

# ANDHRA PRADESH STATE COUNCIL OF HIGHER EDUCATION

# MINOR

# **Subject: Bioinformatics**

# w.e.f. AY 2023-24

# **COURSE STRUCTURE**

Year	Semester	Course	Title of the Course	No. of Hrs /Week	No. of Credits
	II	1	Cell Biology and Microbiology	3	3
Ι			Cell Biology and Microbiology Practical Course	2	1
	III IV	2	Basics of Computer application inBiology	3	3
			Basics of Computer application inBiology Practical Course	2	1
II		3	Genomics and Proteomics	3	3
			Genomics and Proteomics Practical Course	2	1
			Programming in C	3	3
			Programming in C Practical Course	2	1
	V	5	Structural Bioinformatics	3	3
III			Structural Bioinformatics Practical Course	2	1
		v 6	Programming in Pearl	3	3
			Programming in Pearl Practical Course	2	1

# COURSE 1: CELL BIOLOGY AND MICROBIOLOGY

	Theory	Credits: 3	3 hrs/week
Ι	Learning Objectives:		
	To acquaint students with basic		
2.	To complement students with k	nowledge on chromosome structure and packaging.	
3.	To enable them learn concepts	of Microorganisms and their growth aspects	
II.	. Learning Outcomes:		
	-	completion of the course will be able to:	
1.	Explain cells structure of proka	*	
		osis and structure and organization of chromosomes	
	•	ganisms, basic structure of bacteria and viruses	
		ial media and narrate microbial growth	
		methods of identification of bacteria	
II	I. Syllabus: (Total Teaching Ho	ours: 45)	
	nit 1:	· · · · · · · · · · · · · · · · · · ·	( <b>08h</b> )
-	ell Biology:		(0011)
	Cell as a basic unit of life.		
2.	Cell organization of prokaryotic	c and eukaryotic cells.	
	±	alization of cell:- Mitochondria, Chloroplast, Lysosomes	,
Go	olgibodies, Plasma membrane, (	Cytoskeleton, Cell wall and Nucleus.	
Uı	nit 2:		(12h)
1.0	Cell cycle, cell division - mitosi	s and meiosis.	
2.	Chromosome structure, gene, g	ene number, gene clusters and Pseudogene.	
3.	Polytene and lamp brush chrom	nosomes.	
4.	Packing of DNA, supercoiled I	DNA, nucleosome.	
5.	Inverted repeats, repetitive DN	A sequence, satellite DNA	
U	nit 3:		( <b>10h</b> )
Μ	licrobiology:		~ /
	Introduction to Microbiology a	nd microbial diversity	
	Classification of microorganism		
	Bacterial structure and reprodu-		
4.	Introduction to viruses- plant a	nd animal viruses- structure and life cycle	
Uı	nit 4:		( <b>8h</b> )
		ilization methods in biotechnology, Various sterilization	
me	ethods, Microbial contamination	control and Sterility testing.	

2. Microbial growth: The definition of growth, mathematical expression of growth, growth curve, measurement of growth and growth yield, synchronous growth, continuous culture.

# Unit 5:

1. Principles of microscopy – Light microscopy, Bright field and Electron microscopy (SEM and TEM).

2. Staining Techniques - Simple and Differential staining techniques.

3. Direct methods for measuring microbial growth: viable plate counts, membrane filtration. Indirect methods: Metabolic activity – measurements of DNA, Protein, Microscopic counts, electronic counters, most probable number

# COURSE 1: CELL BIOLOGY AND MICROBIOLOGY

Practical

Credits: 1

2 hrs/week

# IV. Skill Outcomes:

On successful completion of this practical course, student shall be able to:

- 1. Work with microscopes and observe plants cell to microorganisms
- 2. Isolate cell organelles from plant cells
- 3. Culture bacteria and fungi in culture media
- 4. Perform sterilization of media
- 5. Identify stages of mitosis and meiosis from biological samples

