Seat	Set	D
No.	Set	

			Bic	oinformatic	s
			BASIC B	BIOINFORM	ATICS
-			hursday, 16-11-2017 M to 01.00 PM		Max. Marks: 70
Instr	uctio		 Section-1, Question 1 i Attempt any four quest Figures to the right ind Answer to the Section- Booklet only. 	ions from Sec icate full mark	tion-II.
				Section - I	
Q.1	A)		ven alternatives:	compares a peotide sequer b)	e correct answer from the orotein query against the all six aces database. tblastx all of these
		2)	OMIM stand fora) Ontology Mechanismb) Online Machine Interiorc) Online Machine inheriord) Online Mendelian Inh	In Man face mapping ritance in Man	
		3)	GUI stands fora) Graphical User Interface) Genome User Interface	ace b)	Genic User Interphase None of these
		4)	SVM stands for a) Species Vector Mach c) System Vector Mach	nine b)	Support Vector Machine None of these
		5)	The protein secondary dangler (a) NCBI c) NRL-3D	b)	GenBank Pfam
		6)	FTP stands for		File Transfer Practice Field Transfer Protocol
		7)	NBRF it stands fora) National Biomedical Res b) Nation Biological Res c) National Biomedical d) Nation Biological Res	search Fund Research Fu	nd

		SLR-MC	-62
	,	Definitions. 1) Sequence similarity 2) BLASTn 3) Phylip 4) Enterz 5) PIR 6) Neural Network 7) SVM	07
		Section - II	
	Ansv	wer any four of the following.	
Q2	Expla	ain Neural Networks and add a note o Support Vector Machine.	14
Q3		the detailed account on NCBI and explain the nucleotide sequence base in details.	14
Q4		t is String Matching Algorithm and add a note on BLAST and FASTA ence comparison.	14
Q5	a) Eb) G	wer any two of the following. xplain the Gene array analysis and its applications. ive an account on Entrez and SRS tools. xplain ClustalW and TreeView in details.	14
Q6	a) G b) P	e short note any two of the following. senBank File Format IR pplications of Bioinformatics.	14

Seat	Set	D
No.	Set	

	Bioinforma CELL BIOLOGY AN	
•	te: Saturday, 18-11-2017 30 AM to 01.00 PM	Max. Marks: 70
Instructio	only.	Part-II marks. are to be written in same answer Booklet
	Section -	· I
Q.1 A)	Rewrite the sentence after choosing given alternatives: 1) The dihybrid test cross ratio is a) 9:3:2:1 c) 9:3:2:2	
	, .	b) Only dominant allele d) All of these
	3) Poly A tail is a characteristics of _a) Prokaryoticc) Both a & b	mRNA. b) Eukaryotic d) None of these
	4) E site, P site and A site are founda) rRNAc) tRNA	on b) mRNA d) hnRNA
	5) is involved in cell re a) Ribosome c) Nucleus	espiration. b) Mitochondria d) lysosome
	6) Cell growth takes place ina) Mc) G2	phase. b) S d) All of these
	7) is a secondary mesa) Hormonec) Cyclic AMP	ssenger. b) Acetylcholine d) None
В)	 Definitions 1) Point mutation 2) Ligase 3) Transcriptomics 4) Lysosome 5) Autotroph 6) GPCR 7) Monocistronic 	07

Section - II

	Answer any four of the following	
Q2	Explain the process of DNA replication in prokaryote with diagram.	14
Q3	What is operon? Explain the mechanism of lac operon concept	14
Q4	Write a note on various cell organelles.	14
Q5	 Answer any two of the following a) Explain the process of transcription in eukaryotes. b) Write a note on DNA repair mechanism. c) Add a note on regulation of cell cycle. 	14
Q6	Answer any two of the followinga) Post translational modifications.b) Receptors and carriers of cell.c) Signal transduction.	14

