SLR-RP-44



| Seat | |
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| No. | |

M.Sc. (Part – I) (Semester – II) Examination, 2015 BIOINFORMATICS

Paper - I: Advanced Bioinformatics

Day and Date : Thursday, 16-4-2015 Total Marks : 70

Time: 11.00 a.m. to 2.00 p.m.

Instructions: 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 Part II are to be written in same anwer Booklet only.

PART-I

| 1. | , | Rewrite the sentence after choosing lternatives. | ng the correct answer from the given |
|----|---|--|--------------------------------------|
| | 1 | PubMed and Medline are | library databases. |
| | | a) Visual Library | b) Virtual Library |
| | | c) Vertebral Library | d) All of these |
| | 2 | 2)is a tool in EME | BOSS which gives protein statistics. |
| | | a) Showfeat | b) Pepstat |
| | | c) Infoseq | d) None of these |
| | 3 | 3) In Dali-lite program graphical re | esult is viewed byViewer. |
| | | a) Jet | b) Jmol |
| | | c) Jlib | d) All of these |
| | 4 | 4) Each amino acid corresponds t | o a turn in an alpha helix. |
| | | a) 120° | b) 85° |
| | | c) 100° | d) None of these |
| | 5 | 5) N-W algorithm was published ir | 1 |
| | | a) 1981 | b) 1970 |
| | | c) 1980 | d) 1987 |



| | 6) | can act as a capture | e m | olecule. | |
|----|------------------|--|------|---------------------------------|-----|
| | | a) Proteins | b) | Enzymes | |
| | | c) Antibodies | d) | All of these | |
| | 7) | In S-W algorithm the trace back beg | gins | s at the value for | und |
| | | anywhere in the matrix. | | _ | |
| | | a) Minimum | , | Zero | |
| | | c) Maximum | d) | None of these | |
| | • | finitions: | | | 7 |
| | • | Genomics | | | |
| | , | PHD method | | | |
| | • | Rooted tree | | | |
| | | Megablast | | | |
| | , | KEGG | | | |
| | - | Capture molecules | | | |
| | 7) | Synteny. | | | |
| | | PART – | Ш | | |
| An | swer a | Iny four of the following : | | | |
| | | is pairwise sequence alignment? Giv | 'e a | detailed description of Smith- | |
| ۷. | | man algorithm hat is pairwise sequen | | - | 14 |
| 3. | Explai | in the prediction of protein structure us | sin | g PHD method. | 14 |
| 4. | Explai used ? | in EMBOSS and its utilities and add a r ? | note | e on for what purpose EMBOSS is | 14 |
| 5. | | are promoters splice sites, regulatory i bioinformatics tools and interpretatio | _ | | 14 |
| 6. | Answe | er any two from the following : | | | 14 |
| | a) Ex | plain taxonomy and phylogeny and ac | dd a | note on maximum parsimony. | |
| | b) Ex | plain PSI-BLAST algorithm | | | |
| | c) Giv | ve a description on the secondary stru | ıctı | ıral elements. | |
| 7. | Write | short notes on (any two) : | | | 14 |
| | a) HM | /IMer | | | |
| | b) SA | GE Database | | | |
| | c) Lo | ops and Coils. | | | |
| | | | | | |



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M.Sc. Bioinformatics (Part – I) (Semester – II) Examination, 2015 Paper No. – II: MICROBIOLOGY AND BIOTECHNOLOGY

Day and Date: Saturday, 18-4-2015 Total Marks: 70

Time: 11.00 a.m. to 2.00 p.m.

Instructions: 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 Part II are to be written in same answer booklet only.

PART-I

| | llternatives. | is used for biologi | cal method of gene | e transfer. | - | |
|---|----------------------|-------------------------------------|---------------------|---------------------|---|--|
| | | | c) both a and b | | | |
| 2 | 2) Sanger's metho | d of sequencing m | akes use of | , | | |
| | a) Oligonucleoti | de | b) Deoxynucleotic | de | | |
| | c) Dideoxynucle | eotide | d) All of these | | | |
| 3 | 3) Prions are basic | ally | | | | |
| | a) RNA | b) Proteins | c) DNA | d) Virus | | |
| 4 | l) The locomotory | structures of bacte | eria are | | | |
| | a) Cilia | b) Flagella | c) Both a and b | d) None | | |
| 5) The standard tissue culture medium stands with the name of | | | | | | |
| | a) Haberlandt a | nd Skoog | b) Murashige and | d Skoog | | |
| | c) Cocking and | Skoog | d) Miller and Sko | og | | |
| 6 | 8) | _ is used for biolog | gical method of gen | ne transfer. | | |
| | a) bacteria | b) virus | c) both a and b | d) none | | |
| 7 | · · | process the ge hrough conjugatio | | nsferred from donor | | |
| | a) Transduction | | b) Conjugation | | | |
| | a) Tues of a was at: | on | d) Nama of these | | | |

