# 

#### (Abstract)

M.Sc Biostatistics in the Dept. of Statistical Sciences, Mangattuparamba Campus - Scheme (Credit distribution for courses) & Syllabus of the First Semester -Approved & Implemented w.e.f 2023 admission - Orders issued

#### ACADEMIC C SECTION

#### ACAD/ACAD C3/365/2023

Dated: 21.11.2023

Read:-1. GO (Ms) No 528/2022/HEDN dated 22/10/2022

2. Letter No Acad C3/20143/2022 dated 26/11/2022

3. Email dated 07/01/2023 from the Head, Dept of Statistical Sciences,

Mangattuparamba Campus.

4. U.O of even number dated 07/02/2023

5. Minutes of the online meeting of the Expert Committee dated 21/03/2023.

6. UO of even number dated 22/06/2023

7. UO No ACAD C/ACAD C3/22373/2019 dated 12/09/2023

8. Email dated 05/11/2023 from the Head, Dept of Statistical Sciences,

Mangattuparamba Campus.

9. Minutes of the meeting of the Department Council dated 18/09/2023

#### ORDER

1. As per paper read (1) above, Govt. of Kerala granted Administrative Sanction for starting Project Mode Programme *M.Sc Biostatistics with Specialization in Epidemiology and SAS Programming* in the Dept. of Statistical Sciences, Mangattuparamba campus, Kannur University during the Academic Year 2022-'23.

2. As per paper read (2) above, HoD, Dept. of Statistical Sciences was requested to prepare and submit the draft Scheme & Syllabus for the aforementioned Programme along with a panel of fivemember Experts to constitute a committee to scrutinize the syllabus.

3. As per paper read (3) above, HoD, Dept. of Statistical Sciences submitted the Scheme and Syllabus for the Programme *MSc Biostatistics with Specialization in Epidemiology and SAS Programming* along with a panel of ten experts to scrutinize the syllabus.

4. As per paper read (4) above, a ten member Expert Committee was constituted, with the Head, Dept of Statistical Sciences as the Coordinator, to scrutinize the Scheme & Syllabus of the aforesaid programme. Head, Dept. of Statistical Sciences was authorised to submit the final Scheme & Syllabus of the Programme after incorporating the corrections/modifications, if any, suggested by the Expert Committee.

5. As per paper read (5) above, the Expert Committee suggested to change the nomenclature of the Programme as *MSc Biostatistics* instead of *MSc Biostatistics with Specialization in Epidemiology and SAS Programming*.

6. Accordingly, as per paper read (6) above, nomenclature of the Programme is changed from MSc Biostatistics with Specialization in Epidemiology and SAS Programming to MSc

#### Biostatistics.

7. As per paper read (7) above, Revised Regulations for PG Programmes under CBCSS in the University Teaching Depts./ Schools was implemented w.e.f 2023 admission.

8. As per paper read (8) above, the Head, Dept. of Statistical Sciences submitted the Scheme (Distribution of credits for courses) & Syllabus of the First Semester MSc Biostatistics Programme w.e.f 2023 admission, incorporating the suggestions of the Expert Committee and also in accordance with the revised Regulations implemented in the Teaching Departments / Schools of Kannur University w.e.f 2023 admissions. Department Council in its meeting held on 18/09/2023 vide paper read (9) above approved the aforesaid syllabus.

9. The Vice Chancellor, after considering the matter in detail and in exercise of the powers of the Academic Council conferred under section 11(1), Chapter III of Kannur University Act 1996, approved the Scheme (distribution of credits) & Syllabus of the First Semester MSc Biostatistics Programme subject to reporting to the Academic Council and accorded sanction to implement the syllabus of M.Sc Biostatistics Programme in the Department of Statistical Sciences, Mangattuparamba Campus w.e.f 2023 admission.

10.Scheme (Distribution of Credits) & Syllabus of the First Semester MSc Biostatistics Programme implemented in the Department of Statistical Sciences, Mangattuparamba Campus with effect from 2023 admission, is appended and uploaded in the University website (www.kannuruniversity.ac.in)

11. Orders are issued accordingly.

Sd/-

Narayanadas K DEPUTY REGISTRAR (ACAD) For REGISTRAR

To:

- 1.Head, Dept of Statistical Sciences, Mangattuparamba Campus 2.Convener, Curriculum Committee.
- Copy To: 1. To Exam Branch (through PA to CE)
  - 2. PS to VC/ PA to PVC/ PA to R/PA to CE
  - 3. EP IV / EXCI/SWC 4 . DR/ AR I /AR II (Acad)
  - 5. To Webmanager (to publish in the website) 6. Computer Programmer
  - 7.SF/DF/FC



Forwarded / By Order

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#### (Abstract)

M. Sc Biostatistics Programme in the Dept of Statistical Sciences, Mangattuparamba Campus of Kannur University - Scheme & Syllabus of II, III & IV Semesters - Approved - Implemented w. e. f 2023 admission - Orders Issued.

#### ACADEMIC C SECTION

ACAD/ACAD C3/365/2023

Dated: 31.05.2024

Read:-1. U.O.s No ACAD C/ACAD C3/22373/2019 dated 12/09/2023, 08/11/2023 &

16/02/2023

2. U.O of even number dated 21/11/2023

3. Circulars of No ACAD C/ACAD C3/22373/2019 dated 01/02/2024 & 12/03/2024

4. Email dated 23/04/2024 from the Head , Dept of Statistical Sciences,

Mangattuparamba Campus

5. Minutes of the meeting of the Department Council dated 22/04/2024

#### ORDER

1. The revised Regulations for PG Programmes under CBCSS in the University Teaching Depts/ Schools were implemented w. e. f 2023 admissions vide paper read (1) above.

2. As per paper read (2) above, Revised Scheme (Credit distribution of courses) & Syllabus (I Semester Only) of M. Sc Biostatistics Programme was approved and implemented in the Dept of Statistical Sciences, Mangattuparamba Campus w. e. f 2023 admission.

3. As per paper read (3) above, Heads of all Teaching Depts who had not submitted the syllabi in full, were requested to submit the syllabi of the remaining semesters in accordance with the approved Regulations and along with a copy of the Department Council Minutes.

4. As per paper read (4) above, the Head, Dept of Statistical Sciences submitted the Scheme & Syllabus (II, III & IV Semesters) of M. Sc Biostatistics Programme to be implemented in the University Teaching Dept w.e.f 2023 admission.

5. Dept Council vide paper read (5) above, recommended the aforementioned Syllabus of M. Sc Biostatistics Programme to be implemented in the Dept of Statistical Sciences, Mangattuparamba Campus w.e.f 2023 admission.

6. The Vice Chancellor, after considering the matter in detail and in exercise of the powers of the Academic Council conferred under Section 11(1), Chapter III of Kannur University Act 1996, approved the Scheme & Syllabus (II, III & IV Semesters) of M. Sc Biostatistics Programme and accorded sanction to implement the same in the Dept of Statistical Sciences, Mangattuparamba Campus w. e. f 2023 admission, subject to report to the Academic Council.

7. The Scheme & Syllabus (Ist, II nd, III rd & IV th Semesters) of M. Sc Biostatistics Programme

under CBCSS, implemented in the Dept of Statistical sciences, Mangattuparamba Campus w. e. f 2023 admission, is appended and uploaded in the University website (www.kannuruniversity.ac.in).

8. Orders are issued accordingly.

Sd/-

# Narayanadas K DEPUTY REGISTRAR (ACAD) For REGISTRAR

To: 1. Head, Dept of Statistical Sciences, Mangattuparamba Campus

2. Convenor, Curriculum Committee

#### Copy To: 1. PS to VC/ PA to R

- 2. PA to CE (to circulate among the sections of the Examination Branch concerned)
- 3. EP IV/ EX C1
- 4. Computer Programmer
- 5. Webmanager (to publish in the website)
- 6. SF/DF/FC



Forwarded / By Order



# **KANNUR UNIVERSITY**

# **M.Sc. BIOSTATISTICS**

# **SCHEME & SYLLABUS**

(Under Choice Based Credit & Semester System)

2023 Admission Onwards NURU

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# DEPARTMENT OF STATISTICAL SCIENCES

Mangattuparamba Campus

Scheme and Syllabus of M Sc. Biostatistics- 2023 Admission onwards- Kannur University

# **KANNUR UNIVERSITY**

### Post Graduate Programme in Biostatistics

M.Sc. Biostatistics programme is a two-year programme divided into four semesters. A student is required to complete at least 80 credits for the completion of the programme and the award of degree.

#### **DURATION**: 2 Years (4 semesters)

**INTAKE**: 15.

#### **OBJECTIVES OF THE PROGRAMME**

- 1. Gain sound knowledge in theoretical and practical aspects of Biostatistics.
- 2. Acquire the working knowledge of various statistical software and programming languages.
- 3. Acquire skills and competencies in Biostatistical computing methods and develop algorithms and computer programmes for analyzing complex datasets.
- 4. Communicate effectively complex statistical ideas to people working in diverse spheres of academics and organizational setups.
- 5. Handle and analyze large databases related to various biomedical research and make meaningful interpretations of the results.
- 6. Get wide range of job opportunities in industry as well as in government sector.
- 7. Make unique contribution for the development of discipline by addressing complex and challenging problems in emerging areas of the discipline.
- 8. Imbibe effective scientific and/or technical communication in both oral and writing.
- 9. Continue to acquire relevant knowledge and skills appropriate to professional activities and demonstrate highest standards of ethical issues in Biostatistical sciences.

#### **ELIGIBILITIES:**

The selection procedure will be based on an entrance examination by the University. The eligibility criteria for appearing entrance examination is any of the following degree with overall 50% marks:

- 1. B.Sc. Statistics/Biostatistics as core course.
- 2. B.A./B.Sc. Mathematics
- 3. B.Sc. Computer Science with Statistics/Mathematics as complementary course
- 4. B. Tech/B.E degree.
- 5. B.Sc. with Mathematics and Statistics as core courses.

#### **ADMISSION:**

- The selection of the candidate is mainly based on the marks secured in the Degree Course/Admission test.
- The admission test will cover statistics and mathematics at the undergraduate level.

#### **Relaxation & Weightage**

Relaxation and weightage will be as per Kannur University rule.

#### **COURSE DETAILS:**

A student must register for the required number of courses at the beginning of each semester.

No students shall register for more than 28 credits and less than 16 credits per semester.

A total of 80 credits shall be the minimum for successful completion of the course in which minimum of 50% of credits have to be earned from Discipline Specific Courses including dissertation for any programme. Those who secure only minimum credit for core/ elective subjects has to supplement the deficiency for obtaining the minimum total credits required for successful completion of the program from the other divisions.

#### **EVALUATION:**

The faculty member who teaches the course shall do evaluation of the students for each course on the basis of Continuous Evaluation and End Semester Examination shall be evaluated by External Examiners. The proportion of the distribution of marks among the continuous evaluation and end semester examination shall be **40:60**.

Continuous Evaluation includes assignments, seminars, written examination and viva voce for each course. Weightage to the components of continuous evaluation shall be given for all theory papers of the course as follows:

Components of CE	Minimum Number	Weightage	Grade Points	Practical Weightage	Grade Points
Test paper	2	40	16	80	-
Assignments	SPY 1	20	08	—	
Seminar presentation, Viva Voce, Discussion, Debate etc.		40	16		
Record				20	-

Test Paper: For each course there shall be at least two class tests during a semester.

Assignments: Each student shall be required to do one assignment for each course.

**Seminar:** Students are required to present a seminar on a selected topic in each paper. The evaluation of the seminar shall be done by the concerned teacher handling the course.

**Viva Voce** – End semester theory Viva Voce examination will be conducted for each paper before the commencement of public examination.

Attendance: Minimum attendance required for each paper shall be 75% of the total number of classes conducted for that semester. Those who secured the minimum requirement of attendance only be allowed to register/appear for End Semester Examination.

Condonation of attendance to a maximum of 10 days in a semester subject to a maximum of two times during the whole period of the PG program may be granted by the university as per university rules.

#### **Conduct of Examination:**

The Vice Chancellor will approve the panel of examiners submitted by the Head of the Department. All the teachers of the Department will be the members of the Board of examiners with Head of the Department as the Chairperson, there shall be an external examiner.

#### **Research Project:**

The students have to complete a research project during IV Semester in collaboration with any of the authorized research institutions located within or outside the state or within their own Department.

# KANNUR UNIVERSITY

#### **DEPARTMENT OF STATISTICAL SCIENCES**

### VISION

Motivated by optimism and responsibility, the vision is to develop an exemplary centre for studies, practice and research in Statistics which will be beneficial to the stakeholders and

the society.

#### **MISSION**

To develop an excellent centre of quality teaching and research in Statistics To develop an international centre for advanced statistical computing and data analysis.

#### **PROGRAMME OUTCOMES**

- PO 1 : Critical Thinking: Take informed actions after identifying the assumptions that frame our thinking and actions, checking out the degree to which these assumptions are accurate and valid, and looking at our ideas and decisions (intellectual, organizational, and personal) from different perspectives.
- **PO2 : Problem Solving:** Identify, formulate, conduct investigations, and find solutions to problems based on in-depth knowledge of relevant domains.
- **PO 3** : Communication: Speak, read, write and listen clearly in person and through electronic media in English/language of the discipline, and make meaning of the world by connecting people, ideas, books, media and technology.
- **PO 4 : Responsible Citizenship:** Demonstrate empathetic social concern, and the ability to act with an informed awareness of issues.
- **PO 5** : Ethics: Recognize different value systems including your own, understand the moral dimensions of your decisions, and accept responsibility for them.
- **PO 6 : Self-directed and Life-long Learning:** Acquire the ability to engage in independent and life-long learning in the broadest context sociotechnological changes.

#### PROGRAMME SPECIFIC OUTCOME

- **PSO 1:** Expertise in the field of biostatistical theory and its practical applications.
- **PSO 2:** Expertise to take up responsibilities as efficient Biostatisticians/Statistical Officers/Research Officers/ Statistical Analytics.
- **PSO 3:** Expertise on techniques of biostatistics and in the field of data analysis.
- **PSO 4:** Make Awareness on recent trends in biostatistical theory and applications.
- **PSO 5:** Utilize statistical methods and tools to analyze data sets, draw meaningful conclusions, and make informed decisions based on biostatistical inferences.
- **PSO6:** Demonstrate proficiency in using statistical software such as R and SPSS, to perform statistical computations, visualize data, and facilitate biostatistical analysis.

#### **COURSE OUTCOME**

- CO1 : Demonstrate an in-depth understanding of Biostatistical concepts, including advanced clinical trials, statistical epidemiology, demography, sampling and design, statistical inference, regression analysis, probability and distribution theory.
- **CO 2** : Apply biostatistical techniques to analyze real life data using statistical packages such as SPSS, SAS and free software R and Python.
- **CO3**: Formulate suitable models for pharmaceutical research and drug development.
- **CO 4** : Apply statistical techniques to analyze medical data which enables the students to develop critical thinking skills and draw meaningful conclusions from complex datasets.
- **CO 5** : Develop research skills, including literature review, problem formulation, data collection, experimental design, and statistical analysis, to conduct independent biomedical research.

Distribution of Grades for the M. Sc. Biostatistics Programme with effect from 2023-24 Onwards									
	1	2	3	4	5	6	7	8	Total Credit s
	Discipli	ne Specific	Electives						
Semester	Core Courses (DSC)	Electives (DSE)	Interdisciplinary/ Multidisciplinary Elective	AEC 2 Credits	SEC(SE C) 2Credits	VAC /MOO C 2Credits	Internship /Field Visit /Minor Project /Institutional/In dustrial Visit 2Credits	Dissertation / Major Project	
1	MSBST01DSC01 MSBST01DSC02 MSBST01DSC03 MSBST01DSC04	<b>Pool A</b> MSBST01DSE01 to 02 (any 1)	1.000	57					10
	4 Credits x 4 = 16 Credits	3 Credits	\</td <td></td> <td></td> <td></td> <td></td> <td></td> <td>19</td>						19
2	MSBST02DSC05 MSBST02DSC06 MSBST02DSC07 MSBST02DSC08	Pool B MSBST02DSE03 to 04 (any 1)		Pool C	Pool D				23
	4 Credits x 4 = 16 Credits	3 Credits		2 Credits	2 Credits				
	-	Pool E MSBST03DSE05 to 06	Pool G						
3	MSBST03DSC09 MSBST03DSC10	(any 1) <b>Pool F</b> MSBST03DSE07 to 12 (any 2)	To be obtained from other Departments			VAC/ MOOC	MSBST03DSC 11		23
	4 Credits x 2 = 8 Credits	3Credits x 3= 9 credits	4 Credits			2* Credits	2 Credits		
4		Pool H MSBST04DSE13 to 14 (any 1) Pool I MSBST04DSE15 to 21 (any 1)						MSBST04DS C12	18
		3Credit x 2= 6 Credit	S					12 Credits	
	Total Credit for M. Sc. Biostatistics Program       83								

\*Credits are over and above the total credit requirement.