# V. Practical Syllabus: Hours 2 hours per week= 30 hours

- 1. Microbiology Good Laboratory Practices and Biosafety.
- 2. Preparation of culture media for cultivation of bacteria
- 3. Preparation of culture media for cultivation of fungi
- 4. Sterilization of medium using Autoclave
- 5. Sterilization of glassware using Hot Air Oven
- 6. Light compound microscope and its handling
- 7. Study of mitosis on onion root tips
- 8. Study of meiosis on onion buds
- 9. Isolation and separation of cell organelles from plant cell.
- 10. Study of growth curve of E.coli

# VI. References:

1. Pelczar, M.J., Chan, E.C.S. and Kreig, N.R. (1993). Microbiology. 5th Edition, Tata Mc Graw Hill Publishing Co., Ltd., New Delhi.

2. Dube, R.C. and Maheswari, D.K. (2000) General Microbiology. S Chand, New Delhi. Himalaya Publishing House, Mumbai.

3. Power, C.B. and Daginawala, H.F. (1986). General Microbiology Vol I & II

4. Prescott, M.J., Harley, J.P. and Klein, D.A. (2010). Microbiology. 5th Edition, WCB Mc Graw Hill, New York.

5. Reddy, S.M. and Reddy, S.R. (1998). Microbiology Practical Manual, 3 rd Edition, Sri Padmavathi Publications, Hyderabad.

6. Singh, R.P. (2007). General Microbiology. Kalyani Publishers,

7. Stanier, R.Y., Adelberg, E.A. and Ingram, J.L. (1991). General Microbiology, 5th Ed.,

Prentice New Delhi.Hall of India Pvt. Ltd., New Delhi.

8. Microbiology Edited by Prescott

9. Jaya Babu (2006). Practical Manual on Microbial Metabolisms and General Microbiology.

Kalyani Publishers, New Delhi.

10. Gopal Reddy et al., Laboratory Experiments in Microbiology

# VII. Co-Curricular Activities

# Suggested Co-Curricular Activities

1. Training of students by related aspects using pure cultres

2.Assignments on handling microscopic techniques with safety and precautions)

3.Seminars, Group discussions, Quiz, Debates etc

4. Preparation of videos on tools and techniques on mitosis and meiosis

5. Collection of material/figures/photos related to stages of cell division, microbial growth curve and staining methods.

6. Visits to advanced laboratories to get exposure to SEM and TEM

7.Invited lectures and presentations on related topics by experts.

#### SEMESTER-III

# **COURSE 2: BASICS OF COMPUTER APPLICATION IN BIOLOGY**

Theory	Credits: 3	3 hrs/week
I Learning	Objectives:	
-	c knowledge about computers and internet.	
	computational methods to utilize expression data's of cellular biology.	
3. To study of the	e inherent structure of biological information.	
4. To analyze the	gene and protein sequences to reveal protein evolution.	
II. Learning	Outcomes:	
e	after successful completion of the course will be able to:	
1. Identify compo	onents of computer	
2. Connect to inte	ernet and email or download information	
3. Understand ba	sic history of bioinformatics	
4. Illustrate about	t biological databses	
5. Retrieve PDB	formats of proteins from respective databases	
III. Syllabus: (T	otal Teaching Hours: 45)	
Unit 1:		(10h)
Computers –		
	of Computers – Areas of computer applications- I-P-O Cycle.	
2. Components o Software -Operat	f Computers – Memory and control units-Input devices and output device ing Systems.	es- Hardware and
Unit 2:		( <b>10h</b> )
	y of Internet-Uses of internet. Connection to Internet - Getting connection	
	Service providers-E-mail and Voice Mail, Creating E-mail Address.	
Unit 3:		( <b>10h</b> )
	<b>bioinformatics</b> – history and its development – Scope and applications of	. ,

Unit 4:

**Biological database** – NCBI-GenBank, EMBL, DDBJ. Sequence Alignment-Pairwise (BLAST and FASTA) and Multiple sequence alignment (ClustalW).