| В. | Definitions: | 7 |
|----|---|----|
| | 1) Probe | |
| | 2) HindIII | |
| | 3) Primers | |
| | 4) YACs | |
| | 5) Histone proteins | |
| | 6) Sex pilli | |
| | 7) Viroids. | |
| | PART – II | |
| Ar | nswer any four of the following. | |
| 2. | Explain general characteristics and classification of plant viruses. | 14 |
| 3. | Explain the structure and replication of bacteriophage lambda. | 14 |
| 4. | Describe the structure and function of cloning vectors-pUC 18 and pBR322. | 14 |
| 5. | Explain the different methods of gene transfer in plant and animal systems. | 14 |
| 6. | Answer any two from the following. | 14 |
| | a) Explain the organization of viral genome. | |
| | b) Applications of recombinant DNA Technology. | |
| | c) Write a note on somatic and germline gene therapy. | |
| 7. | Write short notes on (any two). | 14 |
| | a) Transformation | |
| | b) Bacterial growth kinetics | |
| | c) Cosmids. | |
| | | |

SLR-RP - 45



| Seat | |
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| No. | |

C) Neutrophil

M.Sc. (Part - I) (Semester - II) Examination 2015

| | · | BIOINFORM Basic Biochen | ΙΑΤ | ICS | | | |
|-------|--|--|---------------------------------|---|--------|-----------------|----------|
| • | ate : Tuesday, 21-4 0 a.m. to 2.00 p.m. | | | | | Total Ma | rks : 70 |
| Insti | 3) Figur 4) Ansv | – I , Question 1 is mpt any four que res to the right in vers to the Part - e answer bookle | estic ndica - I al | ns from Part ate full marks nd Part – II ar | S. | oe written in | 1 |
| | | PART - | - I | | | | |
| • | vrite the sentence a | after choosing th | e co | rrect answer | from t | the given | 7 |
| 1) : | Standard free ener | gy is denoted by | | · · · · · · · · · · · · · · · · · · · | | | |
| , | A) ΔE° | B) ∆G° | C) | ΔH° | D) | ΔF° | |
| 2) _ | is the n | nain source of fre | ee e | nergy in biolo | gical | system. | |
| , | A) Glucose | | B) | Fatty acids | | | |
| (| C) ATP | | D) | All | | | |
| 3) , | Amino acids posse | ssing both the ch | narg | es are called ₋ | | | |
| , | A) Divalent ions | | B) | Zwiter ions | | | |
| (| C) Dipole ions | | D) | None | | | |
| 4) _ | is a | structural polysa | acch | aride. | | | |
| , | A) Starch | | B) | Cellulose | | | |
| (| C) Glycogen | | D) | Sucrose | | | |
| 5) _ | is an 6 | example of agran | uloc | yte. | | | |
| , | A) B cell | | B) | Basophil | | | |

D) Eosinophil

| SLI | R-RP – | 46 | | | | | |
|-----|----------------|-----------------------------|----------------------|------------|------------|----|------|
| | 6) | is | a secondary lymp | hoic | organ. | | |
| | | A) Bursa of Fabri | cius | B) | Lymph node | | |
| | | C) Thymus | | D) | None | | |
| | 7) | Antibodies are pro | - | | | | |
| | | A) B cells | B) T cells | C) | NK cells | D) | None |
| | • | finitions : | | | | | 7 |
| | - | Free energy | | | | | |
| | - | Glycosidic bond | | | | | |
| | - | Enzyme | | | | | |
| | 4) | Phagocyte | | | | | |
| | 5) | IgM | | | | | |
| | 6) | Cytokine | | | | | |
| | 7) | CMI. | | | | | |
| | | | PART - | - II | | | |
| Ar | nswer a | ny four of the follo | owing: | | | | |
| 2. | Explai | in the structural cla | assification of prot | eins | | | 14 |
| 3. | Define | e carbohydrate. Ad | d a note on their c | lass | ification. | | 14 |
| 4. | Write | a detailed note on i | innate immunity. | | | | 14 |
| 5. | Explai | in different types of | antigen-antibody | inte | ractions. | | 14 |
| 6. | Answe | er any two from the | e following : | | | | 14 |
| | a) Wr | ite a note on ATP a | as energy source. | | | | |
| | b) Ad | d a note on factors | affecting enzyme | e act | ivity. | | |
| | c) Ex | plain the humoral n | nediated immunity | / . | | | |
| 7. | Write | short notes on (an | y two) : | | | | 14 |
| | | otein folding | - , | | | | |
| | , | ıcrophages | | | | | |
| | - | sorders of immune | system. | | | | |
| | - | | - | | | | |