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	FIRST SEMESTER								
SI No	Course Code	Title of Paper	Contact Hours/Week		Marks				
			L	T/S	P	ESE	CE	Total	Credits
	D	DISCIPLINE SPECIFIC COR	E COU	JRSES	(DCI	E)			
1.1	MSBST01DSC01	Mathematical Methods for Biostatistics	4	(a)	V	60	40	100	4
1.2	MSBST01DSC02	Probability and Distribution Theory	4	2		60	40	100	4
1.3	MSBST01DSC03	Sampling Methods	4	1		60	40	100	4
1.4	MSBST01DSC04	Introduction to Biostatistics	4	2		60	40	100	4
DISCIPLINE SPECIFIC ELECTIVE COURSE (DSE)									
1.5	MSBST01DSExx	Elective-I-DSE (Pool A)	1	2	6	60	40	100	3
		Total credits	1.0						19

L=Lecture, T/S=Tutorials/Seminar, P/I=Practical/Internship, CE =Continuous Evaluation, ESE = End Semester Evaluation

SI No	POOL A:- List of Courses for Elective -I DISCIPLINE SPECIFIC ELECTIVES (DSE)								
1.5a	MSBST01DSE01	Biostatistical Computing Using R - I ( <b>Practical</b> )		2	6	60	40	100	3
1.5b	MSBST01DSE02	Biostatistical Computing Using SPSS - I ( <b>Practical</b> )		2	6	60	40	100	3

THUR UNIVERS

### FIRST SEMESTER M.Sc. BIOSTATISTICS PROGRAMME

Course Code & Title	MSBST01DSC01-MATHEMATICAL METHODS FOR BIOSTATISTICS				
Programme	M.Sc. Biostatistics	Semester I			
_	• To introduce the concept	of sequence and series.			
Courses	• To understand the improper integrals, beta and gamma functions.				
	• Learn Taylor's Theorem with applications.				
Objectives	<ul><li>Describe optima of functions using examples.</li><li>To achieve ideas on vector space, subspaces, independence of vec</li></ul>				
Objectives					
	basis and dimension.				
	• Establish the relation betw	veen algebraic and geometric multiplicity.			
	• To achieve ideas on quadratic forms and reduction of quadratic for				
	and gets ability for solvin	g problems in these areas.			

Modules	Content	Module Outcome
Module I: Sequence and series (15 Hours )	Sequences, series and their convergence, limit superior, limit inferior, limit of sequences, Cauchy sequence. Comparison test, D'Alembert's ratio test, Cauchy's root test, Raabi's test, Gauss test, Cauchy's integral test, Absolute convergence of series, Leibnitz's test for the convergence of alternating series, conditional convergence, indeterminate form, L'Hospital 's rule (problems only).	<ul> <li>The students will be able to:</li> <li>Explain convergences of sequences and series.</li> <li>Solve problems using various tests to examine the convergences of series.</li> <li>Explain the concept of L Hospital's Rule</li> </ul>

Module II: Special functions (15 Hours ) Module III: Vectors and Matrices (15 Hours )	The beta and gamma functions, duplication formula for gamma function, incomplete beta and gamma functions, functions of several variables, Limits and continuity, Taylor's theorem and its applications, Conditions for the optima of multivariate functions, Lagrange's method for constrained optimum, examples (bivariate case only) Vector space, Subspaces, Linear dependence and independence, Basis and dimensions, Matrices and determinants, symmetric, orthogonal and idempotent matrices, Row and column space of matrix, Rank, inverse, Characteristic polynomial, Cayley- Hamilton Theorem (statement and problem).	<ul> <li>Explain proper and improper beta and gamma functions.</li> <li>Understand the calculus of multivariable functions</li> <li>To find local and global optima of functions.</li> <li>To be familiar with vector space, subspace and examples.</li> <li>Explain linear dependence and independence.</li> <li>State Cayley-Hamilton theorem and solve problems.</li> </ul>
Module IV: Eigen values and spectral decomposition (15 Hours ) Reference	<ul> <li>Eigen values and eigen vectors, Spectral decomposition, Algebraic and geometric multiplicities, Generalized inverse, Quadratic forms, Classification of quadratic forms, Properties and reductions.</li> <li><i>Text Books</i> <ol> <li>Malik, S.C &amp; Arora, S. (2006). <i>Math</i> New-age international publishers.</li> <li>Mathai, A. M. &amp; Haubold, H. J. (2017). <i>Physicists and Engineers</i>, De Gruyter,</li> </ol> </li> </ul>	<ul> <li>Determine the Eigen values and Eigen vectors of the given matrix</li> <li>Write down the spectral decomposition of the given matrix</li> <li>Explain different types of quadratic forms.</li> </ul>

	Reference Books						
	1. Rudin, W. (2013). Principles of Real Analysis (3rdEd.)McGraw Hill.						
	2. Ramachandra Rao & Bhimasankaran (1992). Linear Algebra. Tata						
	McGraw Hill, New Delhi.						
	3. Apsostol, T. M. (1974). Mathematical Analysis, Second Edition. Narosa,						
	New Delhi.						
	4. Rao, C. R. (2002). Linear Statistical Inference and Its Applications,						
	Second Edition, John Wiley and Sons, New York.						
	After successful completion of this course, student will be able to:						
	1. Understand the concepts of limit and continuity of functions and their						
Course	properties						
Outcomos	2. Understand convergence of sequences and series of real numbers and						
Outcomes	functions.						
	3. Understand the vector space, matrices and its properties						
	4. Understand the properties of quadratic forms and its reduction.						

#### • Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

#### **ASSESSMENT RUBRICS**

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Components	Weightage
End Semester Evaluation	60
Continuous Eva	luation
Tests	16
Assignment	08
Seminar	16
Total	40

#### Sample Questions to Test Outcomes:

1. Define limit of sequence of real numbers. Give an example of a sequence for which limit does not exist.

- 2. What is meant by absolute convergence of series?
- 3. What is incomplete gamma function?.
- 4. State conditions for the optima of a multivariate function.
- 5. State Cayley-Hamilton theorem.
- 6. Explain Gram-Schmidth orthogonalization process.
- 7. Write a short note on different types of quadratic forms.

Course Code & Title	MSBST01DSC02- PROBABILITY AND DISTRIBUTION THEORY					
Programme	M.Sc. Biostatistics	Semester	Ι			
	The Course aims					
	• To introduce the basic concepts of	of probability				
Course	• To understand the connection between three approaches of					
Objectives	definitions of probability.					
X	• To get an idea on important theorems in probability using axiomatic definition of probability					
	• To learn about various discrete and continuous probability					
$\leq 1$	distributions needed for biostatist		5-7			
	OR UNIVE	/				

#### **DISCIPLINE SPECIFIC CORE COURSE**

Modules	Content	Module Outcome
<b>Module I:</b> Probability and Random Variables ( <b>15 Hours</b> )	Computation of probability based on classical and empirical definitions. Axiomatic approach to probability, probability space, conditional probability space, independence of events, Bayes' theorem and examples, random variable, distribution function, density function, expectation, variance and moments of a random variable and properties.	<ul> <li>Understand various definitions of probability</li> <li>Conditional probability and Bayes' theorem</li> <li>Concept of random variable and their distributions</li> </ul>
Module II: Important large sample theorems (15 Hours )	Definition of moment generating function and its limitations, characteristic function, elementary properties, characteristic functions and moments. Sequence of random variables, various modes of convergence of sequence random variables (definition only), Weak law of large numbers, strong law of large numbers, central limit theorem, DeMoivre-Laplace and Lindbergh- Levy forms of CLT. Applications of CLT in biostatistics.	<ul> <li>Definition of characteristic function</li> <li>Concept of weak and strong laws of large numbers</li> <li>Concept of central limit theorem and its applications in biostatistics</li> <li>Explain different discrete</li> </ul>
Special Discrete Distributions (15 Hours )	Discrete Uniform, Bernoulli, Binomial, Poisson, Geometric, Negative binomial, Hyper geometric, Multinomial. Properties of these distributions. Sample simulation and fitting of discrete distributions.	<ul> <li>Displain unificient discrete distributions.</li> <li>Properties of discrete distributions</li> <li>Simulation of samples from standard discrete distributions</li> </ul>

Module IV: Special Continuous Distributions (15 Hours )	Beta, Gamma, Normal, Weibull, Pareto, Laplace, Logistic, Cauchy and log-normal distributions. Properties of these distributions. Sample simulation and fitting of continuous distributions	<ul> <li>Explain different continuous distributions</li> <li>Properties of continuous distributions</li> <li>Simulation of samples from standard continuous distributions</li> </ul>
References	<ul> <li>Text Books <ol> <li>Krishnamurthy, K.(2006). Handbook of Applications .Chapman &amp; Hall/CRC, Ne</li> <li>Schinazi, R.B. (2010). Probability with Second Ed . Springer, New York.</li> </ol> </li> <li>Reference Books <ol> <li>Bhat, B.R. (2004). Modern Probability T New Delhi.</li> <li>Rohatgi, V. K. (2020). An Introduction to Mathematical Statistics, Wiley Eastern.</li> <li>Johnson, N.L., Kotz, S. and Balakrishnan Univariate Distributions, Vol. I &amp; Vol. II York.</li> <li>Johnson, N.L., Kotz. S. and Kemp. A.W. Distributions, John Wiley and Sons, New</li> </ol> </li> </ul>	Statistical Distributions with w-York Statistical Applications- Theory, New Age Publishers, o Probability Theory and N, N. (1995). Continuous I, John Wiley and Sons, New- .(1992). Univarite Discrete w York.
Course Outcomes	<ul> <li>After successful completion of this course, st</li> <li>1. Understand the concepts of probability and</li> <li>2. Understand characteristic function and its</li> <li>3. Understand various laws of large numbers</li> <li>4. Understand the concepts of discrete and co</li> <li>5. Understand the normal distribution and various their properties and applications for scientification.</li> </ul>	tudent will be able to: I properties. properties and central limit theorems. ontinuous distributions. rious non-normal distributions, ific research.

• Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

#### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation	60
Continuous Eval	luation
Assignment	08

#### Sample Questions to test Outcomes:

1. Define Poisson random variable. Find the moment generating function of a Poisson random variable.

2. Show that in the case of binomial distribution mean is always greater than variance,

however, mean equal to variance in the case of Poisson distribution.

3. Define t-statistic and explain its important applications. Write down the probability density function of Student's t-distribution.

4. Obtain the characteristic generating function of a standard normal distribution.

5. Define bivariate normal distribution. Show that linear combination of independent normal variables is normally distributed.

6. Define chi-square distribution. Obtain the MGF of the Chi-square distribution. Use it to obtain the mean and variance.

Course Code &			
Title	MSBST01DSC03-SAMPLI	NG METHODS	
Programme	M.Sc. Biostatistics	Semester	Ι
	• Explain different types of s	ampling	
Course	• Explain different errors in a	sampling	
Objectives	Difference between SRSW	R and SRSWOR	
	• Concept of stratified rando	m sampling	
	• Explain systematic sampling	ng	
	• Explain ratio and regressio	n estimators	

### DISCIPLINE SPECIFIC CORE COURSE

Modules	Content	Module Outcome
Module I: Sampling theory and Simple random sampling (15 Hours )	Introduction to sampling theory, Errors in sampling, simple random sampling (with and without replacement)-estimation of population mean and population mean square, determination of sample size, comparing efficiency of SRSWOR with SRSWR, simple random sampling with attributes.	<ul> <li>Concept of sampling theory</li> <li>Explain different types of errors</li> <li>Differentiate between SRSWR and SRSWOR</li> <li>Explain SRS with attributes</li> </ul>
Module II: Stratified random sampling and allocations. (15 Hours )	Stratified random sampling- estimation of population mean and variance, methods of allocation of sample size to different strata, comparison of allocations.	<ul> <li>Concept of stratified random sampling</li> <li>Explain methods of allocations.</li> </ul>

Module III: Complex sampling schemes (15 Hours )	Systematic sampling, circular systematic sampling. Cluster sampling, multistage sampling, multiphase sampling.	<ul> <li>Explain circular systematic sampling</li> <li>Explain cluster sampling</li> <li>Explain two stage cluster sampling</li> </ul>
Module IV: Auxiliary information based sampling (15 Hours )	Ratio and regression methods of estimation- bias and appropriate variances, unbiased ratio estimator, difference estimator, comparison of ratio estimator with regression estimator, Probability proportional to size sampling.	<ul> <li>Concept of ratio estimator</li> <li>Explain regression estimator</li> <li>Explain difference estimator</li> <li>Concept of PPS sampling</li> </ul>
References	<ul> <li><i>Text Books</i> <ol> <li>Singh, D and Chowdhary, F.S. (1986). <i>Theo Survey Designs</i>, New Age International, Net</li> <li>Cochran. W.G. (2007). <i>Sampling Technique</i> York.</li> </ol> </li> <li><i>Reference Books</i> <ol> <li>Des Raj, D. and Chandhok, P. (1998). <i>Samp</i> Publishing House, New Delhi.</li> <li>Gupta and Kapoor (2010). <i>Fundamentals of</i> Chand &amp; Sons.</li> <li>Murthy, M.N. (1967). <i>Sampling Theory &amp; Publishing Society, Calcutta.</i></li> <li>Parimal Mukopadhyay (2012). <i>Theory &amp; N</i> PHI Learning, New Delhi.</li> </ol> </li> </ul>	ory and Analysis of Sample ew Delhi. es, John Wiley & Sons, New ple Survey Theory, Narosa of Applied Statistics. Sulthan Methods. Statistical Methods of Survey Sampling,

	After successful completion of this course, student will be able to:
	1. Understand the concepts of probability and non-probability sampling.
Course	2 Understand the estimation methods for population mean, total and
Outcomes	proportion under various sampling schemes.
	3 Understand the use of auxiliary information for the estimation various
-	population parameters

• Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

#### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation	60
Continuous Eva	luation
Tests	16
Assignment	08
Seminar	16
Total	40
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#### Sample Questions to test Outcomes:

1. Explain probability sampling and non probability sampling.

2. Define standard error of sample mean and explain its uses in the construction of confidence interval, testing hypothesis and to obtain p-value.

3. Explain circular systematic sampling.

4. Prove that sample mean is an unbiased estimate of population mean under stratified random sampling.

5. Distinguish between ratio estimators and regression estimators.

6. Explain the difference between the methods of SRS and varying probability scheme.

### **DISCIPLINE SPECIFIC CORE COURSE**

Course Code & Title	de & MSBST01DSC04– INTRODUCTION TO BIOSTATIST		FATISTICS
Programme	M.Sc. Biostatistics	Semester	Ι
Course Objectives	<ul> <li>Introduce the concept of proper</li> <li>Introduce the concept of risk, if</li> <li>Explain applications of various research</li> <li>Explain concept of estimation</li> <li>Introduce the concept of hypot clinical research.</li> </ul>	ortions, ratio and odds relative risk and their r s probability models in and its applications in thesis testing and appli	neasurement n medical n biostatistics cations in

Modules	Content	Module Outcome
Module I:	Proportions:- Comparative Studies, Screening Tests Displaying Proportions	• Explain proportions, ratios
Descriptive	Rates:- Changes, Measures of Morbidity	and rates
Methods for	and Mortality, Standardization of Rates.	• Explain risk, relative risk and odds ratio.
Categorical Data	Ratios:- Relative Risk, Odds and Odds Ratio Generalized Odds for Ordered 2xk	• Explain Mantel-Haenszel
(15 Hours )	Tables, Mantel-Haenszel Method,	Method and Standardized
A.	Standardized Mortality Ratio.	Monanty Ratio

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Module II: Descriptive Methods for Continuous Data (15 Hours )	Tabular and Graphical Methods:- One- Way Scatter Plots, Frequency Distribution, Histogram and the Frequency Polygon, Cumulative Frequency Graph and Percentiles, Stem-and-Leaf Diagrams, Measures of Location, Measures of Dispersion, Box Plots, Special Case of Binary Data, Coefficients of Correlation, Pearson's Correlation coefficient, Nonparametric Correlation coefficients.	<ul> <li>Explain graphical and tabular methods</li> <li>Explain measures of central tendency and dispersion</li> <li>Explain parametric and non parametric correlations</li> </ul>
Module III: Probability Models for data and estimation (15 Hours )	Practical applications of Normal, Binomial and Poisson distributions in bio medical research, Pair-Matched Case–Control Study, Introduction to Confidence interval Estimation, Estimation of Proportions, Estimation of Odds Ratios, Estimation of Correlation Coefficients.	<ul> <li>Explain various applications         <ul> <li>of probability models in             medical research</li> </ul> </li> <li>Estimation of proportions,         <ul> <li>odds ratio and correlation</li> <li>coefficients</li> </ul> </li> <li>Give an introduction to Pair-         <ul> <li>Matched Case-Control Study             and confidence interval             estimation</li> </ul> </li> </ul>
Module IV: Introduction to Statistical Tests of Significance (15 Hours )	Hypothesis Tests, Statistical Evidence, Errors, Summaries and Conclusions, Rejection Region, P Values, Type I and Type II Errors, Relationship to Confidence Intervals. One- Sample Problem with Binary Data, Analysis of Pair-Matched Data, Comparison of Two Proportions, Mantel–Haenszel Method, Inferences for General Two-Way Table, Fisher's Exact Test, Ordered 2x k Contingency Tables.	<ul> <li>Explain basic concepts of hypothesis testing and P value</li> <li>Comparison of Population proportions</li> <li>Understand , Mantel– Haenszel Method, and Fisher's Exact Test.</li> </ul>

	Tast Pook
	Text Dook
	1. Chap T.L. (2003). Introductory Biostatistics, John Wiley & Sons.
	Reference Books
	1. Rosner, B. (2010). Fundamentals of Biostatistics, Cenage Learning,
	Harvard University.
References	2. Chernick, M.R. and Fris, R.H. (2003). Introductory Biostatistics for the
	Health Sciences, John Wiley & Sons.
	3. Peter Armitage, Geoffrey Berry, J. N. S. Matthews (2008). Statistical
	Methods in Medical Research. John Wiley & Sons
	4. Daniel, Wayne W (2009). Biostatistics: A Foundation for Analysis in the
	Health Sciences. John Wiley & Sons.
	After successful completion of this course, student will be able to:
	1. Understand the descriptive methods for different types of data.
Course	2. Understand the concepts of risk, odds and odds ratio.
Outcomes	3. Understand the concept of inferential procedures for medical research.
	4. Understand different methods of comparison of proportions for biostatistical
	studies.

