application of Pharamcovigilance  3) Challenges of HGP.	14

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## M.Sc. Bioinformatics (Semester-IV) (New) (CBCS) Examination, 2017 ADVANCED MOLECULAR BIOLOGY

		AD TAITOLD MOLLO	OLAN BIOLOGI	
Day (	& Da	te: Monday, 24-04-2017	Ma	ax. Marks: 70
Time	: 02.3	30 PM to 05.00 PM		
		3 Figures to the <b>r</b> 4) Attempts <b>any tv</b>	our questions from part-III right indicate full marks. vo questions from Q.5 to Part-I and Part-II are to be	Q. <b>7</b> .
Q.1	A)	PART Rewrite the sentence after che		ver 07
<b>Q</b> . 1	~,	from the give alternatives:  1) Western Blotting is used for _ a) DNA c) RNA	_	01
		tool is used to analyze     a) Mass spectroscopy     c) ExPassy		
		<ul><li>3) Molecular based diagnosis is</li><li>a) FTIR</li><li>c) UV-VIS</li></ul>	done using instrum b) PCR d) All	ient.
		<ul><li>4) Localization of gene in cell or</li><li>a) In vitro</li><li>c) In Situ</li></ul>	r tissuetechnique is b) In Vivo d) All	s used.
		5) technique is used to from. a) SDS PAGE c) IEF	b separate proteins in its r b) Native PAGE d) 2D PAGE	native
		<ul><li>6) technique is used to</li><li>a) In situ hybridizatio</li><li>c) Plaque hybridization</li></ul>	<ul><li>b) Colony hybridizati</li></ul>	
		7) Molecular marker is in genetic material using PCF a) Western Blot c) RFLP		<sup>-</sup> phism

	<ul> <li>Answer the following</li> <li>1) Molecular Markers</li> <li>2) Nitrocellulose membrane</li> <li>3) Dansyl chloride</li> <li>4) Dialysis</li> <li>5) Plaque Hybridization</li> <li>6) Microarray</li> <li>7) Authoradiography</li> </ul>	07
	PART-II	
Q.2	Explain the Primer designing parameters using Oligo 4 software.	14
Q.3	Explain in detail the instrumentation an applications of HPLC and GLC.	14
Q.4	Explain protein sequencing by Edman degradation method.	14
Q.5	<ul> <li>Answer any two of the following.</li> <li>a) Describe variable number tandem repeat (VNTR).</li> <li>b) Write a note on affinity chromatography with neat labeled diagram.</li> <li>c) Describe 2D gel electrophoresis.</li> </ul>	14
Q.6	Short notes (Any two): a) SNP b) cDNA Library c) Site Directed Mutagenesis.	14

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7) GBIF

### M.Sc. Bioinformatics (Semester – IV) (New CBCS) Examination, 2017 EMERGING AREAS OF BIOINFORMATICS

Day 8	ay & Date: Wednesday, 26-04-2017 Max. Marks: 70			70		
Time:	02.3	80 PM to 05.00	PM			
		N.B. :	<ol> <li>Attempt any</li> <li>Figures to th</li> <li>Answers to the</li> <li>same answer</li> </ol>	r Booklet only.	om part-II	
Q.1	A)	Rewrite the se		<u>tion - I</u> hoosing the corr	ect answer from	07
		the given alte 1) ChEMBL is				
		a) EBI	b) NCBI	c) SIB	d) ExPasy	
				icate the molecula	ar graph to and from	
		the comput a) Linear no c) SMILES		b) Conne d) MOL	ction table	
		3) IMGT/HLA a) NCBI	database is pre b) DDBJ		d) DNAS	
		4) a) IMGT is	the immunogen b) IPD	ic database. c) IT IS	d) All	
		5) Every subr a) SS#		receives a submit c) SS	ted SNPID is d) S#	
		b) Ocean b c) Ocean b	iology informati iogeographic in iogeographic in iogeographic in iology initiative	formation system itiative system		
		7)first a) Richard c) Eric Dre	Feynman	nanotechnology. b) Norio d) Sumio	_	
	B)	Definition 1) IMGT 2) 3D compout 3) SIFT 4) Top Down 5) dbSNP 6) SDF				07

	Section - II	
Q.2	Answer any four of the following.  Explain the different chemical database and add a note on SMILES notation in details.	14
Q.3	Give a detailed account on immunoinformatics and explain the bioinformatics strategies for better understanding of immune function.	14
Q.4	Explain the brief principles of Taxonomy and add a note on phylogeny in biodiversity informatics.	14
Q.5	Answer any two from the following.  1) Explain SNP and its application in details. 2) Describe the Properties of nanoparticals. 3) Explain Chemical structure representation.	14
Q.6	Write short notes on (any two)  1) FTIR  2) MOL file format  3) TDWG	14

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## M.Sc. (Bioinformatics) (Semester-IV) (New) (CBCS) Examination, 2017 MOLECULAR MEDICINE

		MOLECULA	R MEDICINE		
•		te: Wednesday, 26-04-2017 30 PM to 05.00 PM		Max. Mark	(s: 70
		4) All questions	ns of <b>Section-I</b> are <b>c Four</b> questions fron carry equal marks. nd labeled diagrams	n section <b>II</b> .	ssary.
•	•	_	ION – I		
Q.1	A)	Rewrite the sentence after of the from the given.	choosing the corre	ct answer	07
		<ul> <li>1) In DNA fingerprinting</li> <li>used.</li> <li>a) Variable number of tand</li> <li>b) Verified number of tand</li> <li>c) Versatile number of trand</li> <li>d) Variable number of trand</li> </ul>	dem repeats dem repeat nsverse repeat	es of DNA is	
		<ul> <li>2) Stem cell exhibitsp</li> <li>a) Only potency</li> <li>b) Potency and self renew</li> <li>c) Potency and non renew</li> <li>d) Only self-renewable</li> </ul>	vable		
		3) MHC Antigen in mouse is a) HLA b) H-2	known as c) ASB	d) HSB	
		4) The -globin gene of haem number	oglobin is located or	n chromosome	
		a) 11 b) 12	c) 16	d) 18	
		5) Gene is mutated i a) CFTR b) Actin	n cystic fibrosis. c) Cadherin	d) Fibrin	
		<ul> <li>6) Stem cell exhibitsp</li> <li>a) Only potency</li> <li>b) Potency and self renew</li> <li>c) Potency and non renew</li> <li>d) Only self-renewable</li> </ul>	vable		
		<ul><li>7) Hematopoietic stem cells a</li><li>a) Pluripotent</li><li>c) Unipotent</li></ul>	are b) Totipotent d) Oligopotent.		

	B) Define the terms:  1) Totipotency 2) Recombination 3) Amniocentesis 4) DNA fingerprinting 5) Microarray. 6) Plasmids 7) Sickle cell anemia	07	
	SECTION – II Answer any four of the following:		
Q.2	Write a brief account on Stem cells and its properties.	14	
Q.3	Explain in detail Cystic fibrosis with labeled diagram.	14	
Q4	Explain in detail steps involved in during discovery and its design.	14	
Q.5	<ul> <li>Answer any two of the following</li> <li>A) Give an account on Huntinston's disease.</li> <li>B) Explain different nature and sources of drug.</li> <li>C) Explain in brief human genome project.</li> </ul>		
Q.6	<ul> <li>Write Short notes on any TWO of the following</li> <li>A) Write a note on induced pluripotent stem cells.</li> <li>B) Describe alzheimer's disease.</li> <li>C) Explain in-vivo and ex-vivo gene therapy.</li> </ul>	14	