Seat	Set	D
No.	Set	

		Bioinform INTRODUCTION TO HTM		
•		Гuesday, 21-11-2017 AM to 01.00 РМ	Max. Mar	ks: 70
Instruc	etions:	 1) Section-1 is compulsory. 2) Attempt any four questions from 3) All Questions carry equal marks 3) Figures to the right indicate full of the properties. 4) Draw neat and labeled diagram. 	narks.	
		Section	ı - I	
Q.1 A	alt 1)	complete the sentences by selecting ternatives: A tag is used to display a) picture c) img	the image. b) image d) src	07
	2)	a) <hr/> c) c) = tag inserts a line horizon a) representation = "horizontal" > tag inserts a line horizon a) representati	b) <line></line>	
		a) <html> c) <title></th><th>displayed on the page are written in b) <head> d) <body></th><th></th></tr><tr><td></td><td>4)</td><td>a) Connector c) Hyperlink</td><td>b) Lined) None of the above</td><td></td></tr><tr><td></td><td>5)</td><td>The first page of a website is called a) Design Page c) First page</td><td>d b) Home page d) Main page</td><td></td></tr><tr><td></td><td>6)</td><td>The number of occurrences of data a) Class limits c) Cumulative frequency</td><td>a value is called b) Frequency d) Relative frequency</td><td></td></tr><tr><td></td><td>7)</td><td>The most frequently occurring value) Spread c) Skewness</td><td>ue in the data set is called b) Mode d) Median</td><td></td></tr><tr><th>В</th><th>1)
2)
3)
4)
5)
6)</th><th>Anchor Anchor WWW Hyperlink Head tag Sample space Cumulative frequency Class mark</th><th></th><th>07</th></tr></tbody></table></title></html>		

Section - II

Q2 Explain briefly advantages and disadvantages of World Wide Web (WWW).
Q3 Explain in detail HTML History.
Q4 Find the correlation co-efficient between X and Y from the following data:14

Χ	17	18	19	19	20	20	21	21	22	23
Υ	12	16	14	11	15	19	22	16	15	20

Q5 Answer any two of the following

14

- a) Explain basic tags with attributes and example.
- **b)** Explain Frame tag attributes with example.
- c) Find the mode from the following data.

Marks	0-10	10-20	20-30	30-40	40-50	50-60
No. of students	3	8	15	20	10	4

Q6 Answer any two of the following

14

- a) Write a note on Ordered & Non-Ordered lists
- **b)** Merits and demerits of mean deviations.
- c) Explain Table tag attributes with example.

Seat	Set	D
No.	Set	

M.Sc. (Semester - I) (CBCS) Examination Oct/Nov-2017 Bioinformatics INTRODUCTION TO PROGRAMMING LANGUAGES & PROGRAMMING THROUGH C & C++

11411	NOL			GH C & C++	,
			hursday, 23-11-2017 M to 01.00 PM	Max. Marks: 7	70
Instru	ictioi		 Part-1, Question 1 is compted Attempt any four questions Figures to the right indicate All questions carry equal m Draw neat and labeled diag 	from Part-II e full marks. parks.	
			Sect	tion – I	
Q.1	A)	alt	ternatives:-	function keyword is used. b) int d) void	07
		2)	,	is a logical AND represented. b) d) &&	
		3)	String constants should be en a) Single Quotes c) Double Quotes	nclosed between b) Both A and B d) None of these	
		4)	To avoid return value in main a) Const c) function	b) int d) void	
		5)	The friend function in C++ is that class. a) Public c) Protected	b) Private d) Virtual	
		6)	OOPs stand fora) Oracle Oriented programmb) Object Oriented programmc) Operand Oriented programmd) None of these	ming ming	
		7)	is an operator winside the block. a) :: c) +	which is used to access global variable b) ++ d) >	

B) **Definitions** 07 1) Variable 2) Algorithm 3) Keyword 4) Compiler 5) Object 6) Class 7) Pointer Section - II Answer any four of the following Q2 Difference between POP and OOP. 14 14 Q3 Write a C program for sum of 10 numbers along with algorithm & flowchart. Q4 Describe concept of inheritance & types of inheritance with its example in C++ 14 Answer any two of the following **Q5** 14 a) What is operator? Explain with examples different types of operator in C programming. **b)** Briefly explain the structure of C Program. c) Difference between entry controlled & exit controlled loops. Short answer (Any two):-14 Q6 a) File handling in C programming. **b)** Polymorphism c) Write a C++ program larger of three numbers.

Seat	Set	D
No.	Set	

			Bioinform PLANT BREEDING AND			
			「hursday, 23-11-2017 ∖M to 01.00 PM		Max. Marks	: 70
nstr	uctio	ns:	 Part-1, Question 1 is compulsory Attempt any four questions from Figures to the right indicate full r Draw neat and labeled diagram. All questions carry equal marks. 	Par nark		
			Section -	– I		
Q.1	A)	gi	ewrite the sentence after choosing ven alternatives: Synthetic seed is produced by enc	-		07
			a) Sodium alginate c) Sodium acetate	,	Sodium nitrate Sodium sulphate	
		2)	The production of secondary metal a) Protoplas c) Sodium acetate	b)	es requires the use of Cell suspension Auxiliary bud	
		3)	Shoot regeneration is promoted by a) NAA c) BAP	b)	2, 4-D IAA	
		4)	A plant breeder wants to develop a should do first a) Mutation c) Hybridization	b)	ease resistant variety, what he Selection Production of crop	
		5)	Hybrids which are superior over pa a) Inbreeding c) Recessive	b)	s are called Dominant Heterosis	
		6)	Major food crops have originated n a) Ocean c) Desert	b)	ly from Mountain Plain	
		7)	In micropropagation, virus free plana) Shoot tip culture c) Protoplast culture	b)	can be obtained through Haploid culture Embryo culture	
	B)	1) 2) 3) 4) 5) 6)	efinitions Ideotype breeding MAS Protoplast Somaclonal Cryopreservation Secondary metabolites Molecular pharming.			07