SLR-RP - 47



| Seat | |
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| No. | |

M.Sc. (Part – I) (Semester – II) Examination, 2015 BIOINFORMATICS

| Paper – | IV : Programming | in Object Oriente | d Languages | | | |
|---|-----------------------------|--|------------------------------|--|--|--|
| Day and Date : Thur Time : 11.00 a.m. to | - | | Total Marks : 70 | | | |
| Instructions : | , , | r questions from Part I ht indicate full mark Part – I and Part – II a | : - II . :s. | | | |
| | PA | ART – I | | | | |
| A) Rewrite the alternatives | sentence after choosir : | ng the correct answe | r from the given 7 | | | |
| | of the follow | | ret and execute java | | | |
| | asses hosted by HTMI | | | | | |
| a) Apple | | b) Appletwatcher | | | | |
| c) Apple | | d) None of these | | | | |
| | is a single se | | | | | |
| a) Vecto | r b) Thread | c) Applet | d) Program | | | |
| 3) The clas | s at the top of exception | on class hierarchy is _ | | | | |
| a) Arithm | netic Exception | b) Throwable | | | | |
| c) Class | | d) Exception | | | | |
| 4) my \$val= | ='x'; | | | | | |
| print \$va | l; | | | | | |
| a) scalar | b) empty value | e c) string | d) none of these | | | |
| 5) Java sou | ırce code is complied ir | nto | - | | | |
| a) .obj | b) .exe | c) source code | d) bytecode | | | |

SLR-RP-47

| | 6) PERL stands for | |
|----|--|----|
| | a) Practical Extraction Report Language | |
| | b) Preparation Extraction Report Language | |
| | c) Practical Extraction Review Language | |
| | d) None of these | |
| | 7) The applet class is in package. | |
| | a) java.util b) java.io c) .java.lang d) java.applet | |
| | B) Definitions: | 7 |
| | 1) Constants | |
| | 2) Split function | |
| | 3) JVM | |
| | 4) Perl | |
| | 5) Scalar | |
| | 6) Perl Hash | |
| | 7) Applet. | |
| | PART – II | |
| Ar | nswer any four of the following : | |
| 2. | Explain hash variables and functions and write a Perl script using to display three letter and one letter amino acid code. | 14 |
| 3. | Explain array in Perl with its example. | 14 |
| 4. | Explain applet and applet life cycle of with example. | 14 |
| 5. | Explain array and hash variable in Perl with example. | 14 |
| 6. | Answer any two from the following: a) Explain constants in java with example. b) Design a registration page using applet. c) Explain interface in java. | 14 |
| 7. | Write short notes on (any two): a) Java features b) Perl in bioinformatics c) Exception handling in java. | 14 |
| | | |



| Seat | |
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| No. | |

a) Insulin

c) Polymerase

M.Sc. (Bioinformatics) (Part – II) (Semester – IV) Examination, 2015 Paper – I: BIOLOGICAL SIMULATION AND MODELING (New) (CGPA Pattern)

| | .,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | (CGPA Pati | ern) | |
|----|--|---|--|-------------------------|
| • | Date : Thursday, 16 00 p.m. to 6.00 p.m. | | | Total Marks : 70 |
| In | 2) Ai 3) Fi 4) Ai | art – I , Question 1 is ttempt any four que igures to the right in nswers to the Part - ame answer booklet | estions from Part – I ndicate full marks. - I and Part – II are t | |
| | | PART – I | | |
| | ewrite the sentence ternatives : | after choosing the | correct answer from | n the given 7 |
| 1) | | types of language b) semi-dynamic | | d) none of these |
| 2) | Thefur a) fopen() | nction creates a Pyth b) open() | _ | d) none of these |
| 3) | x = 4.5 y = 2 print a) 2.0 | x//y ? What will be to b) 10.0 | the output ? c) 5.0 | d) 1.0 |
| 4) | The first step in si a) calculation | mulation is b) processing | c) model building | d) all |
| 5) | | se of any simulation. b) Mathematics | | d) Chemistry |
| 6) | MD in simulation s a) Microbial Dyna c) Molecular Dyna | mics | b) Macroscopic Do | ynamics |
| 7) | The first protein si | mulated was | | |