(**8h**)

(7h)

Unit 5:

Structure of Protein, Classification –PDB, Swiss-PROT, SCOP, CATH. Protein visualization tools-RASMOL, Swiss PDB viewer.

#### **SEMESTER-III**

# **COURSE 2: BASICS OF COMPUTER APPLICATION IN BIOLOGY**

Practical	Cradits: 1	2 hm/maal
Practical	Credits: 1	2 nrs/week

# IV. Skill Outcomes:

On successful completion of this practical course, student shall be able to:

- 1. Login computer and perform booting
- 2. Learn about operating systems
- 3. Retrieve data from databases
- 4. Perform emailing of retrieved data from databases
- 5. work with basics of pair wise alignment and multiple sequence alignment

# V. Practical Syllabus: Hours 2 hours per week= 30 hours

- 1. Login operations of a computer and its characteristics
- 2. Hardware, software and booting of computers
- 3. Operating systems UNIX and LINUX
- 4. Emailing
- 5. Exposure to Databases- NCBI and DDBJ
- 6. Retrieve data from Databases
- 7. Pairwise alignment
- 8. Multiple sequence alignment

# VI. References:

1. Computer basic knowledge; hardware, connection, cables, typing, Windows98/XP, Internet browsers, search engines.

2. LAN connections, setting up the IP address, network security

- 3. Internet surfing and searching information, downloading and installing software
- 4. T.A.Attwood and D.J.Parry-smith, 2001, Introduction of Bioinformatics.

5. A.D.Baxevaris,1998, Bioinformatics: A practical guide to the analysis of Genes and proteins,(Edited) B.F.Publication.

6. David W, 2005, Bio-informatics; sequence and Genome Analysis, 2<sup>nd</sup> Edition By Mount CBS publishers

# VII. Co-Curricular Activities

# **Suggested Co-Curricular Activities**

- 1. Training of students on working with computers
- 2. Help them know about booting and also operating systems in labs
- 3. Assignments on data retrieval and alignments studies
- 4. Groups discussion, quizzes and video making on basic concepts
- 5. Invited lectures on the course

# **COURSE 3: GENOMICS AND PROTEOMICS**

Theory Credits: 3	
I Learning Objectives:	
1. To introduce students to the concepts of Genomics, Human genome project and databas	es
2. To enhance knowledge on gene structure and identification	
3. to enable them learn advanced concepts of proteomics	

# II. Learning Outcomes:

Students after successful completion of the course will be able to:

1. Imbibe concepts of Nucleotide sequence databases

2. Understand identification of gene and its functional sites

3. Illustrate concepts of gene expression, microarrays and concepts of Proteomics

4. Explain concepts of functional proteomics and phylogenetic analysis

5. Illustrate applications of Bioinformatics in various fields

# III. Syllabus: (Total Teaching Hours: 45)

Unit 1:

1. Genomics: Nucleotide sequence Databases, its Analysis and Identification.

2. Goals of the Human Genome Project, cloning vectors, concept of maps, physical maps, shotgun libraries, DNA polymorphism, nucleotides, DNA sequences.

3. **Sequence databases:** GeneBank, EMBL Nucleotide sequence databank, DNA Data Bank of Japan (DDBJ), database formats.

# **Unit 2:**

1. Recombinant DNA technology, restriction enzymes, resource for restriction enzyme (REBASE), similarity search. Polymerase chain reaction, primer selection for PCR, BLASTn,application of BioEdit.

2. Genome information and special features, coding sequences (CDS), untranslated regions(UTR's), cDNA library, expressed sequence tags (EST).

3. Approach to gene identification; masking repetitive DNA, database search, codon-bias detection, detecting functional sites in the DNA.

4. Internet resources for gene identification, detection of functional sites, gene expression.

# Unit 3:

# Gene expression, DNA microarray and Proteomics:

1. Gene expression: Introduction, Basic steps for gene expression.

2. **Microarray:** Concept of microarrays; spotted arrays, oligonucleotide arrays, designing the experiment, Two-color microarray experiments.