Section - II

Answer a	ny four	of the	follo	wing
----------	---------	--------	-------	------

Q2	Give a detailed account on breeding methods for self-pollinated, crosspollinated and clonally propagarted.	14
Q3	Explain about genetic and physiological basis of abiotic stress tolerance.	14
Q4	Discuss in details plant regeneration pathways.	14
Q5	 Answer any two of the following a) Use of somaclonal and gametoclonal variation in crop improvement. b) What is mutation? Explain in-vitro mutagenesis. c) Principles and types of cryopreservation method. 	14
Q6	 Answer short notes any two of the following:- a) Biotransformation b) Metabolic engineering for production of secondary metabolities. c) Role of markers in stress resistance breeding. 	14

Seat	Sat	D
No.	Set	

IV	ı.Sc.	(Semeste		CBCS) Examination Oct/Nov-2017
		BIOLOGI		SE MANAGEMENT SYSTEM
•		Thursday, 10 PM to 05.00		Max. Marks: 70
Instruct	tions	2) Attempt3) Figures4) Draw ne	Question 1 is com any four question to the right indica eat and labeled di tions carry equal	ns from Part-II. ate full marks. iagram.
			Р	PART – I
Q.1 A	-	ewrite the siven alterna		thoosing the correct answer from the 0
	1		is a collecti	ion of recorded data. b) Software d) Scheme
	2	Relational a) Chris I c) E. F. C	Date	nted by b) Hugh Darwen d) Bill Gates
E	3		nt transaction. .ck	akes 'permanent' all changes performed in b) Truncate d) None of these
_	4	a) Describ) Manipc) Addition	ulation & process on of new structur	tructure of database.
	5		essing power of F L	e data manipulating power of SQL with the Procedural languages. b) SQL d) PQL
	6	or view: a) Start c) Join	is a query th	hat retrieves rows from more than one table b) End d) All of the mentioned
	7	a) Equijo c) Both		ed in a join query have no join condition: b) Cartesian d) None of the mentioned

SLR-MC-75 Definitions. 07 B) 1) Table 2) Foreign Key. 3) Data Independence. 4) Cardinality. 5) Procedure. 6) Rollback 7) View. PART - II Answer any four of the following. Q2 Explain Limitations of traditional file processing system & Advantages of 14 DBMS? Q3 Explain Components of DBMS. 14 Q4 Explain Data mining with Types of Data mining techniques and different types 14 of application. Q5 Answer any two of the following. 14 a) Users of DBMS. b) ER Symbols. c) Normalization. Answer any two of the following. Q6 14 a) DML Commands. **b)** Join Operations. c) What is PL/SQL? Features of PL/SQL?

Seat	So.	D
No.	Set	

		Bioinform ADVANCED BIOPHYSI	
		Saturday, 18-11-2017 PM to 05.00 PM	Max. Marks: 70
Instructi	ons:	only.	m Part-II I marks. I are to be written in same answer Booklet
		PART -	-1
Q.1 A)		ewrite the sentence after choosi iven alternatives: The wavelength Visible light is a) 200-780nm c) 200-400nm	_
	2)) The type of IR which deals with v a) Near IR c) Far IR	vibrational energy is b) Mid-IR d) All
	3)	Non-metals are usually a) Electron donar c) Both a & b	b) Electron acceptor d) none
	4)	In X-ray crystallography, moleculea) Solidc) Gaseous	les are in state. b) Liquid d) All
	5)) Radio waves are used as sourcea) UVc) IR	e in spectroscopy. b) Visible d) NMR
	6)	Population inversion is associatea) Spectroscopyc) Crystallography	
	7)	invented the compoa) Antony Van Leeuwenhoek'sb) Hans Jansssen	
B)	1) 2) 3) 4) 5)	efinitions.) Electromagnetic wave) Atomic orbit.) Quartz Cuvette.) Optical activity.) MALDI.) Bragg's Law.) TEM.	07