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• Lecturing, Visualization, Team Learning

# **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

# **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation	60
Continuous Eva	luation
Tests	16
Assignment	08
Seminar	16
Total	40

#### Sample Questions to Test Outcomes:

- 1. Define odds and odds ratio.
- 2. Explain sensitivity and specificity.
- 3. Describe relative risk. How to quantify it.
- 4. Explain Mantel-Haenszel Method.
- 5. Explain contingency table.
- 6. Describe Pair-Matched Case-Control Study.

Course Code & Title	MSBST01DSE01 - BIOSTATISTICAL COMPUTING USING R-I (PRACTICAL)						
Programme	M.Sc. Biostatistics	Semester	I				
Course Objectives	<ul> <li>Define the basic concepts of R software and R packages</li> <li>Describe various concepts required for developing the R Language</li> <li>Build our new functions in R</li> <li>Illustrate different R-Graphics facilities</li> <li>Find rank and inverse using R software</li> <li>Describe different sampling methods using R software</li> </ul>						
Modules	Content	Module C	Jutcome				
Module I: Basic Concepts of R Programming (20 Hours )	Introduction to R- Objects and their classes, operators, vectors and matrices, list and data frames, indexing and accessing data, importing and exporting data. Common built-in functions, R- Graphics.	<ul> <li>Define bas software R</li> <li>Demonstra data structu arrays, mat Class funct</li> <li>Design an R Languag Expression Symbols, F</li> </ul>	ics of statistical ate the important ares such as rix, data frames, tion etc. overview of the e such as s, Objects, Functions.				

### POOL A: DISCIPLINE SPECIFIC ELECTIVES

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Module II: Matrices and Standard Probability Distributions (25 Hours )	Matrices, rank, determinants and inverse. Eigen values and vectors, power of matrices, g-inverse, system of linear equations, roots of algebraic and transcendental equations. Plotting of cdf and pdf of standard distributions. Generations of random samples from standard distributions, demonstrations of the sampling distributions	<ul> <li>How to find rank and inverse using R software.</li> <li>How to solve system of linear equations using R software</li> <li>Plotting pdf and cdf curve of different distributions</li> </ul>
Module III: Biostatistical Sampling Methods (25 Hours )	Random samples elections, estimation of mean pro-portion, variance, confidence interval and efficiency under SRS, stratified random sampling, Various kind of allocation, stratification, estimators based on ratio and regression methods pps sampling, two stage cluster sampling, and systematic sampling.	<ul> <li>How to draw random samples using different sampling techniques</li> <li>PPS sampling techniques using R softwares</li> <li>Ratio and regression methods using R softwares.</li> </ul>
Module IV: Biostatistical data analysis (20 Hours )	Measures of Morbidity and Mortality in R, Relative Risk, Odds and Odds Ratio, Generalized Odds for Ordered 2 x k Table, Mantel– Haenszel Method, Box Plots, Estimation of Proportions and Odds Ratios, testing of hypotheses.	<ul> <li>Compute Morbidity and Mortality in R</li> <li>Computation of Odds and odds ratio using R</li> <li>Data description using Box plot</li> <li>Inference procedures in R</li> </ul>

References	<ol> <li>Text Books         <ol> <li>Maria D.U., Ana F.M. and Alan T.A. (2008): Probability and Statistics with R. CRC Press.</li> <li>Dalgaard, P. (2008): Introductory Statistics with R, (Second Edition), Springer.</li> </ol> </li> <li>Reference Books         <ol> <li>Purohit, S.G, Ghore, S.D and Deshmukh, S.R. (2004): Statistics Using R. Narosa.</li> <li>Babak Shahbaba. (2012). Biostatistics with R: An Introduction to Statistics through Biological Data. Springer New York.</li> </ol> </li> </ol>
Course Outcomes	<ul> <li>After successful completion of this course, student will be able to: <ol> <li>Understand various built in functions in R programming for biostatistical data analysis.</li> <li>Understand different functions in R programming for writing compute r programmes and develop computer programmes for different problems</li> <li>Understand the usage of packages in R for drawing various diagrams and computing descriptive statistics, comparison of means, ANOVA, non-parametric tests, simple correlation and regression procedures</li> </ol> </li> </ul>

• Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Hands on training

#### **ASSESSMENT RUBRICS**

Components	Weightage				
End Semester Evaluation	60				
Continuous Eva	aluation				
Practical Tests	32				

Record	08
Total	40

#### Sample Questions to Test Outcomes:

1. Write an R program to create a matrix taking a given vector of numbers as input. Display the matrix.

2. Import a given dataset in R, and conduct its descriptive analysis.

3. Select a simple random sample of 50 numbers without replacement from the numbers 1 to 2000.

4. Generate a random sample of size 100 from a standard normal distribution.

5. Illustrate the law of large numbers using R.

6. Enter the given 2 matrices, and find their product.

Course Code & Title	MSBST01DSE02-BIOSTA SPSS -I (PRACTICAL)	TISTICAL COMPU	TING USING
Programme	M.Sc. Biostatistics	Semester	Ι
Course Objectives	<ul> <li>The main focus of the corresearch question using S</li> <li>Illustrate different toolbo</li> <li>Data definition and acces</li> <li>Apply SPSS software to a</li> <li>Students get awareness to technique and interpret research</li> </ul>	urse will be on to solve SPSS oxes in SPSS and data analysis and develop different statis o chose appropriate stat esults using SPSS.	e biostatistical l presentation. tical tools tistical

### POOL A: DISCIPLINE SPECIFIC ELECTIVE

Modules	Content	Module Outcome		
Module I: SPSS Environment, Basic Concepts of SPSS Programming (20 Hours )	Introduction to SPSS- Starting SPSS, Working with data file, SPSS windows, Menus, Dialogue boxes. Preparing the Data file, Creating data file and entering data, Defining the variables, Entering data, modifying data file, import file. Variable types in SPSS and Defining variables – Creating a Codebook in SPSS. Screening and cleaning data, Manipulation of data.	<ul> <li>Understand the installation and familiar with toolboxes of SPSS.</li> <li>Data management and modifications of data.</li> </ul>		
<b>Module II:</b> Preliminary Analysis in SPSS ( <b>25 Hours</b> )	Computing Variables- Recoding (Transforming) Variables: Recoding Categorical String Variables using Automatic Recode - Sorting Data - Grouping or Splitting Data. Categorical variables, continuous variables. The Explore procedure - Frequencies Procedure – Descriptive - Compare Means - Frequencies for Categorical Data, different statistical distributions	<ul> <li>Working with Data types</li> <li>Recoding and sorting</li> <li>Descriptive statistics</li> <li>Explore procedure, graphics in SPSS</li> </ul>		
Module III: Inferential Statistics (25 Hours )	Pearson Correlation, Chi-square Test of Independence – Inferential Statistics for Comparing Means: One Sample t Test, Paired Samples T Test, Independent Samples T Test, One-Way ANOVA. Two way ANOVA, Multivariate ANOVA.	<ul> <li>Compute and interpret correlation coefficients</li> <li>Learn how to conduct various statistical tests using SPSS</li> <li>Preparing ANOVA</li> </ul>		

Module IV: Non- Parametric statistics (20 Hours )	<ul> <li>Learn how to perform non parametric tests</li> <li>Kruskal- Wallis test. Interpreting the output of tests, p-value computation.</li> <li>Learn how to perform non parametric tests</li> <li>Get p value of various tests</li> <li>Interpretation of test results</li> </ul>				
References	<ol> <li>Text Books         <ol> <li>Hinton, P. R., Brownlow, C, Mc Murray, I. and Cozens, B. (2004): <i>SPSS Explained</i>, Routledge, Taylor and Francis group, New York.</li> </ol> </li> <li>Reference Books         <ol> <li>Field, A. (2011); Discovering Statistics Using SPSS, Sage Publications.</li> <li>William E. Wagner. (2015). Using IBM SPSS statistics for research methods and social science statistics, Fifth edition, SAGE         </li></ol> </li> </ol>				
Course Outcomes	<ul> <li>After successful completion of this course, student will be able to:</li> <li>1. Build capacity to analyzing complex information with the help of SPSS.</li> <li>2. Understand with the tool box of statistical software SPSS</li> <li>3. Summarize variables using frequencies and descriptive analysis.</li> <li>4. Understand to producing cross tabulation tables and testing for significant relationships with chi square test.</li> <li>5. Understand the usage of assessing relationships between continuous variables through plots and correlations.</li> </ul>				

• Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Hands on training

#### **ASSESSMENT RUBRICS**

Components	Weightage					
End Semester Evaluation	60					
Continuous Evaluation						
Practical Tests	32					
Record	08					
Total	40					

#### Sample Questions to Test Outcomes:

1. Compute t-test for difference of means on a given continuous variable, based on a categorical variable.

2. Compute descriptive statistics of a given continuous variable of a dataset.

3. Perform an appropriate ANOVA for the given data.

4. Perform appropriate non parametric test for the given data.

5. Load the given external spreadsheet data into SPSS, and obtain the pie chart, histogram of the variables.

6. Perform MANOVA for the given data set.

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	SECOND SEMESTER										
Sl No	Course Code	Title of Paper		Contact ours/Week		Contact Hours/Week		Mar	∶ks		
			L	T/S	P	ESE	CE	Total	Credits		
	DI	SCIPLINE SPECIFIC CORE	COL	JRSES	(DC)	E)					
2.1	MSBST02DSC05	Biostatistical Inference	4		1	60	40	100	4		
2.2	MSBST02DSC06	Applied Regression Analysis	4	1	V	60	40	100	4		
2.3	MSBST02DSC07	Statistical Epidemiology	4	1	-	60	40	100	4		
2.4	MSBST02DSC08	Survival Analysis	4	1		60	40	100	4		
	I	DISCIPLINE SPECIFIC ELF	ЕСТГ	VES (I	DSE)	l		I			
		Elective-II-DSE (One course		-							
2.5	MSBST02DSExx	has to be chosen from <b>Pool</b>		2	6	60	40	100	3		
		<b>B</b> )	P	-							
ABILITY ENHANCEMENT COURSE(AEC)											
		Offered to other									
	MSBST02AECxx	Departments. ( One	2				100	100	2		
2.6		course has to be chosen					100	100	-		
2.0		from <b>Pool C</b> )	1		_						
		(To be obtained from				P		*	2		
		other Departments)			100	-			2		
	S I	SKILL ENHANCEMENT C	COUR	RSE(SE	C)	7					
	14	Offered to other	1	1	A	$\langle \Lambda \rangle$	5				
	MSBST02SECxx	Departments. ( One	2	13	S .	60	40	100	2		
27	MODUTOZOLCXX	course has to be chosen		29	1	00		100	2		
2.7	1	from <b>Pool D</b> )	E	5/	C						
		(To be obtained from	-					*	2		
		other Departments)				_					
		Total Credits		·					23		

L=Lecture, T/S=Tutorials/Seminar, P/I=Practical/Internship, CE =Continuous Evaluation, ESE = End Semester Evaluation.

(\*Note: Evaluation is determined by respective Department)

SUNO		POOL	L B									
SINU		<b>DISCIPLINE SPECIFI</b>	C ELE	CTIV	E (DSI	E)						
251	MSBST02DSF03	Biostatistical Computing		2	6	60	40	100	3			
2.3.1	WISDS102DSL05	Using SPSS-II (Practical)		2	0	00	10	100	5			
252	MSBST02DSF04	Biostatistical Computing		2	2	2	2 6	6	60	40	100	3
2.3.2	Using SAS-I (Practical)	Š.		00	10	100	5					
SI No		POOI	LC	1.0	1							
SITU		ABILITY ENHANCEM	ENT C	OURS	SE (AE	<b>C</b> )						
		A foundation course in										
2.6.1	MSBST02AEC01	LaTex for scientific	100	2	6	60	40	100	2			
		documentation		Þ								
2.6.2	MSBST02AEC02	Basic Statistical data analysis		2	6	60	40	100	2			
		using EXCEL		1 and 1	Ū	00		100	_			
SI No		POOI	LD									
		SKILL ENHANCEME	NT CO	DURS	E (SEC	<b>C)</b>						
2.7.1	MSBST02SEC01	Exploratory Data Analysis		2	6	60	40	100	2			
		Using SPSS										
2.7.2	MSBST02SEC02	Regression Analysis Using		2	6	60	40	100	2			
		SPSS					-					



#### SECOND SEMESTER M.Sc. BIOSTATISTICS PROGRAMME

Course Code & Title	MSBST02DSC05-BIOSTATISTICAL INFERENCE						
Programme	M.Sc. Biostatistics Semester II						
Course Objectives	<ul> <li>A thorough understanding of parameter such as sufficience efficiency.</li> <li>Understanding the notion of for sufficiency, minimal and</li> <li>Derivation of the Cramer-Raexistence of MVB estimator Lehmann-Scheffe theorems</li> <li>To introduce the concept of significance level, power of significance level, power of simple hypothesis against si</li></ul>	f important properties of estimators of a y, consistency, unbiasedness and Fisher-Neymann factorization theorem complete sufficient statistics. to lower bound and the conditions for the apply the concept of Rao-Blackwell and to obtain UMVUE of a parameter. Testing of hypothesis, critical region, The test and p-value. Funiformly most powerful test for testing imple alternative and obtain sequential ting the hypothesis. ikelihood ratio test and confidence polications.					

Modules	Content	Module Outcome
1	Unbiasedness, consistency,	• Derive the important
V	consistent asymptotically normal (CAN) estimators, efficiency,	<ul> <li>Determine the sufficient</li> </ul>
Module I:	sufficiency, invariance property of	statistic
Properties of	consistent estimators, Fisher-	• Determine consistent
estimators.	Neymann factorization theorem for	estimators and consistent
	sufficiency (proof for discrete	and asymptotically normally
(15 hours)	distributions only), joint sufficient	distributed estimators.
	statistics, minimal and complete	• Identify efficient
	sufficient statistic.	estimators.

	Minimum variance unbiased	• State and prove Cramer-
	estimator (MVUE), Likelihood and	Rao inequality.
Module II:	score functions. Fisher information,	• Examples of Minimum
Minimum	Cramer-Rao inequality and its	variance bound estimator.
Variance	applications, Cramer-Rao Lower	Apply Rao-Blackwell
Unbiased	Bound (CRLB), Minimum variance	and Lehmann-Scheffe
Estimation.	bound unbiased estimator (MVB).	theorems to find UMVUE.
	Rao-Blackwell and Lehmann-	• Understand the concept
(15 hours)	Scheffe theorems. Method of	of MLE and its properties.
	moments and method of MLE and	• Examples of moment
	their properties.	estimators.
	Null and alternative hypotheses,	• Identify null, alternative,
	simple and composite hypotheses,	simple and composite
	two types of errors in testing of	hypothesis
Module III:	hypothesis, p-value, level of	• Find critical region, size
Tests of	significance and size of test, power	and power of the test.
Hypotheses and	function, Neymann-Pearson lemma,	Apply Neymann-
Most Powerful	most powerful and uniformly most	Pearson lemma to find most
Tests.	powerful tests. Sequential	powerful test.
	Probability Ratio Test	• Derive SPRT for test the
(15 hours)	(SPRT)(Concept only).	parameters of normal
N.		distribution, exponential and
A	4	Poisson distributions,
	7.	binomial distribution.
-1/	Different parametric test (Z ,t and F	• Apply different parametric
all	tests), large sample tests. Likelihood	test for various real life
Module IV:	ratio test, monotone likelihood ratio	applications.
Parametric test,	property. Confidence interval	• Understand likelihood ratio
Likelihood ratio	estimation of mean and variance,	property and its applications.
test and SPRT	difference of mean and population	• Compute the confidence
(15 hours)	proportion, and difference of	interval for mean, population
	population proportion.	proportional and variance.
	Text Books	
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	1. Hogg, R. V., McKean, J. W., & Craig, A. T. (2013). Introduction to	
	Mathematical Statistics. Pearson Education India.	
	2. Vijay K. Rohatgi, A. K. Md. Ehsanes Saleh (2015). An Introduction to	
	Probability and Statistics, 3rd Edition, John Wiley and Sons, NewYork.	
	3. Mood, A. M., & Graybill, F. A. (6). Boes, DC (1974). Introduction to	
References	the Theory of Statistics. Third edition. McGraw Hill.	
	Reference books	
	1. Casella,G.andBerger,R.L.(2002). <i>StatisticalInference</i> , <i>SecondEdition</i> , Dux bury, Australia.	
	2. Lehman, E. L. (1986): <i>Testing of Statistical Hypotheses</i> . John Wiley, New York.	
	3. Lehmann, E. L(1983). <i>Theory of Point Estimation</i> , John Wiley and Sons, New York.	
	After successful completion of this course, student will be able to:	
	1. Understand the concepts of Sufficiency and Completeness	
	2. Understand the concepts of Minimum Variance Unbiased	
	Estimation.	
Course	3. Understand various estimation methods and applications in real	
Outcomes	life problems	
5	4. Apply various parametric and sequential testing procedures	
A	to deal with real life problems.	
1	5. Understand Most Powerful Tests for testing simple null hypothesis	
<1/	and developing MP tests for different problems.	
100	VIII ISC	

• Lecturing, Hands on Training, Visualization, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Discussion, Questioning and Answering ASSESSMENT RUBRICS

Components	Weightage
End Semester	60
Evaluation(ESE)	
Continuous	Evaluation
Tests	16
Assignment	08
Seminar	16
Total	40

### Sample Questions to Test Outcomes:

- 1. State factorization theorem for sufficiency.
- 2. Let  $X_1, X_2, ..., X_n$  be a random sample of n observations from Gamma distribution with shape parameter  $\theta$ . Find sufficient statistic for  $\theta$ .
- 3. State and prove Basu's theorem.
- 4. Let X<sub>1</sub>, X<sub>2</sub>, ..., X<sub>n</sub> be random sample from a Poisson population with parameter  $\lambda$ . Show that the  $\overline{X}^2$  is a UMVUE for  $\lambda^2$ .
- 5. Let  $X_1, X_2, ..., X_n$  be a random sample of size n observations from beta first kind distribution with parameter  $\alpha$  and  $\beta$ . Find the estimators of  $\alpha$  and  $\beta$  by method of MLE.
- 6. Let p be the probability that a coin will fall head in a single toss in order to testH<sub>0</sub> : p = 1/2against H<sub>1</sub> : p = 3/4. The coin is tossed 5 times and H<sub>0</sub> is rejected if more than 3 heads are obtained. Find the probability of type-I error and power of the test.
- 7. Define a most powerful test and explain the utility of Neyman-Pearson lemma.
- 8. Given the nine sample values 4.5, 6.5, 3.8, 4.2, 7.7, 8.5, 9.4, 5.3, 3.9 from a normal distribution with mean μ and variance 4. Find the best critical region for testingH<sub>0</sub> : μ = 4 versus H<sub>1</sub> : μ = 5 of size 0.05. Also calculate the power of the test.
- Obtain OC function for testing H<sub>0</sub> : p = p<sub>0</sub> versus H<sub>1</sub> : p = p<sub>1</sub> using SPRT with strength (α, β) based on observations from b(n, p).
- Show that the likelihood ratio test for testing the equality of variances of two normal distributions is the usual F-test.