3. **Proteomics**:-Protein sequence information, composition and properties, physico-chemical properties based on sequence, sequence comparison,

4. Primary databases, Secondary databases.

5. Pair-wise sequence alignment, gaps, gap-penalties, scoring matrices, PAM250,BLOSUM62, local and global sequence alignment, multiple sequence alignment, useful programs, ClustalW, BLASTp.

#### (10h)

(10h)

(10h)

3 hrs/week

#### Unit 4:

1. **Proteomics classification;** Tools and techniques in proteomics; 2-D gel electrophoresis, gelfiltration, PAGE, isoelectric focusing, affinity chromatography, HPLC, ICAT, fixing and spot visualization, Mass spectroscopy for protein analysis, MALDI-TOF, Electrospray ionization(EST),Tandem mass spectroscopy (MS/MS) analysis; tryptic digestion and peptidefingerprinting (PMF).

2. Protein Micro array in protein expression, profiling and diagnostics, drug target discovery. Database searching, 3-dimensional structure determination by X-ray and NMR.

3. **Phylogenetic analysis:** Evolution, elements of phylogeny, methods of phylogenetic analysis, Phylogenetic tree of life, comparison of genetic sequence of organisms, phylogenetic analysis tools-Phylip, Clustal W.

# Unit 5:

(5h)

# Applications of Bioinformatics in various fields:

Environment, biotechnology, molecular biology, neurobiology, agriculture, drug designing, biomedical genome medicines, medical microbiology.

# **COURSE 3: GENOMICS AND PROTEOMICS**

Practical	Credits: 1	2 hrs/week

# IV. Skill Outcomes:

On successful completion of this practical course, student shall be able to:

- 1. Retrieve biological information from various databases
- 2. Retrieve literature from PUBMED and related databases
- 3. Perform Pair-wise and multiple sequence alignments and compare biological sequences
- 4. Analyze 3-dimensional protein structure
- 5.Performphylogenetic analysis

# V. Practical Syllabus: Hours 2 hours per week= 30 hours

- 1. Introduction of National Center for Biotechnology Information (NCBI).
- 2. Introduction of biological search engine- Entrez

3. Introduction to literature database at NCBI and querying the PUBMED centraldatabase using the ENTREZ search engine

- 4. Analysis of 3D structure of protein using RasMol through command line.
- 5. Analysis of 3D structure of protein and nucleic acid using Cn3D.
- 6. Pair-wise sequence alignment by using ClustalW.
- 7. Multiple sequence alignment by using ClustalW.
- 8. Introduction of BioEdit. Effect of insertion INDEL from given amino acid using
- 9. Pairwise and Multiple sequence alignment using BioEdit.
- 10. Phylogenetic analysis using web tool.

# VI. References:

1. Daniel, 2006, Biostatistics, Eighth Edition. John Wisley and sons.

2. Durbin, Eddy, Krogh, Mithison, Biological sequence analysis.

3. T.A.Attwood and D.J.Parry-smith, 2001, Introduction of Bioinformatics.

4. A.D.Baxevaris,1998, Bioinformatics: A practical guide to the analysis of Genes and proteins,(Edited) B.F.Publication.

5. David W, 2005, Bio-informatics; sequence and Genome Analysis, 2<sup>nd</sup> Edition By Mount CBS publishers

# VII. Co-Curricular Activities

# a) Suggested Co-Curricular Activities

1. Training of students in accessing data bases and retrive information

2. Assignments on data retrieval and structure prediction

3.Seminars, Group discussions, Quiz and projects on aspects of genomics and proteomics

4. Preparation of videos on tools and their usage

5. Visits to facilities and organizations working on advanced concepts of Genomics and proteomics

7.Invited lectures and presentations on related topics by field experts.