b) Trypsin inhibitor

d) Protease

| SLR-RP - 60 | |
|---|----|
| B) Definitions: 1) Python 2) Dynamic 3) Static 4) Class 5) System 6) SIR 7) Energy. | 7 |
| PART – II | |
| Answer any four of the following: | |
| 2. Explain string functions in python with example. | 14 |
| 3. Explain working with files in python. | 14 |
| 4. Write a note on principles and applications of simulations. | 14 |
| 5. Explain population model in simulation with examples. | 14 |
| 6. Answer any two from the following: 1) Write a note on simulation software. 2) Explain python editor in details. 3) Add a note on Molecular mechanics. | 14 |
| 7. Write short notes on (any two):a) Biological simulation.b) Functions in python.c) Examples of molecular dynamics. | 14 |



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c) Sculpture

| • | – II) Semester – IV (BIOINFORM per No. II – Biodivei | IATICS (New) | |
|----------------------------------|---|--|---|
| Day and Date : Sat | urday, 18-4-2015 | | Total Marks: 70 |
| Time: 3.00 p.m. to | 6.00 p.m. | | |
| Instructions: | Part – I, Question 1 Attempt any four q Figures to the right Answer to the Partanswer Booklet only | uestions from Part t indicate full mark – I and Part – II are | S. |
| | PA | RT-I | |
| 1. A) Rewrite the salternatives. | sentence after choosing | the correct answe | r from the given 7 |
| 1) Floras, Pla a) BGBM | ants and Peoples, and La | andscape Ecology c) GBIF | _ |
| more taxa | enetically informative | | nges are shared by two or ally uninformative |
| | investigates plants and appe Ecology | | eoples |
| a) Landso | t of can be me cape diversity c diversity | easured by the num b) Community o d) All of these | ber of links in the food web. liversity |
| 5) Which of the a) Literary | he following is not a type y works | e of copyright work b) Furniture | :? |

d) Musical works

SLR-RP - 61 6) World intellectual property organization was created in _____ year. a) 1967 b) 1960 d) 1980 c) 1957 7) _____ of the following is not specifically protected by intellectual property legislation. a) Trade marks b) Patents c) Copyright d) Trade secrets B) Answer the following. 7 1) Taxonomy 2) Patch density 3) Biodiversity informatics 4) GBIF 5) Industrial design 6) Farmer's right 7) WIPO. PART - II Answer any four of the following. 2. Explain in brief principles of Taxonomy and add a note on phylogeny in Biodiversity Informatics. 14 3. Explain goals of WIPO. Add a note on Protection of Intellectual property. 14 4. What is Biodiversity Informatics? Explain in detail national, regional and global biodiversity information systems and network. 14 5. Explain patenting of biological material with case study. 14 14 6. Answer **any two** of the following. 1) Explain Conservation of Biodiversity. 2) Write a note on advantages and disadvantages of PBR. 3) Explain technology transfer. 7. Short notes (any two). 14 1) Types of phylogenetic tree. 2) Trade mark. 3) Biodiversity data availability.



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| No. | |

M.Sc. (Bioinformatics) (Part – II) (Semester – IV) (CGPA Pattern) Examination, 2015

Paper – III: ADVANCED MOLECULAR BIOLOGY (New)

Day and Date: Tuesday, 21-4-2015 Total Marks: 70

Time: 3.00 p.m. to 6.00 p.m.

- *Instructions*: 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 Part II are to be written in same answer booklet only.