# **DISCIPLINE SPECIFIC CORE COURSE**

Course Code &	MSBST02DSC06-APPLIED REGRESSION ANALYSIS	
Title		
Programme	M.Sc. Biostatistics	Semester II
Course Objectives	<ul> <li>Describe simple linear</li> <li>Apply principle of leas in simple linear regress</li> <li>Describe multiple linea model adequacy.</li> <li>Identify multicollineari problem of estimation of and also Identify auto of</li> <li>Explain polynomial reg</li> <li>Understand the notion</li> <li>Explain generalized line</li> <li>Describe the logistic reg</li> </ul>	regression models and its properties t square method to estimate the parameters ion models. r regression models and its properties and ty problem, its consequences, discuss the of parameters when multicollinearity occurs correlation and its consequences. gression in one and several variables. of nonlinear regression. ear model. gression, Poisson regression.

Modules	Content	Module Outcome
	The simple linear regression	• Explain simple linear
5	models, least square estimation,	regression model
Module I:	statistical assumptions and	• Describe least square
Linear	properties of estimators, standard	estimators.
Regression	error of estimates, tests of	• Articulate to inference
Models	significance and confidence	regarding regression
(15 Hours)	intervals for the parameters, error	parameters.
	and residual plots.	• Explain ANOVA.
	Multiple regression models, OLS	• Explain multiple linear
Module II:	and ML estimators, testing and	regression models.
Regression	prediction. Multicollinearity,	• Explain multicollinearity.
Diagnostics	heteroscedasticity, autocorrelation:	• Discuss detection and
(15 Hours)	their nature, consequences,	remedial measures of

	detection, remedial measures and	multicollinearity.
	estimation in the presence of them.	• Explain heteroscedasticity
		and remedial measures of
		heteroscedasticity.
Module III:	Polynomial regression in one and	Explain polynomial
Polynomial and	several variables. Linearization	regression.
nonlinear	transforms, Diagnostic checks and	• Discuss non-parametric
regression	correction. Nonparametric	regression.
(15 Hours)	regression and concept of spline	• Explain concept of spline
	smoothing.	smoothing.
	Generalized linear models.	Discuss generalised linear
Module IV:	Logistic regression, Poisson	model.
Non-Linear	Regression. Estimation, model	• Explain logistic
Regression	adequacy of GLM and diagonostic	regression.
(15 Hours)	tests.	• Introduce the concept of
		Poisson regression.
References	<ul> <li><i>Text Books</i></li> <li>1. McCullagh, P. (2019). General Kingdom: CRC Press.</li> <li>2. Hosmer, D.W. and Lemeshow, S. (1) John Wiley.</li> <li>3. Montgomery, D. C, Peek, E.A. <i>Introduction to Linear Regression A</i></li> <li><i>Reference books</i></li> <li>1. Seber, G. A. F. and Lee, A. J. (20) Wiley</li> <li>2. Draper ,N.R. and Smith,H.(1998): <i>Ed.</i> John Wiley.</li> <li>3. Goon,Gupta,DasGupta(2001):<i>An Ot</i> World Press.</li> <li>4. Chatterjee, S., &amp; Hadi, A. S. (2013)</li> </ul>	Alized Linear Models. United (989): Applied Logistic Regression, and Vining, G.G. (2006): nalysis, John Wiley. 003): Linear Regression Analysis, Applied Regression Analysis, 3 <sup>rd</sup> utline Series in Statistics Vol11, . Regression Analysis by Example.

	After successful completion of this course, student will be able to:
	1. Understand various regression models including logistic regression
	models.
Course	2. Understand consequences of multicollinearity, heteroscedasticity,
Outcomes	autocorrelation, their detection and remedial measures.
Outcomes	3. Apply statistical techniques to model relationships between
	variables and make predictions.
	4. Acquire knowledge of various advanced regression models
	with applications in biostatistics.

• Lecturing, Hands on Training, Visualization, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Discussion, Questioning and Answering

### ASSESSMENT RUBRICS

Components	Weightage
End Semester	60
Evaluation(ESE)	
Continuous Ev	aluation
Tests	16
Assignment	08
Seminar	16
Total	40

### Sample Questions to Test Outcomes:

1. Derive the OLS estimators of intercept and slop coefficients of a simple linear regression model.

2. Explain measure of goodness of fit in regression analysis. obtain its relationship with correlation coefficient.

- 3. Define multicollinearity.
- 4. How to detect heteroscedasticity using Spearman's rank correlation test.
- 5. Discuss Durbin-Watson test for autocorrelation.
- 6. Explain the difference between  $R^2$  and adjusted  $R^2$  in multiple regression. Mention its uses

- 7. What is logistic regression model?
- 8. Explain link function and linear predictor?
- 9. Explain the parameter estimation and inference of GLM.
- 10. Explain orthogonal polynomial regression.

Course Code & Title	MSBST02DSC07-STATISTICAL EPIDEMIOLOGY	
Programme	M.Sc. Biostatistics	Semester II
	• To understand basic conc study designs in epidemic	epts of epidemiology and explain different blogy.
	• Identifying achievements disease.	in epidemiology and measuring health and
	• To aquire the knowledge	of cohort study designs, case control study
Course	designs.	
Objectives	• Develop the knowledge o	f various matched case control studies and
	cross over study designs.	
	• To know about statistical	concepts and inference.
	• Understand relationship b	between variables.
	• State Mendal's law and ex	stimation of allele frequency, estimation of
	allele frequencies, Hardy-	Weinberg law.
5	• Introduce the concept of a	detection and estimation of linkage,
- A.	inheritance of quantitative	e traits, stochastic models of carcinogenesis

# DISCIPLINE SPECIFIC CORE COURSE

Modules	Content	Module Outcome
1	Basic concepts of	
	epidemiology: definition and	• Explain the basic
Module I:	scope of epidemiology,	concepts of epidemiology.
Basic concepts	achievements in epidemiology,	• Measuring health and
of	measuring health and disease,	disease
Epidemiology	definition of health and disease,	• Comparing disease
(15 Hours)	measures of disease frequency,	occurrence.
	comparing disease occurrence.	

	Types of study: observations and	
Module II:	experiments, observational	• Explain different study
Study designs	epidemiology, cohort study designs,	designs in epidemiology.
in	case control study designs,	• Understand various
epidemiology	randomized, field trails, control trials,	cohort and case control
(15 Hours)	potentials errors in epidemiological	studies in biostatistics.
	studies, ethical issues	20.
	Distribution and summary measures:	101
Madula III.	distribution, measures of central	• To understand statistical
Exploratory	tendency, measures of variability,	
exploratory	normal and log normal distributions,	concepts.
(15 Hours)	estimation, testing, CI, ANOVA,	• 10 know about
(13 110018)	relationship between two variables: chi-	relationship between two
	square test, correlation, regression,	variables.
	logistic regression.	<i>c</i>
		• Understand the concept
	Concept of cause, establishing the	of Concept of cause.
Module IV:	cause of disease, scope of	• Identifying the scope of
	prevention, levels of prevention,	prevention.
epidemiology	Introduction to clinical	• Understand the basic
(15 Hours)	epidemiology.	concepts of clinical
5		epidemiology.
- A		A/
	Text Books	
<1/	1. Beaglehole, R., Bonita, R. and	Kjellstorm, T. (1993). Basic
1	Epidemiology. World Health Organiza	ation, Geneva.
	2. Newman, S.C. (2001). Biostatistical M	Iethods in Epidemiology. John
References	Wiley & Sons, New York.	
	3. Virasakdi, C. (2010). Analysis of Epi	demiological Data Using R and
	Epicalc. Epidemiological Unit, Songl	a University,Thailand
	Reference books	
	1. Rothan, K. J., Greenland, S. and Lash	, T. L. (2008). Modern
	<i>Epidemiology, 3<sup>rd</sup> Edition</i> . Wokers K1	uver.

	2. Clayton, D.and Hills, M.(1993). Statistical Methods in Epidemiology.
	Oxford University Press.
	After successful completion of this course, student will be able to:
Course	1. Understand Basics of epidemiology.
Outcomos	2. Understandtypes of study used in epidemiology
Outcomes	3. Understand the concept of clinical epidemiology.
	4. Identify the scope of cause and prevention of disease.

• Lecturing, Hands on Training, Visualization, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Discussion, Questioning and Answering ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Ev	aluation
`ests	16
Assignment	08
Assignment Seminar	08 16

# Sample Questions to Test Outcomes:

- 1. Explain Basic concepts of epidemiology along with its scope.
- 2. Narrate the notion of measures of disease frequency.
- 3. Differentiate between health and disease.
- 4. Elaborate various types of study in epidemiology.
- 5. Differentiate between cohort study and case control study.
- 6. Describe the potential error in epidemiological studies.
- 7. Define ANOVA and mention its application.

8. Why we need chi-square test? Elaborate the concept.

9. Explain the concept of clinical epidemiology.

10. Differentiate between scope of prevention and levels of prevention in epidemiological study.

Course Code & Title	MSBST02DSC08-SURVIVAL ANALYSIS		
Programme	M.Sc. Biostatistics	Semester II	
	• Understand the basic notion	on of survival analysis	
	• Understanding application	s of survival function, hazard function,	
	mean residual life function	as and other aging concepts.	
	• Understand and examine	• Understand and examine the properties of standard lifetime	
Course	distributions.		
Objectives	• Develop the concept of nonparametric tests like KS test, sign test,		
Mann-Whitney and Wilcoxon U tests.		xon U tests.	
	• Introduce the concepts of censoring and truncation and		
	classifications.		
	• Introduce the notion of Estimating survival rates using large scale data		
	like DHS, NFHS, DLHS, etc. Comparing survival curves.		
	• Introduce the notion of Kaplan-Meier estimation technique, life tables,		
Α.	Mantel-Haenszel test.		

Modules	Content	Module Outcome
<b>Module I</b> Basics of survival analysis ( <b>15 Hours</b> )	Basics of survival analysis- discrete and continuous time models, survival function, hazard rate function, probability density function, mean residual life time. Aging classes-IFR, IFRA and their duals, Bathtub failure rate.	<ul> <li>Understand the basic concepts and ideas of survival analysis.</li> <li>Understand the basic concepts of ageing classes.</li> </ul>
<b>Module II</b> Life	Life distributions-exponential, Weibull, lognormal and gamma distributions,	• Examine the properties and methods for standard

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distributions	characterizations. Nonparametric tests-	survival time distributions.	
(15 Hours)	Kolmogorov-Smirnov test, sign test and	• Understand the basic	
	signed-rank test, Mann-Whitney,	concepts of nonparametric	
	Wilcoxon U test, chi- square test for	tests.	
	goodness of fit, test for independence of		
_	attributes.	1-	
	Concepts of censoring Mechanism -Type-		
	I, Type-II and random censoring,	• Estimate survival	
Madada III	Progressive censoring, Truncation,	functions using parametric	
	Methods for truncated and interval	and non-parametric	
Censoring and	censored data. Likelihood construction and	methods.	
I runcation	estimation of Censored and Truncated	• Understand the basic	
(15 Hours)	Data. Estimating survival rates using large	concepts of censoring and	
	scale data like DHS, NFHS, DLHS, etc.	truncations.	
	Comparing survival curves.		
	Kaplan-Meier estimation technique,	Apply and interpret	
Module IV	life tables, Mantel-Haenszel test.	regression models for survival	
Estimation of	Interval estimation of survival	data .	
Survival	probabilities. Introduction to	• Understand the	
function	survival regression. Cox	concept of Cox-Proportional	
(15 Hours)	proposional hazard model.	hazard model.	
5		17	
A.	1 Lowless LE (2002): Statistical Math	a da fan Lifetina (Sacand	
	I. Lawless, J.F. (2003): Statistical Methods for Lifetime (Second		
<1/	<i>Edition)</i> , John Wiley & Sons Inc., New Jersey.		
1	2. Kalbfleisch, J. D. and Prentice, R.L. (1980): The Statistical		
References	Analysis of Failure Time Data, John Wiley & Sons Inc. New		
	Jersey.		
	3. Moore, D.F. (2016): Applied Survival Analysis Using R,		
	Springer.		
	Kejerence Books		
	1. Klein J.P. and Moeschberger M.L. (2003) Survival Analysis -		
	Techniques for Censored and Truncated Data, Second Edition,		

	Springer-Verlag, New York.
	2. Miller, R. G.(1981): Survival Analysis, John Wiley & Sons Inc.
	3. Bain, L. G.(1978): Statistical Analysis of Reliability and Life testing
	Models, Marcel Decker.
	4. Cox, D.R and Oakes, D.(1984): Analysis of Survival Data. Chapman
~	and Hall.
	5. Fraser, D.A.S.(1957): Non-parametric Methods in Statistics, Wiley,
	New York.
	After successful completion of this course, student will be able to:
	1. Understand various lifetime probability distributions and
Course	their structural properties
Outcomos	2. Understand different methods for the estimation of
Outcomes	survival function.
	3. Conduct analysis of life time data.
	4. Apply statistical techniques to model lifetime data and make predictions.

• Lecturing, Hands on Training, Visualization, Team Learning

# **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

# **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous E	valuation
Tests	16
Assignment	08
Seminar	16
Total	40

### Sample Questions to Test Outcomes:

- 1. Define survival function and hazard function.
- 2. Explain aging classes and their dual.
- 3. What do you mean by bathtub failure rate?
- 4. Explain any three life distributions.
- 5. What is goodness of fit test? Explain any two goodness of fit test.
- 6. Define censoring. Elaborate various types of censoring.
- 7. Explain truncation. Differentiate between censoring and truncation.
- 8. Explain Kaplan-Meier estimation technique.

### POOL B: DISCIPLINE SPECIFIC ELECTIVE COURSE (Practical)

Course Code &	MSBST02DSE03- BIOSTATISTICAL COMPUTING USING SPSS –		
Title	II (Practical)		
Programme	M.Sc. Biostatistics	Semester	II
	• To introduce some advanced statistical computing techniques in		ng techniques in
Course	applied statistics to extract information and visualization thereby		
Objectives	enabling them to perform data analysis effectively and efficiently		
	in SPSS.		
	• Illustrate different statistical techniques based on all the elective		
	course in second semester.		

Modules	Content	Module Outcome
	Statistical Computing II is a	• Describe different statistical
- 7 N	practical course. The practical	technique to solve problems
/ /	is based on all the elective	coming under all the elective
V	courses in the second semester.	courses in second semester.
Course	After successful completion of this course, student will be able to:	
Outcomes	1. Equipped with different theoretical methods in biostatistics to	
	achieve the objectives.	
	2. Enhanced with the basic concepts of biostatistical theories besides	
	developing their ability to handle real world problems with large	
	scale data.	

- Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION
  - Lecture, Seminar, Hands on training

### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation	60
Continuous Eval	uation
Practical Tests	32
Record	08
Total	40

### **POOL B: DISCIPLINE SPECIFIC ELECTIVE COURSE (Practical)**

Course Code &	MSBST02DSE04- BIOSTATISTICAL COMPUTING USING SAS –		-
Title	I (Practical)		
Programme	M.Sc. Biostatistics	Semester II	
	• To introduce some advanced statistical computing techniques in		n
Course	applied statistics to extract information and visualization thereby		y
Objectives	enabling them to perform data analysis effectively and efficiently		
- A.	in SAS programming.		
	• Illustrate different statistical techniques based on all the elective		
<1/	course in second semester.		
K	UDININE		

Modules	Content	Module Outcome
	Statistical Computing II is a	• Describe different statistical
	practical course. The practical is	technique to solve problems
	based on all the elective courses	coming under all the elective
	in the second semester.	courses in second semester.

	After successful completion of this course, student will be able to:	
	1.Equipped with different theoretical methods in biostatistics to	
Course	achieve the objectives.	
Outcomes	2. Enhanced with the basic concepts of biostatistical theories besides	
	developing their ability to handle real world problems with large	
	scale data.	

- Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION
  - Lecture, Seminar, Hands on training

# ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation	60
Continuous Eva	luation
Practical Tests	32
Record	08
Total	40

POOL C:- List of Ability Enhancement Courses (Offered to other Departments)

ABILITY ENHANCEMENT COURSE (AEC)

Course code: MSBST02AEC01

Name of the Course: A foundation course in LaTex for scientific documentation.

Department Offering the Course: Department of Statistical Sciences

Mode of Delivery: Hybrid

# Credit Distribution, Eligibility and Pre-Requisites of the Course

Credits	Contact hours per week				Pre-requisite			
	Lecture	Tutorial	Practical/	Eligibility Criteria	(if any)			
	Lecture	Tutoriai	Internship		(II ally)			
2	1	1	2					

**Skill Outcomes:** To introduce students with a software that is being widely used for scientific typesetting, To make students know importance of this software for publishing research articles, letters, project reports, books and beamer/slide presentation and thereby help them to be comfortable with the software .

### **Course Contents:**

**Module 1:** Installation of Kile and MikeTeX. Class and packages. Latex programming and commands, sample packages. Error messages, Some sample errors, list of LaTeX error messages.

**Module 2:** Fonts, symbols, Indenting, paragraphs, line spacing, word spacing, titles and subtitles. Document class, page style, parts of the documents, table of contents. Command names and arguments, environments, declarations. Theorem like declarations, comments within text.

**Module 3**: Mathematical environments, math mode, mathematical symbols. Graphic package, multivalued functions, drawing matrices. Tables, tables with captions. References to figures and tables in text.

**Module 4:** Picture environments. Extended pictures, other drawing packages. Preparing book, project report in LaTeX, LaTeX Beamer for Technical Presentations.

### **Suggested Readings:**

- Kottwitz, S. (2021). LaTeX Beginner's Guide: Create Visually Appealing Texts, Articles, and Books for Business and Science Using LaTeX. United Kingdom: Packt Publishing.
- 2. Lamport (1994). Latex: A Document Preparation System, 2/E. India: Pearson Education.

3. Kopka, H., Daly, P. W. (2003). *Guide to LaTeX*. United Kingdom: Pearson Education. **TEACHING LEARNING STRATEGIES** 

• Hands on training, Lecturing, Visualization, Team Learning.

### **MODE OF TRANSACTION**

• Lab session, Lecture, Seminar, Discussion, Questioning and Answering

### Assessment Rubrics: Evaluation by Department

### Sample Questions to Test Outcomes:

- 1. How do you install Kile and MiKTeX for LaTeX editing and compilation, and what are their roles in the LaTeX ecosystem?
- 2. Explain the concept of classes and packages in LaTeX, and how they contribute to document formatting and customization.
- 3. Discuss LaTeX programming and commands, including sample packages, and common error messages encountered during compilation.
- 4. What are some sample errors in LaTeX, and how can they be addressed? Provide a list of common LaTeX error messages.
- 5. Describe the role of fonts and symbols in LaTeX, and how they can be customized for document appearance.
- 6. How do you manage indentation, paragraphs, line spacing, and word spacing in LaTeX documents?
- 7. Discuss titles, subtitles, document classes, page styles, and the creation of table of contents in LaTeX.
- 8. Explain command names and arguments, environments, and declarations in LaTeX, including theorem-like declarations and comments within text.
- 9. What are mathematical environments and math mode in LaTeX, and how can mathematical symbols be utilized?
- 10. Describe the usage of the graphic package in LaTeX, including drawing matrices, creating tables with captions, and referencing figures and tables within text.

# POOL C: ABILITY ENHANCEMENT COURSE -II

Course code: MSBST02AEC02

Name of the Course: Basic Statistical data analysis using EXCEL.

Department Offering the Course: Department of Statistical Sciences

Mode of Delivery: Hybrid

### Credit Distribution, Eligibility and Pre-Requisites of the Course

Credits	Contact hours per week	Eligibility	Pre-requisite
---------	------------------------	-------------	---------------

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	Lecture	Tutorial	Practical/ Internship	Criteria	(if any)
2	1	1	2		Basic knowledge of statistics

#### **Skill Outcomes:**

To build a strong understanding on the Basics of Microsoft Excel, To understand data crunching, Understand core analytic techniques that work in Excel, Data visualization in Excel.

### **Course Contents:**

**Module 1**: Excel Introduction, Basic Navigation Tab, Concept of Cell and Cell address, row Column concept, Basic mathematical and statistical functions in Excel.

**Module 2:** Min, Max, Trim, Lower, Upper, Proper, Left, Right, Mid Exact, Randbetween, Rand, Len (Length of character) Paste special, SQRT, If function with Example of IF, More function like And, OR with their example, Conditional Formatting basic and advance level with OR, AND, Nested IF function, Index, Offset, Match.

**Module 3**: Graphics in excel-pie chart, bar chart, multiple bar diagram, sub-divided bar diagram, histogram, line chart, scatter diagram, box plot.

**Module 4**: Median, Mode, Standard Deviation (SD), Correlation, Large, Small, Pivot Table, Pivot Charts, Slicing, Sparkling.

### **Suggested Readings:**

- 1. Linoff, Gordon S (2015). Data analysis using SQL and Excel. John Wiley & Sons.
- Guerrero, Hector, Rauscher Guerrero, and Rauscher (2019). Excel data analysis. Springer International Publishing.

### **TEACHING LEARNING STRATEGIES**

• Hands on training, Lecturing, Visualization, Team Learning.

### **MODE OF TRANSACTION**

• Lab session, Lecture, Seminar, Discussion, Questioning and Answering

### Assessment Rubrics: Evaluation by Department

### Sample Questions to Test Outcomes:

- 1. What is the purpose of Excel and how is it commonly used in data management and analysis?
- 2. Describe the basic navigation tab in Excel and explain the concept of cells, cell addresses, rows, and columns.
- 3. Discuss basic mathematical and statistical functions in Excel, including examples of their usage.
- 4. Explain the functions MIN, MAX, TRIM, LOWER, UPPER, PROPER, LEFT, RIGHT, MID, EXACT, RANDBETWEEN, RAND, and LEN, and how they are used in Excel.
- 5. How does the Paste Special function work in Excel, and what are its common applications?
- 6. Walk through the usage of the SQRT function and demonstrate its application in Excel.
- 7. Provide examples of the IF function in Excel and discuss its importance in conditional logic.
- 8. Explain the functions AND and OR in Excel with examples, and discuss their usage in conjunction with conditional formatting.
- 9. What are nested IF functions in Excel, and how are they implemented? Provide examples.
- 10. Discuss advanced functions such as INDEX, OFFSET, MATCH, and their applications in Excel data analysis and manipulation.
- 11.

# POOL D:- List of Skill Enhancement Courses (Offered to other Departments) SKILL ENHANCEMENT COURSE (SEC)-I

Course code: MSBST02SEC01

Name of the Course: Exploratory Data Analysis Using SPSS

Department Offering the Course: Department of Statistical Sciences

Mode of Delivery: Hybrid

Credit Distribution, Eligibility and Pre-Requisites of the Course

Credits Lec	Cont	tact hours	per week		Pre-requisite (if any)		
	Lecture	Tutorial	Practical/	Eligibility Criteria			
	Lecture	Tutoriai	Internship				
					Knowledge of basic		
2	1 1		2	0.001.0	statistics		

**Skill Outcomes:** To introduce students with a software that is being widely used for Statistical data analysis. To make students know importance of this software for data analysis in research articles and thereby help them to be comfortable with the software.

### **Course Contents:**

**Module 1:** What is SPSS?, Opening SPSS, Layout of SPSS, Structure of SPSS Exiting SPSS, inputting data, An overview of SPSS.

**Module 2:** Exploring data distributions using descriptive statistics, Creating frequency distributions and summary tables, Generating basic visualizations (e.g., histograms, box plots) in SPSS.

**Module 3**: Understanding correlation and covariance, Performing correlation analysis in SPSS, Introduction to linear regression and its application in SPSS, Understanding hypothesis testing principles, Conducting hypothesis tests in SPSS, Interpreting SPSS output for hypothesis testing.

**Module 4: Generating** various types of charts and graphs in SPSS, Customizing visualizations for clarity and impact, exploring the SPSS Chart Builder tool.

### Suggested Readings:

- 1. Landau, S., & Everitt, B. S. (2003). *A handbook of statistical analyses using SPSS*. Chapman and Hall/CRC.
- 2. Tukey, J. W. (1977). Exploratory data analysis (Vol. 2).
- 3. Aldrich, J. O. (2018). Using IBM SPSS Statistics: An interactive hands-on approach. Sage Publications.

### **TEACHING LEARNING STRATEGIES**

• Hands on training, Lecturing, Visualization, Team Learning.

### **MODE OF TRANSACTION**

• Lab session, Lecture, Seminar, Discussion, Questioning and Answering

# Assessment Rubrics: Evaluation by Department Sample Questions to Test Outcomes:

- 1. What is the purpose of SPSS and how does it contribute to statistical analysis?
- 2. Describe the layout and structure of SPSS interface, including its main components and functions.
- 3. How do you open and exit SPSS, and what considerations should be taken into account?
- 4. Explain the process of inputting data into SPSS and discuss common formats accepted.
- 5. What are descriptive statistics, and how can they be used to explore data distributions in SPSS?
- 6. Describe the steps involved in creating frequency distributions and summary tables using SPSS.
- 7. How can basic visualizations such as histograms and box plots be generated in SPSS, and what insights can they provide?
- 8. What are correlation and covariance, and how are they calculated and interpreted in SPSS?
- 9. Discuss the principles of hypothesis testing and how it is conducted in SPSS.
- 10. Explain how to customize visualizations for clarity and impact in SPSS, and explore the functionalities of the SPSS Chart Builder tool.

# POOL D: SKILL ENHANCEMENT COURSE (SEC)-II

Course code: MSBST02SEC02

Name of the Course: Regression Analysis Using SPSS

Department Offering the Course: Department of Statistical Sciences

Mode of Delivery: Hybrid

# Credit Distribution, Eligibility and Pre-Requisites of the Course

Credits	Contact hours per week				Pre-requisite		
	Lecture	Tutorial	Practical/	Eligibility Criteria	(if any)		
			Internship		(II ally)		

				Basic knowledge in
2	1	1	2	regression.

**Skill Outcomes:** To introduce students with software that is being widely used for regression analysis. To make students know importance of this software for data analysis in research articles and thereby help them to be comfortable with the software.

#### **Course Contents:**

**Module 1:** What is SPSS?Opening SPSS, Layout of SPSS, Structure of SPSS Exiting SPSS, inputting data, an overview of SPSS.

**Module 2:**Introduction to SPSS regression procedures, Importing and preparing data for regression analysis, Overview of the SPSS regression dialog box, Conducting simple linear regression in SPSS, Interpreting regression output in SPSS, Assumptions and diagnostics in simple linear regression

**Module 3**: Understanding and formulating multiple linear regression models, Conducting multiple linear regression analysis in SPSS, Interpreting output and assessing model fit

**Module 4:** Assumption testing for multiple regression, Dealing with multicollinearity in SPSS. Interpreting diagnostic plots and statistic, Organizing and documenting regression analysis in SPSS, Creating comprehensive reports with SPSS output, Best practices for presenting regression findings to diverse audiences

### **Suggested Readings:**

- 1. Landau, S., & Everitt, B. S. (2003). *A handbook of statistical analyses using SPSS*. Chapman and Hall/CRC.
- **2.** Aldrich, J. O. (2018). Using IBM SPSS Statistics: An interactive hands-on approach. Sage Publications.
- **3.** Chatterjee, S., & Hadi, A. S. (2013). *Regression analysis by example*. John Wiley & Sons.

### TEACHING LEARNING STRATEGIES

- Hands on training, Lecturing, Visualization, Team Learning. MODE OF TRANSACTION
- Lab session, Lecture, Seminar, Discussion, Questioning and Answering Assessment Rubrics: Evaluation by Department

### Sample Questions to Test Outcomes:

- 1. What are the key components of SPSS and how do they contribute to statistical analysis?
- 2. How do you input data into SPSS and what are the common formats accepted?
- 3. Describe the process of exiting SPSS and any considerations to keep in mind.
- 4. What are the steps involved in importing and preparing data for regression analysis in SPSS?
- 5. Explain the regression dialog box in SPSS and its various options for analysis.
- 6. Walk through the process of conducting simple linear regression in SPSS, including data input and interpretation of results.
- 7. What are the assumptions and diagnostics involved in simple linear regression analysis, and how can they be assessed in SPSS?



		THIRD SE	MEST	ER					
SI No	Course Code	Title of Paper	C Hot	Contact urs/We	ek	Marks			
			L	T/S	Р	ESE	CE	Total	Credits
	DIS	SCIPLINE SPECIFIC CORI	Ε COL	JRSES	(DSC	C)		1	I
3.1	MSBST03DSC09	Design of Experiments and Quality Control	4	a start	ý	60	40	100	4
3.2	MSBST03DSC10	Analysis of Clinical Trials	4	1		60	40	100	4
	]	DISCIPLINE SPECIFIC EL	ECTIV	VES (D	SE)				
3.3	MSBST03DSExx	(One course has to be chosen from <b>Pool E</b> )	3	2		60	40	100	3
3.4	MSBST03DSExx	Elective-II (DSE) (Any two courses have to be chosen from <b>Pool F</b> )	3	2		60	40	100	3
	INTE	RDISCIPLINARY ELECT	IVE CO	OURSI	E (ID)	C)*			·
3.5	MSBST03IDCxx	IDC Elective (One course has to be chosen from <b>Pool G</b> ) (Offered to other Departments)	2	2	4	60	40	100	4
		To be obtained from other departments	-	/	4	A			
*Note: C	compulsory course	1	r .,	1	21	0	~		1
	-	VAC/MOOC COU	RSE**	*	/	1	~		
	моос	Offered by external agencies	E	~		60	40	100	At least 2
3.6	MSBST03VACxx	VAC (To be decided by Department as per requirements)	2	2	2	60	40	100	2
* C	*Note: The course shall GPA.	be considered as additional cre	dits and	d shall :	not be	e consid	ered fo	br compt	itation of

	INTERSHIP/FIELD VISIT/ MINOR PROJECT/ INDUSTRIAL VISIT								
3.7	MSBST03DSC11	Any one of: internship, field visit, minor project or industrial visit.	2	-	-	60	40	100	2
	Tota	(D)200 0	19		1				23

L=Lecture, T/S=Tutorials/Seminar, P/I=Practical/Internship, CE =Continuous Evaluation, ESE = End Semester Evaluation

Sl No	POO	OL E: DISCIPLINE SPECIFIC	CELE	CTIVE	ES (DS)	E) (Pra	actical	l)	
3.3.1	MSBST03DSE05	Biostatistical Computing Using R -II		2	6	60	40	100	3
3.3.2	MSBST03DSE06	Biostatistical Computing Using SAS -II		2	6	60	40	100	3
		POOL F : DISCIPLINE SPE	CIFIC	ELEC	CTIVE	S (DSI	E)		
3.4.1	MSBST03DSE07	Stochastic Process and Time Series Analysis	4	2	-	60	40	100	3
3.4.2	MSBST03DSE08	Applied Multivariate Analysis	4	2	-	60	40	100	3
3.4.3	MSBST03DSE09	Machine Learning Techniques for Biostatistics	4	2	~	60	40	100	3
3.4.4	MSBST03DSE10	Categorical Data Analysis	4	2	5	60	40	100	3
3.4.5	MSBST03DSE11	Operations Research	4	1	52	60	40	100	3
3.4.6	MSBST03DSE12	Data Visualisation and analysis using Python	2	ľ	4	60	40	100	3

Sl No	POOL G:	POOL G: INTERDISCIPLINARY ELECTIVES (for other Departments)(IDC)							
3.5.1	MSBST03IDC01	Statistical Data Analysis using SPSS		2	6	60	40	100	3
3.5.2	MSBST03IDC02	Statistical Data Analysis using R		2	6	60	40	100	3

	DISCIPLINE SPECIFIC CO	DRE COURSE							
Course Code &	MSBST03DSC09- DESIGN OF	EXPERIMENTS AND QUALITY							
Title	CONTROL								
Programme	M.Sc. Biostatistics	Semester III							
Course Objectives	<ul> <li>This course provides the study conduct experiments, as we successful completion of this</li> <li>Apply ANOVA for one way models with equal and un Random and Mixed effect models</li> <li>Design and analyse incomplication of orthogonality, control of the effects of differ analyse factorial experiments.</li> <li>Understand basics of static charts.</li> </ul>	ents the ability to understand the design and ll as to analyze and interpret data. After course, student will be able to: and two-way classification, fixed effect equal number of observations percell, odels. lete block designs, understand the nnectedness and balance. ent factors and their interactions and istical quality control and various control							

D				
Module I Basic concepts of design of experiments (15 Hours)	Basic terminology efinitions, Randomiz deplication and local co vixed, mixed and random nodels, Gauss Markov the undamental principles of f experiments, Analys ariance-one way and two	and zation, ontrol, effect eorem, design is of ways.	<ul> <li>Understand concepts of experiments.</li> <li>Differentiate mixed, and models.</li> <li>Explain the theorem.</li> <li>Perform analy for one-way</li> </ul>	the basic design of between fixed, random effect Gauss-Markov ysis of variance and two-way