# COURSE 4: PROGRAMMING IN C

Theory	Cradita: 2	2 has/mast
2. To gain the know	sic concepts of Programming. wledge of C- programming language. cs which will help them to create programs and applications in C	<u>3 hrs/week</u> C using functions,
<ol> <li>Explain about C +</li> <li>Understand the c problems.</li> <li>Describe the con</li> <li>Explains about the</li> </ol>	ter successful completion of the course will be able to: and their applications in bioinformatics analysis. concepts of Arrays, Strings and Structures which helps them to s	solve the real time
Unit 1: Introduction to C 1. History of C 2. Characteristics of 3. Program Structure		(10h) ion and Execution.
<ol> <li>Arithmetic Unary</li> <li>If Statement, Nes</li> <li>Ladder switch, S</li> </ol>	<b>hing &amp; Looping Statements</b> y Assignment Relational & Logical Conditional sted if, Statement else-if. Statement Looping. op while loop do-while loop Jump Statements.	(10h)
<ol> <li>2. Single &amp; Multi-E</li> <li>3. Types of Functio</li> </ol>	Rules & Restrictions. Dimensional Arrays ons, Functions and Arrays Function. ope of Variables Built-in Functions	(10h)
Unit 4: 1. String Functions,	, String Manipulation.	(10h)

2.Pointer Concepts, Pointers and Functions Pointers and Arrays.Array of Pointers, Static Initialization, Pointers and Structures, Illegal indirection.

3. Defining New Data types, Unions Type Casting Enumerated, Data types Static Variables, Type

Definition.

Unit 5:

# **Pointers**

- Null pointers, pointers and settings
   Pointer and two dimensional arrays
- Function philosophy
   Function basics, Function prototype

# COURSE 4: PROGRAMMING IN C

Practical	

Credits: 1

2 hrs/week

# IV. Skill Outcomes:

On successful completion of this practical course, student shall be able to:

- 1. Understand how to write the programs by using C-language.
- 2. Expertise in solving the problems by application of C- programming.
- 3. Perform Multithreading
- 4. Write program for stacking
- 5. Work with Applet

# V. Practical Syllabus: Hours 2 hours per week= 30 hours

- 1. Find the prime numbers between 1 to 50
- 2. Write a program which uses switches and break case statements.
- 3. Find out length of given string
- 4. Write a program of insertion sort.
- 5. Write a program which implements stack operation.
- 6. Multithreading using get property.
- 7. Multithreading using sleep property.
- 8. Write a program which implements mouse listener and mouse motion listener.
- 9. Creating a frame window in an applet.
- 10. Draw line, rectangle, oval in an applet.

# VI. References:

- 1. Let us C by Yashwant K., 4th Ed.
- 2. The C programming language by Ritchie, D.M., 2nd Ed.
- 3. C: The Complete Reference is written by Herbert Schildt.
- 4. Programming in ANSI C is written by E Balagurusamy.

# VII. Co-Curricular Activities

# a) Suggested Co-Curricular Activities

- 1. Training of students on C programming
- 2. Assignments on Topics related in course
- 3.Seminars, Group discussions, Quiz and projects on aspects C programming
- 4. Preparation of videos on tools and their usage

5.Invited lectures and presentations on related topics.

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#### **COURSE 5: STRUCTURAL BIOINFORMATICS**

#### Credits: 3 Theory 3 hrs/week Ι **Learning Objectives:** 1. To introduce students to structural bioinformatics and structural analysis 2. To enhance knowledge on structural prediction 3. To educate them on the topic Macro-molecular interaction II. **Learning Outcomes:** Students after successful completion of the course will be able to: 1. Understand the structural basis for biological phenomena 2. Explain conformational analysis of proteins using computational methods 3. Describe features about forces that determine the conformational analysis of nucleic acids 4. Learn aspects of protein structure prediction 5. Examine the aspects of genome sequencing **III. Syllabus:** (Total Teaching Hours: 45) Unit 1: (10h)**Introduction:** Overview of structural bioinformatics – understanding structural basis for biological phenomena– challenges in structural bioinformatics - integration of structural data with other data. **Protein structures** Conformational Analysis of proteins – Forces that determine protein structure – polypeptide chain geometries - Ramachandran Map - potential energy calculations - observed values for rotation angles. (10h) **Unit 2: Structural analysis** Conformational analysis of nucleic acids – general characteristics of nucleic acid structure – geometries, glycosidic bond – rotational isomers and ribose puckering - forces stabilizing ordered forms – base pairing–

base stacking. Unit 3:

# **Structural Prediction**

Structure Prediction Methods – Homology Modeling – Fold Recognition Methods – ab initio methods – Rosetta – CASP – prediction of secondary structure – Chou-Fasman, Garnier, Osguthorpe-Robson (GOR) methods (qualitative aspects only) – transmembrane structure prediction – solvent accessibility calculations and prediction

# Unit 4:

# **Macro-molecular interactions**

 $Interactomes - macromolecular\ interactions - protein-protein\ interactions - protein-DNA\ interactions - protein-ligand\ interactions - interactions\ databases - BIND,\ ProNIT - Docking - principles\ and\ methods$ 

# Unit 5:

#### **Current contours**

Genome sequencing - Proteomics - Phylogeny - Gene expression - Protein-protein interaction network

(10h)

#### (10h)

(**5h**)

# **COURSE 5: STRUCTURAL BIOINFORMATICS**

Credits: 1		

2 hrs/week

# IV. Skill Outcomes:

Practical

On successful completion of this practical course, student shall be able to:

- 1. Access free online repositories of structural data
- 2. Retrieve right PDB structures and visualize them using PyMOL
- 3. Identify structural and functional domains of proteins
- 4. Predict structure of transmembrane proteins
- 5. Understand about protein-protein interaction networks

# V. Practical Syllabus: Hours 2 hours per week= 30 hours

1. Access public repositories of structural data- Protein Data Bank (PDB) and Electron Microscopy Data Bank (EMDB)

- 2. Access repositories and retrive information from Protein Data Bank in Europe (PDBe) and PDBe-KB
- 3. Finding the right structures UniProt and PDB
- 4. Viewing the structure PyMOL
- 5. Domain identification in proteins using HMMER, Inter Pro and CATH
- 6. Access MemBrain for trans membrane protein prediction
- 7. DNAproDB for DNA protein interactions
- 8. Access STRING for protein-protein interaction networks

# VI. References:

- 1. C.R.Cantor & P.R.Schimmel, Biophysical Chemistry Part I, W.H. Freeman & Co., San Fransisco, 1980.
- 2. C. Branden and J. Tooze, Introduction to Protein Structure, Garland Publishing Inc., NewYork, 1999.
- 3. P.E. Bourne and H. Weissig (Eds.) Structural Bioinformatics, John-Wiley and Sons, 2003
- 4. Molecular Modeling and Simulation, Tamar Schlick, 2002
- 5. Structural Bioinformatics, 2nd Edition, Jenny Gu and Philip Bourne, 2009
- 6. Protein Structure Prediction: a practical; approach, M. Sternberg, 1996
- 7. Textbook of Structural Biology, Liljas et all, 2010
- 8. Molecular Biophysics, Michael Daune, 1999
- 9. Introduction to Proteins: Structure, Function, and Motion, Kessel et al, 2010
- 10. Understanding Bioinformatics, M. Zvelebil, J. Baum, 2007

# VII. Co-Curricular Activities

# a) Suggested Co-Curricular Activities

- 1. Training of students on aspects of course using free online tools
- 2. Assignments on protein structure prediction and importantly domain identification
- 3.Seminars, Group discussions, Quiz, online tests etc.
- 4. Preparation of videos on tools as assignments for better understanding of students as peer group teaching
- 5.Collection of material related to every topic and share in google classroom
- 6. Visits to research organizations or firms on the aspects of topic
- 7. Invite guest lectures and presentations on related topics of structural Bioinformatics.

#### **COURSE 6: PROGRAMMING IN PEARL**

Theory

#### Credits: 3

3 hrs/week

# I Learning Objectives:

- 1. To acquaint with on basics in perl and more on usage of scalar, arrays and hashes.
- 2. To gain knowledge of regular expressions concepts in perl and its major role in bioinformatics.
- 3. To understand the significance of perl modules in the advance programming skills.