PART-I

| | | | | ı Altı – | - 1 | | | |
|--|----|--------------|---------------------------------------|-------------------|-----------------------|------------------|-------------------|---|
| 1. | A) | | ewrite the sentence a ternatives : | fter choosing the | e cc | orrect answer fr | om the given | 7 |
| | | 1) | PCR is used in a) RAPD | b) AFLP | c) | RFLP | d) Both a) and b) | |
| | | 2) | cDNA is prepared fro a) rRNA | | c) | tRNA | d) snRNA | |
| | | 3) | Western Blotting is use a) DNA | | c) | RNA | d) Both a) and b) | |
| | | 4) | Dialysis is used to rea a) Salt | | | Both a) and b) | d) None | |
| 5) Edman degradation is used fora) DNA sequencingc) RNA sequencing | | | | b) | Protein sequer All | ncing | | |
| 6) DNA fingerprint is used to detecta) Paternityc) Genetic test | | ed to detect | , | Crime samples | 3 | | | |
| | | 7) | size and shape. a) HPLC | omatography is | b) | lon exchange | proteins based on | |
| | | | c) Affinity | | d) | Gel filtration | | |

| | B) Answer the following: | 7 |
|----|---|----|
| | 1) Autoradiography | |
| | 2) Taq DNA polymerase | |
| | 3) Biochip | |
| | 4) ddNTPs | |
| | 5) Cy3 and Cy5 | |
| | 6) Nitrocellulose membrane | |
| | 7) Electro Blot. | |
| | PART-II | |
| Ar | nswer any four of the following : | |
| 2. | What is amplicon? Explain different types of PCR techniques. | 14 |
| 3. | Explain in detail the protein sequencing method. | 14 |
| 4. | Explain in detail southern blot and western blot techniques and its applications. | 14 |
| 5. | What is electrophoresis? Describe the separation of proteins using SDS PAGE. | 14 |
| 6. | Answer any two of the following: | 14 |
| | 1) Write a note on Plaque hybridization technique. | |
| | 2) Give a detailed account on DNA finger printing. | |
| | 3) Give a detailed account on DNA microarray. | |
| 7. | Short notes (any two): | 14 |
| | 1) Iso Electric Focusing | |
| | 2) RT-PCR | |
| | 3) cDNA library. | |
| | | |

SLR-RP-62

SLR-RP - 63



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M.Sc. (Part – II) (Semester – IV) Examination, 2015 BIOINFORMATICS New (CGPA Pattern)

| Nev Paper No. IV:En | w (CGPA Pa nerging Area | • | rmatics | |
|--|-----------------------------------|---|----------------------|--|
| Day and Date: Thursday, 23-4-201 Time: 3.00 p.m. to 6.00 p.m. | 5 | | Total Marks : 70 | |
| Instructions: 1) Part – I, Que 2) Attempt any 3) Figures to th 4) Answers to t same answe | four question ne right indicat | s from Part – II . e full marks. | be written in | |
| | PART-I | | | |
| A) Rewrite the sentences after alternatives. | choosing the | correct answer | from the given | |
| is a file format for holding information about the atoms, bonds, connectivity and co-ordinates of a molecule. | | | | |
| a) SDF b) I | MDL Mol c) | SMILES | d) WLN | |
| BeeBase is an online bid to | oinformatics da | atabase that dis | plays data related | |
| a) Apis florea | b) | Apis dorsata | | |
| c) Apis mellifera | d) | None of these | | |
| PCA refers to | | | | |
| a) Particle Component | Analysis b) | Particle Cataly | tic Activity | |
| c) Principal Componer | • , | None of these | | |
| 4)is a patho | gen database. | | | |
| a) VIPR b) l | EuPathDB c) | GOLD | d) Both a and b | |
| 5) is a file form and its associated data | _ | resent chemica | ll structure records | |
| a) CML b) s | SDF c) | XYZ | d) All of these | |

SLR-RP - 63_____ represents molecules by list of the atoms and of the bonds in the molecule. a) Atom look-up table b) Connection table c) Both a) and b) d) None of these 7) _____ first used the term nanotechnology. a) Eric Drexler b) Richard Feynman d) None fo these c) Sumio lijima B) Definitions: 7 1) WLN 2) Vaccine designing 3) Molecular modeling 4) Silver nanoparticle 5) EuPathDB 6) Top down 7) Gene Map. PART - II Answer any four of the following: 2. Define Immuno-informatics and explain the future of computational modeling and prediction system in clinical immunology. 14 3. Enlist the different databases for chemical structure representation and add a note on QSAR.S. 14 14 4. Give a detailed account on various genome databases. 5. Explain synthesis of nanoparticles by mechanical and biological method. 14 14 6. Answer any two of the following: 1) SMILE representation. 2) Give a detailed account on applications of nanoparticles in different area of science. 3) Give a detailed account on analysis of silver nanoparticles by UV and FTIR. 7. Short notes (any two): 14 1) Types of chemical databases and its uses 2) Applications of genome sequencing

3) FTIR.