	Completely randomized design	• Design and analyze CRD,
	analysis, randomized block	RBD, LSD.
Module II	design-analysis, Latin square	• Design and analyze
Randomized	design-analysis, Graeco-Latin	experiments using Graeco-Latin
designs	square designs, analysis of	square designs.
(15 Hours)	missing data. Analysis of	• Understand methods for
	covariance for RBD.	analyzing and handling missing
		data.
	Incomplete block design-	• Design and analyze
	Balanced incomplete block	experiments using incomplete
Madula III	design, construction of BIBD	block designs.
Incomplete	design, intra block analysis of	• Perform intra block analyses
block and	BIBD, Factorialexperiments-2 <sup>n</sup> ,	for balanced incomplete block
factorial	concept of confounding.	designs.
designs		• Design and analyze factorial
(15 Hours)		experiments with 2 <sup>n</sup> levels.
(15 110013)		• Identify and understand the
		presence of confounding in
		factorial experiments.
	Quality and quality assurance,	1
	Methods of quality assurance,	• Define and differentiate between
- A	statistical quality control	quality and quality assurance
Module IV	Control charts, Basic ideas,	• Design and interpret mean charts,
Introduction to	designing of control charts for	median charts, range (R)-charts,
statistical	the number of non-conformities	and standard deviation (S)-charts
quality control	and fraction non-conformities,	for process monitoring.
(15 Hours)	mean charts, Median charts, R-	• Understand the economic
	charts, and S-charts, ARL,	implications of implementing
	Economic design of Shewarts	Shewhart's control charts.
	control charts.	
References	Text Books	
initionety	1. Das, M.N. and Giri, N.S. (2	2002): Design and Analysis of

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	Experiments 2nd Edition New Age International (D) Itd New		
	Experiments, 2 Edition, New Age international (1) Etd., New		
	Delhi.		
	2. Joshi,D.D.(1987):Linear Estimation and Design of Experiments.		
	Wiley Eastern Ltd., New Delhi.		
	3. Montgomery, D.C. (2001): Design and Analysis of Experiments. 5th		
	Edition, John Wiley & Sons- New York.		
	4. Montgomery, R.C. (1985). Introduction to Statistical Quality		
	Control, Fourth edition, Wiley.		
	Reference books		
	1. Gupta, S. C and Kapoor, V. K.(2010). Fundamentals of Applied		
	Statistics. Sulthan Chand & Co, NewDelhi.		
	2. Amitava Mitra - Fundamentals of Quality Control and Improvement –		
	Pearson Education Asia 2001.		
	3. The ISO 9000 book, Second Edition, Rabbit, J T and Bergle, PA Quality		
	resources.		
	After successful completion of this course, student will be able to:		
	1 Demonstrate a mastery of fundamental principles guiding the		
	design of experiments.		
	2 Able to design and analyze experiments using Latin square and		
	Grace Latin square designs		
Course	Diacco-Latin square designs.		
Course	3. Demonstrate expertise in designing and analyzing factorial		
Outcomes	experiments with 2 <sup>n</sup> levels, and will be able to identify and control		
<1/	confounding factors within factorial designs to ensure the accuracy		
1	of results.		
	4. Identify different types of control charts and their applications in		
	monitoring processes. Optimize the design of control charts to		
	minimize costs while ensuring effective quality control.		

### • Lecturing, Visualization, Team Learning

# **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

### **ASSESSMENT RUBRICS**

Components	Weightage	
End Semester Evaluation(ESE)	60	
Continuous Ev	aluation	
Tests	16	
Assignment	08	
Seminar	16	
Total	40	

### Sample Questions to Test Outcomes:

- 1. Explain Gauss-Markov theorem.
- 2. Explain Cochran's theorem.
- 3. Describe the three fundamental principles of experimentation and explain the importance of these principles with respect to designing statistical experiments.
- 4. Briefly explain contrasts and orthogonal contrasts.
- 5. Explain in detail the analysis of two-way classified data with one observation per cell.
- 6. Explain the efficiency of RBD relating to CRD.
- 7. Distinguish between ANOVA and ANCOVA
- 8. What do you understand by "Analysis of Covariance"? Illustrate with suitable example.
- 9. Explain Greaco Latin Square Design and orthogonal latin square design.
- 10. In a LSD a single observation is missing. How will you estimate the missing value and carry out the analysis of the design?

Course Code & Title	MSBST03DSC10- ANALYSIS OF CLINICAL TRIALS	
Programme	M.Sc. Biostatistics Semester III	

### **DISCIPLINE SPECIFIC CORE COURSE**

	The objective of this course is to study more advanced topics in design and	
Course	analysis of clinical trials. After successful completion of this course, student	
Objectives will be able to:		
Objectives	1. Understand Basics of Clinical Trails	
	2. Understand design of clinical trials	
	3. Understand Sample size determination in clinical trials	
	4. Understand the concept of meta analysis in clinical trials.	

Modules	Content	Module Outcome
Module I Introduction to clinical trails (15 Hours)	Introduction to clinical trails, the need and ethics of clinical trials, bias and random error in clinical studies, Protocols, conduct of clinical trials, over view of Phase I-IV trials, Data management-data definitions, standard operating procedure, informed consent form, case report forms, database design, data collection systems for good clinical practice.	<ul> <li>Understand the need and ethics of clinical trial.</li> <li>Describe different types of forms used in clinical trial.</li> <li>Understand about different types of errors that occur during the conduct of clinical trial.</li> <li>Have a clear idea on clinical practice and clinical practice and management.</li> </ul>
Module II Design of clinical trials (15 Hours)	Design of clinical trials- Different phases, Comparative and controlled trials, Random allocation, Randomization, response adaptive methods and restricted randomization. Methods of Blinding, Parallel group designs, Crossover designs, Symmetric designs, Adaptive designs, Group sequential designs, Zelen's designs, design of bioequivalence trials. Outcome measures.	<ul> <li>Understand about different types of randomization procedures used in clinical trials.</li> <li>Have an idea on different types of bliniding.</li> <li>Describe different types of designs used in clinical trials.</li> </ul>
Module III		• Determine the sample

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Statisticalanalysisinclinical trails(15 Hours)	<ul> <li>Sample size determination in one and two sample cases, comparative trials, activity studies, testing and other purposes, unequal sample sizes and case of ANOVA. Surrogate endpoints-selection and design of trials with surrogate, analysis of surrogate end point data. Reporting and analysis-Interpretation of result, multi-center trials.</li> <li>Sample size determination in one and size for different types of response variables.</li> <li>Describe multi-centre trials.</li> <li>Have a clear idea on reporting and interpreting of results in a clinical trial.</li> <li>Describe surrogate endpoints.</li> </ul>
Module IV Meta analysis in clinical trials (15 Hours)	<ul> <li>Meta analysis in clinical trials-concept and goals, fixed and random effect approaches.</li> <li>Bioassay: Direct and indirect assays, Quantal and quantitative assays, Parallel line and slope ratio assays, Design of bioassays.</li> <li>Understand different types of bioassay and their analysis.</li> <li>Describe meta-analysis.</li> </ul>
References	<ol> <li>Text Books         <ol> <li>Chen, D.G. and Peace, K.E. (2011). Clinical Trial Data Analysis Using R. Chapman &amp; Hall</li> <li>Friedman, L. M., Furburg, C. D. Demets, L. (1998): Fundamentals of Clinical Trials, Springer Verlag.</li> <li>Kulinskaya E, Morgeathaler S, Staudte R G(2008). Meta analysis, Wiley.</li> </ol> </li> <li>Reference Books</li> </ol>
	<ol> <li>Das, M. N. and Giri(2008). <i>Design of Experiments</i>, New Age, India</li> <li>Jennison and B.W. Turnbull (1999): <i>Group Sequential Methods</i> with Applications to Clinical Trials, CRC Press.</li> </ol>
Course	<ol> <li>Das, M. N. and Giri(2008). Design of Experiments, New Age, India</li> <li>Jennison and B.W. Turnbull (1999): Group Sequential Methods with Applications to Clinical Trials, CRC Press.</li> <li>After successful completion of this course, student will be able to:</li> </ol>
Course Outcomes	<ol> <li>Das, M. N. and Giri(2008). Design of Experiments, New Age, India</li> <li>Jennison and B.W. Turnbull (1999): Group Sequential Methods with Applications to Clinical Trials, CRC Press.</li> <li>After successful completion of this course, student will be able to:         <ol> <li>Understand Basics of Clinical Trials.</li> </ol> </li> </ol>
Course Outcomes	<ol> <li>Das, M. N. and Giri(2008). Design of Experiments, New Age, India</li> <li>Jennison and B.W. Turnbull (1999): Group Sequential Methods with Applications to Clinical Trials, CRC Press.</li> <li>After successful completion of this course, student will be able to:         <ol> <li>Understand Basics of Clinical Trials.</li> <li>Understand design of clinical trials.</li> </ol> </li> </ol>
Course Outcomes	<ol> <li>Das, M. N. and Giri(2008). Design of Experiments, New Age, India</li> <li>Jennison and B.W. Turnbull (1999): Group Sequential Methods with Applications to Clinical Trials, CRC Press.</li> <li>After successful completion of this course, student will be able to:         <ol> <li>Understand Basics of Clinical Trials.</li> <li>Understand design of clinical trials.</li> <li>Understand Sample size determination in clinical trials.</li> </ol> </li> </ol>

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

# ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Eva	aluation
Tests	16
Assignment	08
Seminar	16

### Sample Questions to Test Outcomes:

- 1. What is a clinical trial?
- 2. Differentiate between a prospective an retrospective study.
- 3. Define response adaptive randomization.
- 4. Distinguish between blinding and masking
- 5. Define sample size calculation for independent continuous response variables.
- 6. Differentiate between analytical dilution assay and comparative dilution assay.
- 7. Distinguish between adaptive and non-adaptive randomization.
- 8. Explain about blocked randomization and stratified randomization.
- 9. Explain in detail about blindness in a clinical trial.
- 10. Describe the design of adaptive trials and group sequential design.

# POOL E: DISCIPLINE SPECIFIC ELECTIVE COURSE (Practical)

Course Code &	MSBST03DSE05- BIOSTATISTICAL COMPUTING USING R – II		
Title	(Practical)		
Programme	M.Sc. Biostatistics	Semester	III

	• To introduce some advanced statistical computing techniques in
Course	applied statistics to extract information and visualization thereby
Objectives	enabling them to perform data analysis effectively and efficiently
	in R programming.
	• Illustrate different statistical techniques based on all the elective
	course in third semester.

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Modules	Content	Module Outcome
	Statistical Computing III is a	• Describe different
	practical course. The practical is	statistical technique to
	based on all the elective courses in	solve problems coming
	the third semester.	under all the elective
		courses in third semester.
Course	After successful completion of this course,	student will be able to:
Outcomes	3. Equipped with different theoretical mathematical mathe	ethods in biostatistics to
	achieve the objectives.	
	4. Enhanced with the basic concepts of	biostatistical theories besides
	developing their ability to handle real	world problems with large
	scale data.	

• Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Hands on training

### ASSESSMENT RUBRICS

Components	Weightage		
End Semester Evaluation	60		
Continuous Evaluation			
Practical Tests	32		
Record	8		
Total	40		

### **POOL E: DISCIPLINE SPECIFIC ELECTIVE COURSE (Practical)**

Course Code & Title	MSBST03DSE06-BIOSTATISTICAL COMPUTING USING SAS-II (PRACTICAL)
Programme	M.Sc. Biostatistics Semester III
Course	• To introduce some advanced biostatistical computing techniques
Objectives	in applied statistics to extract information and visualization
	thereby enabling them to perform data analysis effectively and
	efficiently in SAS programming.
	• Illustrate different biostatistical techniques based on all the
	elective course in third semester.

Modules	Content	Module Outcome
	Biostatistical Computing III is a	• Describe different
	practical course. The practical is	statistical technique to
	based on all the elective courses in	solve problems coming
	the third semester.	under all the elective
		courses in third semester.
Course	After successful completion of this course,	student will be able to:
Outcomes	1.Equipped with different theoretical method	ls in biostatistics to achieve
5	the objectives.	1
- A.	2. Enhanced with the basic concepts of biost	atistical theories besides
	developing their ability to handle real worl	d problems with large scale
<11	data.	51/2
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# **TEACHING LEARNING STRATEGIES**

- Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION
  - Lecture, Seminar, Hands on training

# ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation	60
Continuous Eva	luation
Practical Tests	32
Record	8
Total	40

# POOL F: DISCIPLINE SPECIFIC ELECTIVE COURSE

Course Code &	MSBST03DSE07-STOCHASTIC PROCESS AND TIME SERIES	
Title	ANALYSIS	
Programme	M.Sc. Biostatistics	Semester III
	The course will help the Students to develop a comprehensive	
Course	understanding of stochastic processes, particularly focusing on Markov	
Objectives	chains, including their definition, classification, and real-world examples.	
	It also help the student to delve in to basic concepts of time series analysis	
	and time series modelling. Also Students will develop skills in time series	
	analysis, including autocorrelation, stationarity, and regression modeling.	

Modules	Content	Module Outcome
Module I: Introduction to stochastic process (16 Hours)	Introduction to stochastic process, Markov Chains: Definition, Examples and classification, Discrete renewal equation and basic limit theorem, Absorption probabilities, Criteria for recurrence.	<ul> <li>Define Markov chains and classify them based on various criteria.</li> <li>Understand the discrete renewal equation and its significance in the context of Markov chains.</li> <li>Calculate absorption probabilities for absorbing</li> </ul>

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		<ul> <li>Markov chains.</li> <li>Understand the implications of recurrence on the long-term behavior of Markov chains.</li> </ul>
Module II: Continuous time Markov chains (14 Hours)	Continuous time Markov chains, Examples, General pure birth process, Poisson process, Birth and death process, Finite state continuous time Markov chains, Applications to queuing models.	<ul> <li>Provide examples of continuous-time Markov chains in various domains.</li> <li>Understand the concept of pure birth processes and Poisson processes.</li> <li>Analyze birth and death processes in continuous time Markov chains.</li> <li>Analyze finite-state continuous-time Markov chains using mathematical tools.</li> </ul>
Module III: Introduction to time series analysis (16 Hours)	Characteristics of time series: Time series as a discrete parameter stochastic process, Autocorrelation (ACF) and cross correlations, Stationary time series, Estimation of autocorrelations. Classical regression in time series context, exploratory data analysis, smoothing methods for time series. Wold representation of linear stationary processes.	<ul> <li>Understand time series as a discrete parameter stochastic process.</li> <li>Calculate autocorrelation functions (ACF) and cross-correlations for time series data.</li> <li>Define stationary time series and understand its importance in time series analysis.</li> <li>Estimate autocorrelations and conduct classical regression analysis for time series data.</li> </ul>
Module IV: Linear time series model (14 Hours)	Linear time series models : Autoregressive (AR), Moving Average (MA), Autoregressive Moving Average (ARMA) and	• Understand the concepts and properties of autoregressive (AR) and moving average (MA) models.

| Scheme and Syllabus of M Sc. Biostatistics- 2023 Admission onwards- Kannur University

	<ul> <li>Autoregressive Integrated</li> <li>Moving Average (ARIMA)</li> <li>models. Forecasting and</li> <li>estimation of ARMA models.</li> <li>Seasonal ARIMA models,</li> <li>Residual analysis and diagnostic</li> <li>checking.</li> <li>• Understand the structure and</li> <li>parameters of autoregressive</li> <li>moving average (ARMA) models.</li> <li>• Apply ARIMA models to</li> <li>analyze and forecast time series</li> <li>data with trend and seasonality.</li> <li>• Conduct residual analysis and</li> <li>diagnostic checking to assess the</li> <li>adequacy of time series models.</li> </ul>
	Text Books 1. Karlin.S. and Taylor, H.M. (1975) A First Course in Stochastic
Processes, second edition, Academic Press.	
	2. Bhat, B.R. (2002) Stochastic Processes, second edition, New Age
	Publication.
	3. Shumway, R. H and Stoffer, D. S. (2006). Time series Analysis and its
	Applications. Springer.
	4. Box, G. E. P. Jenkins, G. M. and Reinsel, G. C. (1994). Time Series
	Analysis: Forecasting and Control, Pearson Education.
References	Reference Books
	1. Feller, W. (1965, 1968), An Introduction to Probability Theory and
its Applications, Volume I and II, Wiley	its Applications, Volume I and II, Wiley Eastern.
A	2. Bhat, U.N. (1984) Elements of Applied Stochastic Processes, John
	Wiley.
<1/	3. Cinlar, E. (1975) Introduction to Stochastic Processes, Prentice Hall.
distant of the second s	4. Brockwell, P.J and Davis R.A. (2006) Time Series: Theory and
	Methods, 2ndedn. Springer-Verlag
	5. Chatfield, C. (2004) The Analysis of Time Series - An Introduction,
	Sixth edition, Chapman and Hall.
	6. Anderson, T.W (1971) Statistical Analysis of Time Series, Wiley.
Course	After successful completion of this course, student will be able to:
Outcomes	1. Understand the implications of recurrence on the long-term behavior of
	Warkov chains.
2. Determine key characteristics and parameters of birth and death	
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processes.	
3. Apply AR and MA models to analyze and forecast time series data.	
4. Apply ARIMA models to analyze and forecast time series data with trend	
and seasonality.	

• Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

#### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Ev	aluation
Tests	16
Assignment	08
Seminar	16
Total	40

### Sample Questions to Test Outcomes:

- 1. Define Markov chain.
- 2. Prove that a Markov chain is completely determined by its initial distribution and one step TPM.
- 3. Define periodic and aperiodic Markov chains.
- 4. Consider Markov chain whose TPM is

0	$\frac{1}{3}$	$\frac{2}{3}$
$\frac{1}{2}$	0	$\frac{1}{2}$
$\frac{1}{2}$	$\frac{1}{2}$	0)

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(i)Is the chain is irreducible.