# II. Learning Outcomes:

Students after successful completion of the course will be able to:

- 1. Understand basic concepts of biodiversity and its distribution.
- 2. Gain the knowledge of patterns of the biodiversity.
- 3. Appreciate the Biodiversity present in the India.
- 4. Acquire the knowledge on terminology of Biodiversity.
- 5. understand the importance information technology to identify the biodiversity in global way.

# III. Syllabus: (Total Teaching Hours: 45)

# Unit 1:

# **Introduction to Perl**

1. The Organization of DNA and Organization of Proteins,

2. In Silico, Limits to ComputationGetting started with perl:- A Low and Long Learning Curve.

3.Perl's Benefits, Installing Perl on Your Computer, How to Run Perl Programs, Text Editors and Finding Help.

4. The art of programming: Individual Approaches to programming, Edit-Run-Revise (and Save), An Environment of Programs, Programming Strategies.

# **Unit 2:**

#### Programming

1. The Programming Process, sequences and strings: Representing Sequence Data,

- 2. Transcription: DNA to RNA, Using the Perl Documentation
- 3. A Program toStore a DNA Sequence, Concatenating DNA Fragment.
- 4. Calculating the Reverse Complement in Perl, Proteins, Files, and Arrays, Reading Proteins in Files.

# Unit 3:

# Arrays, Motifs and Loops

1. Arrays Scalar and List Context.

2. Motifs and Loops:-Flow Control, Code Layout, Finding Motifs, Counting Nucleotides, Exploding Strings into Arrays,

3. Operating on Strings Writing to Files.

# Unit 4:

# **Subroutines and Bugs**

1. Subroutines, Scoping and Subroutines

2. Command-Line Arguments and Arrays

(10h)

(8h)

# (10h)

(10h)

3. Passing Data to Subroutines, Modules and Libraries of Subroutines,

4. Fixing Bugs In YourCode

# Unit 5:

# **Mutations and Randomization**

1. Random Number Generators, A Program Using Randomization,

2. .A Program to Simulate DNA Mutation, Generating Random DNA, Analyzing DNA.

3. The genetic code:-Hashes, Data Structures and Algorithms for Biology,

4. The Genetic Code, Translating DNA into Proteins, Reading DNA from Files in FASTA Format, Reading Frames

(7h)

# **COURSE 6: PROGRAMMING IN PEARL**

Practical

Credits: 1

2 hrs/week

# IV. Skill Outcomes:

On successful completion of this practical course, student shall be able to:

- 1. Understand and Expertise in programming in Perl.
- 2. Write the programs for Biological data with the application of Perl programming
- 3. Do MAP construction
- 4. Work with different software tools
- 5. Create databases

# V. Practical Syllabus: Hours 2 hours per week= 30 hours

- 1. Installing Perl on your PC.
- 2. Create Perl script.
- 3. Write a program to store protein sequence.
- 4. Write a program to store DNA sequence.
- 5. Write a program to store RNA sequence.
- 6. Use Perl to concatenation of DNA
- 7. Use Perl to concatenation of protein sequence.
- 8. Perl script for to simulate DNA mutation

# VI. References:

1. Beginning Perl for Bioinformatics by James Tisdall, O-Reilly publication.

- 2. Perl: Complete Reference Perl by Martin C. Brown, McGrawHill publication.
- 3. Mastering Perl for Bioinformatics by James D. Tisdall, O-Reilly Publication.
- 4. Teach Yourself Perl 5 in 21 days by David Till, Sams publishing.

5. Mastering Algorithms with Perl by Jon Orwant, Jarkko Hietaniemi and John Macdonald, O-Reilly Publication

# VII. Co-Curricular Activities

# **Suggested Co-Curricular Activities**

- 1. Invited talks by subject experts of Perl
- 2. Give assignments on programming to store biological sequences
- 3. Quiz, Group Discussions etc.,
- 4. Making of videos and material creating and sharing
- 5. Guest lectures and field visits.