PART - II

	Answer any four of the following.	
Q2	Add a note on instrumentation and applications of IR spectroscopy.	14
Q3	Define chemical bond. Explain different types of bonds.	14
Q4	Write a note on theory and instrumentation of UV spectroscopy.	14
Q5	 Answer any two of the following. a) Explain the principle of NMR. b) Add a note on applications of CD and ORD. c) Explain the types of ionization for mass spectroscopy. 	14
Q6	Answer any two of the following.a) Principle of X-ray crystallography.b) Applications of Laser.c) Microscope	14

M.Sc. (Semester - III) (New) (CBCS) Examination Oct/Nov-2017 Bioinformatics

C	OMF	PU1	TATIONAL STRUCTURE	BIOLOGY AND DRUG DESIGNING
•			uesday, 21-11-2017 M to 05.00 PM	Max. Marks: 70
Instru	ıctio		 Part-1, Question 1 is compe Attempt any four questions Figures to the right indicate Answer to the Part-I and Part only. 	from Part-II.
			PA	RT – I
Q.1	A)	gi	ven alternatives:	oosing the correct answer from the 0
		1)	validation of NMR structure.	service for analysis, visualization and
			a) NMR core	b) NMRclust
			c) Vivaldi	d) VASCO
		2)	Drug metabolism takes place	
			a) Liver	b) Kidney
			c) Brain	d) All of these
		3)	MNT is a) Protein database	b) Pathway database
			c) Interaction database	d) All of these
		4)	,	natic diagram of all interaction between
			a) Nuclpolt	b) Thumbnail Image
			c) Wiring Diagram	d) Ligplot
		5)		otein models generated by Modeller
			Program. a) ModBase	b) 3D Crunch
			c) PDB	d) PDBsum
		6)	•	•
		٠,	a) Keg draw	b) ACD lab
			c) Chem. Sketch	d) All of these
		7)	The 3D structure of Protein of	an be modeled by
			a) PSI PRED	b) Swiss Model
			c) GOR	d) PHD

B) **Definitions** 07 1) Template. 2) Efficacy. 3) PDBeNMR. 4) Potency. 5) Jmol 6) QSPR. 7) Domain PART - II Answer any four of the following Q2 Explain in detail Protein-lipid and protein-DNA interaction with database and 14 applications. 14 Q3 Explain in detail in silico drug discovery process and challenges. Q4 Explain RNA structures prediction method and a note on molecular 14 visualization software's. 14 Q5 Answer any two of the following a) Write a note on QSAR and its applications. **b)** Write a note on mutations in drug targets. c) Write a note on structure visualization with tools. Q6 Answer any two of the following 14 a) CATCH and SCOP. **b)** Drug absorption. c) Protein Folding.

Seat	
No.	

M.Sc. (Semester - III) (New) (CBCS) Examination Oct/Nov-2017 Bioinformatics MOLECULAR MEDICINE

		MOLECULAR MED	ICINE
•		e: Tuesday, 21-11-2017 30 PM to 05.00 PM	Max. Marks: 70
Instr	uctio	ns: 1) All questions of Section I are computed2) Answer any four questions from section3) All questions carry equal marks.4) Draw neat and labeled diagrams who	tion II
		SECTION - I	
Q.1	A)	· · · · · · · · · · · · · · · · · · ·	
		,	ed that demonstrates the desired o) Genome d) Iron
		,	e. o) CFTR d) Marfan
		4) Stem cell exhibits propertion a) Only potency k c) Potency and non renewable contact the contact and contact are contact as a contact and contact are contact as a contact and contact are contact and contact are contact as a contact are contact as a contact are contact and contact are contact and contact are contact as a contact are contact and contact are contact are contact and contact and contact are contact are contact and contact are contact and contact are contact are contact and contact are contact are contact and contact are contact an	Potency and self renewable
		,	obin is located on chromosome o) 12 d) 18
		· · · · · · · · · · · · · · · · · · ·	D) Totipotent d) Oligopotent
		,	 o) Chagas disease d) Cystic fibrosis
	B)	 Definitions 1) Totipotency. 2) Recombination. 3) Microarray. 4) Lead optimization. 5) Magic bullets. 6) Down's syndrome. 7) Functional cloning. 	07

SECTION - II

Q2	Define absorption. Explain in detail factor affecting absorption and add a note on pharamacogenetics.	14
Q3	Explain in detail viral and non viral methods of gene transfer.	
Q4	Explain In brief properties, types and applications of adult stem cells.	
Q5	 Answer any two of the following a) Give an account on phenylketonuria. b) Explain in brief gammaglobelinemia. c) Give an account on human genome project. 	14

Q6 Answer any two of the following

14

- a) Chorionic villus sampling and its applications.
- b) Blood and blood group antigens.c) Parkinson's disease.

Answer any four of the following