(ii)Is the chain is ergodic? Explain.

(iii)Find the stationary distribution of the chain.

- 5. Show that in an irreducible Markov chain all the states are of same type.
- 6. Show that probability of extinction of Galton Watson branching process is the smallest positive root of the equation s = P(s), where P(s) is the PGF of the offspring distribution.
- Suppose that customers arrive at a bank according to a Poisson process with mean rate of 3 per minute. Find the probability that during a time interval of 2 minutes (i)exactly 4 customers arrive (ii) more than 4 customers arrive and (iii) at least one customer arrive.
- 8. Describe the relation between Poisson process and binomial distribution.
- 9. Explain renewal process. Give examples

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10. Describe the stationary behavior of a birth and death processes and hence obtain the system size distribution of M/M/1 and M/M/c queueing models.

Course Code &	MCDCT02DCE00 ADDI IED MILL	
Title	MSBST03DSE08-APPLIED MULTIVARIATE ANALYSIS	
Programme	M.Sc. Biostatistics	Semester III

## POOL F: DISCIPLINE SPECIFIC ELECTIVE COURSE



	• Understand the fundamentals of multivariate data analysis and the
	notion of multivariate distributions.
	• Demonstrate proficiency in analyzing multivariate data using the
	multivariate normal distribution, including understanding marginal
	and conditional distributions.
	• Master the concept of characteristic functions and their application in
Course	multivariate data analysis.
Objectives	• Develop skills in estimating the mean vector and covariance matrix of
	multivariate datasets.
	• Interpret canonical variates and canonical correlations, both in
	population and sample contexts, as part of Canonical Correlation
	Analysis (CCA).
	• Apply orthogonal factor models in Factor Analysis, including
	methods of estimation, factor rotation, and computation of factor
	scores.
	• Utilize various similarity measures and hierarchical and non-
	hierarchical clustering methods in Cluster Analysis.

Modules	Content	Module Outcome		
Module I Multivariate normal distribution (15 Hours)	Multivariate data, preliminary analysis, notion of multivariate distributions, multivariate normal distribution, marginal and conditional distributions, characteristic function, estimation of mean vector and covariance matrix.	<ul> <li>Able to identify and describe the notion of multivariate distributions.</li> <li>Learn about marginal and conditional distributions in the context of multivariate data.</li> <li>Derive characteristic functions and their importance in the analysis of multivariate data.</li> </ul>		
Module II Principal component and canonical correlation	Principal components Analysis: - population principal components, summarizing sample variation by principal components, graphing the principal components; Canonical correlation analysis: - canonical variates	<ul> <li>Students will demonstrate a comprehensive understanding of PCA.</li> <li>Summarize sample variation effectively using</li> </ul>		

analysis	and canonical correlations, interpreting	ns, interpreting principal components.	
(15 Hours)	the population canonical variables, the • Master the princip		
	sample canonical variates and sample	Canonical Correlation Analysis,	
	canonical correlations.		
		• Demonstrate a	
	1000 GEL2001	comprehensive understanding of	
	1000	factor analysis including the	
	10%	orthogonal factor model, methods	
Module III	Factor analysis: - orthogonal factor model;	of estimation.	
Factor and	methods of estimation, factor rotation,	• Develop proficiency in	
cluster analysis	factor scores; Cluster analysis: - similarity	applying factor analysis	
(15 Hours)	measures, hierarchical clustering methods,	techniques, including estimating	
	non-hierarchical clustering methods.	factor loadings, conducting factor	
		rotation to simplify interpretation.	
		• Demonstrate the ability to	
		apply cluster analysis techniques	
		to real-world datasets.	
Module IV:		• Demonstrate proficiency	
MANOVA and		in comparing several multivariate	
Multidimension		population means using one-way	
al scaling	Comparison of several multivariate	Multivariate Analysis of Variance	
(15 Hours)	population means (one-way MANOVA),	(MANOVA).	
A	simultaneous confidence intervals for	• Construct simultaneous	
	treatment effects, two-way multivariate	confidence intervals for treatment	
4)	analysis of variance; Distance methods: - multidimensional scaling, correspondence	effects in multivariate data	
		analysis settings	
1	analysis.	Proficiency in distance	
	- UNIT	based multiveriete enclusion	
		based multivariate analysis	
		methods.	

References	Text Books		
	1. Johnson, R.A. and Wichern, D.W. (2007) Applied Multivariate Statistical		
	Analysis, PHI Learning Private Ltd, New Delhi, Sixth edition.		
	2. Rencher, A.C. (1995) Methods of Multivariate Analysis, John Wiley.		
	3. Dillon, W.R. and Goldstein, M (1984) Multivariate Analysis, John		
	Wiley.		
	Reference Books		
	1. Anderson, T.W. (1984) An Introduction to Multivariate Statistical		
	Analysis, John Wiley.		
	2. Seber G.A.F. (1983) Multivariate Observations, Wiley.		
	3. Tabachnick, B.G. and Fidell, L.S. (2018) Using multivariate statistics,		
	Sixth edition, Pearson India Education Services Pvt Ltd, India.		
	After successful completion of this course, student will be able to:		
	1. Demonstrate a comprehensive understanding of various multivariate data		
	analysis techniques.		
	2. Develop proficiency in dimensionality reduction techniques such as Principal		
Course	Components Analysis (PCA) and Factor Analysis.		
Outcomes	3. Acquire advanced multivariate analysis skills, including conducting		
	comparisons of several multivariate population means using one-way		
	MANOVA, simultaneous confidence interval construction for treatment effects.		
5	4. Master the analysis of multivariate relationships using techniques such as		
A.,	Canonical Correlation Analysis (CCA), cluster analysis, and distance methods.		

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

### ASSESSMENT RUBRICS

Components	Weightage	
End Semester	60	
Evaluation(ESE)		
<b>Continuous Evaluation</b>		

10 W W 12	
Total	40
Seminar	16
Assignment	08
Tests	16

#### Sample Questions to Test Outcomes:

- 1. Define a singular multivariate normal distribution. Give an example of a random vector following singular multivariate normal distribution.
- 2. If  $X \sim Np(\mu, \Sigma)$  and if  $\Sigma$  is a diagonal matrix then show that the components of X are independently normally distributed and conversely.

3. Give an example to show that the marginal distributions are normals does not imply that the joint distribution follows a multivariate normal distribution.

4. Define partial correlation. Explain how do you evaluate the partial correlation coefficients from a non-singular covariance matrix.

- 5. Define multiple correlation. Show that  $0 \le \rho 1.23...p \le 1$ .
- 6. Derive the distribution of sample mean of a sample of size n from Np( $\mu$ ,  $\Sigma$ ).
- 7. Find the MLE of the correlation matrix based on a random sample of size n from  $Np(\mu, \Sigma), \Sigma > O$ , when  $\mu$  is known.
- 8. Define Wishart distribution and derive its characteristic function.
- 9. Derive the characteristic function of a matrix-variate gamma distribution.

10. Derive the characteristic function of a Wishart matrix and show that Wishart distribution is a matrix variate generalization of  $\chi^2$  distribution.

#### **POOL F: DISCIPLINE SPECIFIC ELECTIVE COURSE**

Course Code &	MSBST03DSE09 - MACHINE LEARNING TECHNIQUES FOR		
Title	BIOSTATISTICS		
Programme	M.Sc. Biostatistics	Semester	III

	• To classify variables and articulate their roles within statistical
	modeling, distinguishing between categorical, numerical, and
	ordinal variables, and understanding their implications for
	analysis.
	• Demonstrate competency in applying least squares regression and
Course	nearest neighbors algorithms for data analysis.
Course	• Master supervised learning techniques and function
Objectives	approximation methods, utilizing them to model relationships
	between input and output variables effectively.
	• Demonstrate proficiency in utilizing roughness penalty methods
	and Bayesian approaches to estimate model parameters,
	understanding their roles in controlling model complexity and
	improving generalization.
	• Proficient in employing kernel methods and local regression
	techniques for modeling non-linear relationships in data.

Modules	Content	Module Outcome
Module I Introduction and overview of supervised learning (15 Hours)	Supervised Learning and Function Approximation, A Statistical Model for the Joint Distribution of input and output vectors, Function Approximation, Structured Regression Models, Linear Methods for Regression: Least squares, Subset selection, Shrinkage Methods, Methods using derived input directions, Multiple outcome shrinkage and selection, Lasso and related path algorithms.	<ul> <li>Explore structured regression models and classes of restricted estimators, including roughness penalty and Bayesian methods.</li> <li>Understand the practical considerations in automatic selection of smoothing parameters and nonparametric logistic regression.</li> </ul>
Module II: Linear methods for	Linear methods for classification using linear regression of an indicator matrix, linear discriminant analysis, logistic regression and separating hyperplanes. Basis expansions and	• Explore linear discriminant analysis (LDA) and its role in

classification	regularizations: Piecewise polynomials and	multi-class classification
(15 Hours)	splines, Automatic Selection of the	problems.
	Smoothing Parameters, Nonparametric	• Implement piecewise
	Logistic Regression, Multidimensional	polynomials and splines
	Splines.	for capturing complex
	0000 001001	functional forms in
	1000 1 1 1 1 1	regression and
	VOT NIT I	classification tasks.
		• Explore structured local
		regression models in
		multidimensional spaces
		and their advantages in
		capturing complex data
	Selecting the band width of the Kernel	patterns.
Module III:	Structured Local Regression Models in RP	• Implement kernel
Kernal	Local Likelihood and Other Models Kernel	density classification
smoothing	Density Estimation and Classification:	methods for classifying
(15 Hours)	Kernel Density Estimation, Kernel Density	data points based on their
(10 110115)	classification and the Naïve Bayes classifier.	estimated densities
	Mixture Models for Density Estimation and	• Understand the
	Classification.	mathematical properties of
5		RBEs and kernels and
- A.		their role in non-linear
- 7 N	T	transformations of input
2.1	AN.	data
1	Bias Variance and Model Complexity. The	
Module IV:	Bias-Variance Decomposition Optimism of	• Gain proficiency in
Model	the Training Error Rate, Estimates of In-	techniques such as cross-
assessment,	Sample Prediction Error, The Bayesian	validation and bootstrap
inference and	Approach and BIC, Minimum Description	methods for model
averaging	Length, Cross-Validation, Bootstrap	evaluation and selection.
(15 Hours)	Methods, Conditional or Expected Test Error,	• Understand the
	introducing Model Inference and averaging:	principles behind the
	Local regression in IR, The EM Algorithm,	Bayesian Information

	MCMC for Sampling from the Posterior,	Criterion (BIC) and	
	Bagging, Model Averaging and Stacking,	minimum description	
	Stochastic Search: Bumping.	length for model selection.	
	Text Books		
	1. Hastie, T., Tshibirai, R. and Friedman, J. (2017) The Elements of Statistical		
-	Learning : Data Mining, Inference and Prediction, 2nd edition. Springer, New		
	York.		
Defenences	2. James, G., Witten, D., Hastie, T. and Tibshin	rani, R.(2013) An Introduction to	
Kelerences	Statistical Learning with Applications in R. Springer, New York.		
	Reference Books		
	1. James, G., Witten, D., Tibshirani, R. and Hastie, T. Neural Networks and Deep		
	Learning: A Textbook.		
	2. Introduction to Machine Learning The Wikipedia Guide.		
	After successful completion of this course, stud	ent will be able to:	
	1. Gain proficiency in applying least squares regression, nearest neighbors, and		
	local methods for high-dimensional data analysis.		
C	2. Master linear methods for regression and classification, including subset		
Course	selection, shrinkage methods, and linear discriminant analysis.		
Outcomes	3. Learn techniques for estimating in-sample	e prediction error, including cross-	
	validation and bootstrap methods.		
	4. Explore advanced model inference methods, including maximum likelihood		
	estimation, Bayesian inference, and the EN	A algorithm.	
		1	

• Lecturing, Visualization, Team Learning

# **MODE OF TRANSACTION**

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• Lecture, Seminar, Discussion, Questioning and Answering

### ASSESSMENT RUBRICS

Components	Weightage
End Semester	60
Evaluation(ESE)	
<b>Continuous Evaluation</b>	

16	
08	
16	
40	

#### **Sample Questions to Test Outcomes:**

- 1. How do least squares and nearest neighbors differ in their approach to supervised learning, and what are the advantages and limitations of each method in terms of handling high-dimensional data?
- 2. Can you explain the concept of the bias-variance tradeoff in the context of model selection? How do different methods such as Lasso, subset selection, and shrinkage methods address this tradeoff, and under what conditions would one method be preferred over another?
- 3. What are the key principles behind kernel methods and local regression in function approximation? How do these methods handle non-linear relationships between variables, and what are some practical considerations when choosing between different kernel functions or regression approaches?
- 4. How do linear methods for classification, such as logistic regression and linear discriminant analysis, differ in their approach to separating classes? Discuss the advantages and limitations of each method in terms of handling non-linearly separable data and the assumptions underlying their models.
- 5. What role does feature extraction play in filtering and smoothing techniques, particularly in the context of multidimensional splines and wavelet smoothing?
- 6. What are the key principles behind one-dimensional kernel smoothers, and how do they differ from other smoothing techniques such as splines or local regression?
- 7. How do structured local regression models in multidimensional spaces (Rp) extend the concepts of one-dimensional kernel smoothers?

POOL F: DISCIPLINE SPECIFIC ELECTIVE COURSE			
Course Code &	MSBST03DSE10 CATECODICAL DATA ANALVSIS		
Title	MSBS103DSE10-CATEGORICAL DATA ANALYSIS		
Programme	M.Sc. Biostatistics Semester III		
1	<ul> <li>Demonstrate a comprehensive understanding of categorical data and their measures, including appropriate techniques for analysis.</li> <li>Gain proficiency in conducting inference for contingency tables, applying appropriate statistical tests and interpreting results effectively.</li> </ul>		
Course			
Objectives	• Develop a deep understanding of generalized linear models,		
	<ul> <li>particularly focusing on binary and count data.</li> <li>Gain proficiency in handling longitudinal data, understanding its characteristics.</li> </ul>		
1			
	• Gain practical experience in fitting general linear mixed effect		
	models, conducting inference for random effects, and interpreting		
	results.		

<b>POOL F:</b>	DISCIPLIN	<b>NE SPECIFIC</b>	ELECTIVE	COURSE
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Modules	Content	Module Outcome
		• Demonstrate a thorough
Module I		understanding of categorical
Categorical	Categorical data and their measures,	data and their measures.
data and their	Inference for contingency tables,	• Proficient in conducting
measures	Generalized linear models for binary	inference for contingency
(15 Hours)	and count data. Estimation, Inference	tables.
	and fitting of model.	• Experience in applying
	CA UNING	generalized linear models to
		binary and count data.
Module II	Logistic, logit and log linear models	• Understanding Logistic, Logit,
Logit and	with categorical predictors, Logit and Log-linear Models.	
logistic models	models with multi responses-	• Proficient in specifying and
(15 Hours)	Nominal and ordinal responses.	estimating logistic, logit, and

Module III: Longitudinal data and their characteristics (15 Hours)	Longitudinal data and their characteristics, The general linear model for longitudinal data-ML and REML estimation, EM algorithm, General linear mixed effect model. Inference for the random effects. BLUPs, Empirical Bayes, Shrinkage model building and diagnostics, Generalized additive mixed model.	<ul> <li>log-linear models with categorical predictors.</li> <li>Apply logit models to datasets with multi-responses, encompassing both nominal and ordinal responses.</li> <li>Develop a comprehensive understanding of longitudinal data and their unique characteristics.</li> <li>Proficient in applying the general linear model to longitudinal data.</li> <li>Fitting general linear mixed effect models to longitudinal data.</li> </ul>
Module IV: Generalised linear model for longitudinal data (15 Hours)	Generalised linear model for longitudinal data, Random effect model, Transition models, Poisson and logistic regression models, Analysis and test. Classification of missing data mechanism- intermittent missing values and dropouts, weighted estimating equations, Modeling the drop out process.	<ul> <li>Explain the concepts and principles behind generalized linear models (GLMs) and their application to longitudinal data analysis.</li> <li>Develop a thorough understanding of random effect models and their role in analyzing longitudinal data with correlated observations.</li> <li>Understand the importance of model validation and diagnostic checks in longitudinal data analysis, and be able to apply these techniques effectively.</li> </ul>

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	Text Books
	1. Agresti, A. (2012). Categorical data analysis (Vol. 792). John Wiley &
	Sons.
	2. Diggle, P.J, Heagerty, P., Liang, K., Y & Zeger, S., I (2003), Analysis of
References	longitudinal data, Oxford university press.
Keterences	3. Lindsey, J., K. (1993) Models for repeated measurements, Oxford
	Reference Books
	1. Weiss, R., E.(2005), Modelling longitudinal data, Springer, New York.
	2. Little, R. J. A. & Rubin, D., B(2002), Statistical analysis with missing
	data, Wiley.
Course	After successful completion of this course, student will be able to:
Outcomes	1. Conduct analysis of longitudinal data.
	2. Apply statistical techniques to model longitudinal data and make
	predictions.
	3. Understand analysis of longitudinal data with missing data.
	4. Understand analysis of longitudinal data with time-dependent covariates.

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

### ASSESSMENT RUBRICS

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Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Eva	luation
Tests	16
Assignment	08
Seminar	16
Total	40

#### Sample Questions to Test Outcomes:

- 1. What are categorical data, and what measures are commonly used to summarize and analyze them?
- 2. Discuss common statistical tests used for assessing the association or independence between categorical variables in contingency tables.
- 3. Provide examples of binary and count data scenarios where GLMs are appropriate and discuss the interpretation of model coefficients in these contexts.
- 4. Describe the process of model estimation and interpretation in logistic regression models.
- 5. Discuss strategies for handling issues such as missing data, multicollinearity, and model validation to ensure the reliability and generalizability of the model results.
- 6. Compare and contrast maximum likelihood (ML) and restricted maximum likelihood (REML) estimation techniques in the context of longitudinal data analysis. Provide examples to illustrate your points.
- 7. Explain the concept of a general linear mixed effect model (GLMM) and its relevance in analyzing longitudinal data.
- 8. Define the Generalized Additive Mixed Model (GAMM) and discuss its advantages in modeling longitudinal data with non-linear relationships.
- 9. Explain the concept of the Generalized Linear Model (GLM) for longitudinal data analysis.
- 10. Define transition models and discuss their significance in longitudinal data analysis.

Course Code & Title	MSBST03DSE11-OPERATIONS RESEARCH			
Programme	M.Sc. Biostatistics Semester III			
Course Objectives	<ul> <li>Understand the fundamental concepts of linear programming problems (LPP) and their applications in real-world optimization scenarios.</li> <li>Develop proficiency in graphical solution techniques for visualizing and analyzing LPPs, including identifying feasible regions and optimal solutions.</li> </ul>			

### POOL F: DISCIPLINE SPECIFIC ELECTIVE COURSE

	• Demonstrate the ability to determine feasible, basic feasible, and	
	optimum basic feasible solutions to LPPs, both graphically and	
	analytically.	
	• Gain insight into the theoretical foundations of linear	
	<ul> <li>programming, including the analytical results and theoretical development of the simplex method.</li> <li>Master advanced solution techniques such as the use of artificial variables, the Big-M method, and the two-phase simplex method</li> </ul>	
	to solve complex LPPs.	
	• Explore practical optimization problems such as transportation	
	and assignment problems, and learn to apply appropriate solution	
1	methods to address them effectively.	

Modules	Content	Module Outcome
Module I Introduction to linear programming problem (15 Hours)	Introductiontolinearprogrammingproblem(LPP),graphical solution, feasible,basic feasible, and optimum basicfeasible solution to an LPP.Analytical results in general LPP,theoreticaldevelopmentofsimplex method.	<ul> <li>Define linear programming problems and their significance in optimization.</li> <li>Explain the concept of feasible solutions and their graphical representation.</li> <li>Identify basic feasible solutions and optimal basic feasible solutions in LPPs.</li> </ul>
Module II Simplex methods (15 Hours)	Artificial variables, Big-M method, two phase simplex method Duality, duality theorems, dual simplex methods.	<ul> <li>Derive analytical results for general linear programming problems.</li> <li>Understand the theoretical development of the simplex method for solving LPPs.</li> <li>Apply artificial variables, the Big-M method, and the two-phase simplex method to handle</li> </ul>

	special cases in LPPs.	
Module III Transportation and integer programming (15 Hours)	<ul> <li>Explore the concept of duality in linear programming and understand the duality theorems.</li> <li>Discuss the application of dual simplex methods in solving LPPs.</li> <li>Analyze transportation and assignment problems and apply appropriate solution techniques.</li> </ul>	
Module IV Game theory and applications (15 Hours)	<ul> <li>Game theory, pure and mixed strategies, conversion of two-person zero gain to a linear programming problem. Solution to game through algebraic, graphical and linear programming method.</li> <li>Examine integer programming and its solution methods.</li> <li>Explore network analysis techniques such as Critical Path Method (CPM) and Program Evaluation and Review Technique (PERT).</li> <li>Introduce game theory concepts, including pure and mixed strategies.</li> </ul>	
References	<ul> <li>Text Books</li> <li>1. K.V. Mital and Mohan, C (1996). Optimization Methods in Operations Research and SystemsAnalysis,3<sup>rd</sup> Edition, New Age International(Pvt.)Ltd.</li> <li>2. Kanti Swarup, Gupta, P. K.and John, M. M. (1985): Operations Research., Sultan Chand &amp; Sons.</li> <li>Reference Books</li> <li>1. Hadley, G.(1964).Linear Programming, Oxford&amp; IBH Publishing Co, New Delhi.</li> <li>2. Taha.H.A.(1982): Operation Research, An Instruction, Macmillan.</li> <li>3. HillerF.S.AndLieberman,G.J.(1995).IntroductiontoOperationsResearc h,McGrawHill</li> </ul>	
Course	After successful completion of this course, student will be able to:	

Outcomes	1.	Identify and develop operational research models from the
		verbal description of the real system.
	2.	Understand the mathematical tools that are needed to
		solve optimization problems.
	3.	Understand various methods in Integer programming and
	1	Game theory.

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

## ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Ev	aluation
Tests	16
Assignment	08
Seminar/Viva	16

### Sample Questions to Test Outcomes:

1. Define Linear Programming Problem (LPP) and discuss its significance in optimization. What are the key components of an LPP, and how is it formulated?

2. Explain the graphical solution method for solving LPPs. How is the feasible region identified, and how are optimal solutions determined graphically?

3. Define feasible solutions, basic feasible solutions, and optimum basic feasible solutions to an LPP. What criteria are used to identify these solutions?

4. Discuss the theoretical development of the simplex method for solving LPPs analytically. How does the simplex method iteratively move from one basic feasible solution to another to find the optimal solution? 5. Explain the concept of artificial variables and the Big-M method in the context of solving LPPs. How are artificial variables used to convert an LPP into a form suitable for the simplex method?

6. Describe the two-phase simplex method for solving LPPs. What are the two phases, and how does this method handle special cases such as degeneracy?

7. Discuss duality in linear programming, including duality theorems and the dual simplex method. What is the relationship between the primal and dual problems, and how is duality used in optimization?

8. Define the transportation problem and the assignment problem. How are these types of problems formulated as linear programming problems, and what are their applications?

Course Code & Title	MSBST03DSE12- DATA VISUALIZATION AND ANALYSIS USING PYTHON		
Programme	M.Sc. Biostatistics Semester III		
Course	• Gain proficiency in data acquisition and manipulation using Python		
Objectives	libraries.		
	• Understand and implement different data visualization techniques for		
	comprehensive data analysis.		
	• Learn the regression model building framework in Python, from		
	problem definition to model validation.		
5	• Develop skills in multiple linear regression including handling		
- A.	categorical variables and diagnosing model issues with Python.		

### POOL F: DISCIPLINE SPECIFIC ELECTIVE COURSE

Modules	Content	Module Outcome
Module I Introduction to PYTHON (15 Hours)	Introduction to PYTHON, Data acquisition processes, extraction, cleaning, annotation, integration, reduction, and transformation. Basic charts, multivariate visualization, pixel- oriented, geometric projection, icon- based, and hierarchical visualization.	<ul> <li>Ability to effectively extract, clean, annotate, integrate, reduce, and transform data using Python.</li> <li>Understanding of data preprocessing techniques to prepare datasets for analysis, including data normalization,</li> </ul>

		missing value imputation, and
		feature engineering.
	Data visualization tools, rank analysis,	• Proficiency in utilizing
	trend analysis, multivariate analysis,	advanced visualization
Module II	distribution analysis, correlation	methods such as pixel-
Data	analysis, and geographical analysis.	oriented, geometric
toolo	1000	projection, icon-based, and
(15 Hours)	Ver NIT	hierarchical visualization.
(15 Hours)		• Ability to communicate
		insights and findings derived
		from data visualization
		effectively.
	D	• Understanding of best
Madula III	Regression model building framework,	practices in data visualization
Niodule III Desmossier	covering problem definition, data pre-	design.
Regression	processing, model building,	• Ability to leverage interactive
model building	diagnostics, and validation. Simple	visualization tools and
Iramework	linear regression, coefficients of	dashboards to engage
(15 Hours)	determination, significance tests,	stakeholders and facilitate
	residual analysis, and	exploratory data analysis and
	confidence/prediction intervals.	decision-making processes.
A		• Understanding of the
A	*	regression model building
	Multiple linear regression, coefficients	framework and evaluation
Madula IV	of multiple determination,	using Python.
Multiple lineer	interpretation of regression coefficients,	• Capability to validate
ragrassion	categorical variables, diagnosing issues,	regression models using
(15 Hours)	heteroscedasticity and multicollinearity,	diagnostic techniques,
(13 110013)	outliers, autoregression, and variable	including confidence and
	transformation for robust regression	prediction intervals, to assess
	model building.	model accuracy and
		reliability for predictive
		analysis.

	Text Books	
	1. AndyKirk, Data Visualization a Handbook for Data Driven Design,	
	Sage Publications, 2016	
	2. Philipp K.Janert, Gnuplot in Action, Understanding Data with Graphs,	
	Manning Publications,2010.	
	Reference Books	
	1. Alberto Cordoba, "Understanding the Predictive Analytics Lifecycle",	
	Wiley,2014.	
	2. Eric Siegel, Thomas H. Davenport, "Predictive Analytics: The Power to	
References	Predict Who Will Click, Buy, Lie, or Die", Wiley, 2013.	
	3. James R Evans, "Business Analytics-Methods, Models and Decisions",	
	Pearson 2013.	
	4. R. N. Prasad, Seema Acharya, "Fundamentals of Business Analytics",	
	Wiley,2015.	
	5. Perkovie, L. (2011). Introduction to computing using python: An	
	Application development focus. Wiley Publishing.	
	6. McKinney, W. (2012). Python for data analysis: Data wrangling with	
	Pandas, NumPy, and IPython. "O Reilly Media, Inc."	
1	After successful completion of this course, student will be able to:	
	1. Apply Python libraries for data acquisition to extract, clean, integrate,	
5	and transform datasets efficiently.	
— Λ.	2. Utilize Python libraries for various visualization methods to interpret and	
Course	communicate complex data effectively.	
Outcomos	3. Construct regression models using Python libraries, including appropriate	
Outcomes	pre-processing techniques and diagnostic tools.	
	4. Analyze simple linear regression results, including coefficients of	
	determination and significance tests, using Python.	
	5. Implement multiple linear regression models in Python, addressing issues	
	like multicollinearity and heteroscedasticity for robust predictions.	

### • Lecturing, Visualization, Team Learning

# MODE OF TRANSACTION

• Lecture, Seminar, Discussion, Questioning and Answering ASSESSMENT RUBRICS

Components	Weightage	
End Semester Evaluation(ESE)	60	
Continuous Evaluation		
Tests	16	
Assignment	08	
Seminar	16	
Total	40	

#### Sample Questions to Test Outcomes:

- 1. How would you describe the process of data acquisition and what are its key components?
- 2. Can you explain the difference between data cleaning and data transformation?
- 3. What are some common techniques used for reducing the dimensionality of datasets during data preprocessing?
- 4. Describe a situation where you might encounter missing data during the data acquisition process and how would you handle it?
- 5. How do you decide which visualization technique to use for a given dataset and analysis objective?
- 6. Explain the purpose of rank analysis tools in data visualization and provide an example of when you would use them.
- 7. What is the significance of residual analysis in regression model building and how is it performed?
- 8. Describe the steps involved in conducting a simple linear regression analysis and interpreting its results.
- 9. How do you diagnose multicollinearity in a multiple linear regression model and what are its potential consequences?

10. Can you provide an example of how you would use autoregression in a regression model building process and explain its importance?

Course Code & Title	MSBST03IDC01-STATISTICAL DATA ANALYSIS USING SPSS			
Programme Offered	Department of Statistical Sciences Semester III			
Course	Demonstrate proficiency in navigating SPSS interface components			
Objectives	including the Data Editor window, SPSS Output window, and			
	various drop-down menus.			
	• Able to create, modify, import, and transform datasets using			
	SPSS.			
	• Gain an understanding of different types of variables and how to assign appropriate labels to them within SPSS.			
	• Learn to conduct inferential statistical analysis techniques such as			
	frequencies, cross tabs, independent sample t-tests, paired sample			
	t-tests, ANOVA, correlation, regression, confidence intervals, and			
	non-parametric tests using SPSS.			
	• Develop the ability to interpret and communicate the results of			
5	statistical analyses conducted in SPSS.			

#### POOL G: INTERDISCIPLINARY ELECTIVE COURSE (IDC)

Modules	Content	Module Outcome
Module I Introduction to SPSS and its interface (15 Hours)	SPSS windows, Data editor window, Types of variables and labels, SPSS output window, Drop down menus of SPSS, Creating and modifying data files	<ul> <li>Proficiency in navigate various SPSS windows.</li> <li>Able to differentiate between categorical and continuous variables and understand their significance in data analysis.</li> <li>Creating and modifying data files in SPSS.</li> </ul>
Module II	Import of data files, Transform drop	• Skills to transform variables

Data	down menu, computing variables,	using the Transform drop-down				
management in	Recode option, Data drop down	menu in SPSS.				
SPSS	menu, Split files, Weight cases,	• They will be able to perform				
(15 Hours)	Select cases.	operations such as computing				
		new variables, recoding existing				
	10000 0m100	variables.				
	100 11	• Able to utilize advanced data				
	Ver NIT,	manipulation features of SPSS.				
		• Demonstrate proficiency in				
Madula III	SPSS plots and graphs, Bar	creating various plots and graphs				
Graphics in	diagram, Pie diagram, Multiple	in SPSS.				
SDSS	bar diagram, Histogram, Box	• Develop the skills to interpret and				
(15 Hours)	plot, P-P plot, Q-Q plot, Scatter	analyze graphical representations				
(15 Hours)	diagram.	generated in SPSS.				
		• Demonstrate proficiency in				
	Analyze drop down menu,	conducting various statistical				
	Descriptive statistics,	analyses using the Analyze drop-				
Module IV	Frequencies, Cross tabs,	down menu in SPSS.				
Basic statistical	Compare means-independent	• Gain proficiency in conducting				
analysis	sample t test, paired sample t test,	inferential statistics tests such as				
(15 Hours)	ANOVA, Correlation,	frequencies, cross tabs,				
A	Regression, Confidence intervals,	independent sample t-tests, paired				
	Non-parametric test.	sample t-tests, ANOVA,				
<1/	WAL	correlation, regression.				
Jan Barris	Text Book	61				
	1. Hinton P R, Brownlow C, McMurray, I. and Cozens, B.(2004):					
	SPSS Explained, Routledge, Tay	lor and Francis group, New York.				
Keterences	References Reference Book					
	1. Sabine Landau, Brian S. Ever	itt (2003): A Handbook of				
	Statistical Analyses Using SP	SS, New York.				
Course	After successful completion of this c	course, student will be able to:				
Outcomes	1. Understand the usage of menus in SPSS window for drawing					
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various diagrams.

- Computing descriptive statistics, the comparison of means, ANOVA, non-parametric tests, simple correlation and regression procedures and apply for real data sets.
- 3. Acquire the skills of plotting different graphs using SPSS.

### **TEACHING LEARNING STRATEGIES**

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

Lecture, Seminar, Discussion, Questioning and Answering

### ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Ev	aluation
Tests	16
Assignment	08
Seminar	16
	10

#### Sample Questions to Test Outcomes:

- 1. Describe the use of Recode command in SPSS
- 2. Describe the use of Weight case facility in SPSS
- 3. What is the use of split file option in SPSS?
- 4. Distinguish between Value label and Values options in SPSS.
- 5. Explain the method of constructing and interpreting a Boxplot
- 6. Explain crosstab facility and its uses available in SPSS
- 7. How do you construct a frequency table and histogram using SPSS?
- 8. Explain Transform dropdown menu in SPSS.
- 9. Describe briefly the various options available in SPSS Analyze menu.

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10. Explain different methods for constructing graphs in SPSS.

Course Code & Title	MSBST03IDC02-STATISTICAL DATA ANALYSIS USING R					
Programme offered	Department of Statistical Sciences Semester III					
Course	• Demonstrate a solid understanding of the fundamental concepts in					
Objectives	R programming, including objects and their classes, operators,					
	vectors, matrices, lists, and data frames.					
	• Gain proficiency in indexing and accessing data within R, as well as importing and exporting data from various file formats.					
	• Able to apply R programming to compute descriptive statistics.					
	• Learn to create various graphical representations of data using R.					
	• Develop the skills to plot cumulative distribution functions					
	(CDFs) and probability density functions (PDFs) for various					
	values of parameters in standard probability distributions using R.					
	• Learn to generate random samples from standard probability					
6	distributions in R.					

# POOL G: INTERDISCIPLINARY ELECTIVE COURSE

Modules	Content	Module Outcome
Module I Introduction to R (15 Hours)	Introduction to R- Objects and their classes, operators, vectors and matrices, list and data frames, indexing and accessing data, importing and exporting data. Common built-in functions. Simple applications - Descriptive statistics.	<ul> <li>Demonstrate a solid understanding of the fundamental concepts of R programming.</li> <li>Acquire proficiency in indexing and accessing data within R, enabling them to effectively manipulate datasets for analysis.</li> <li>Able to apply basic descriptive statistics techniques in R to analyze datasets effectively.</li> </ul>

<b>Module II</b> R-Graphics ( <b>15 Hours</b> )	R-Graphics- Histogram, Box-plot, Stem and leaf plot, Scatter plot, Q- Q plot. Looping- for loop, repeat loop, while loop, if command, ifelse command.	<ul> <li>Demonstrate proficiency in creating various types of graphical representations using R, including histograms, box plots, etc.</li> <li>Able to apply their knowledge of R graphics and looping structures to conduct exploratory data analysis and statistical inference tasks.</li> </ul>
Module III Basic probability and distribution (15 Hours)	Basic concepts of probability and random variables, Probability distributions (Binomial, Poisson, Geometric, Uniform, Normal, Gamma, Beta), Plotting of cdf and pdf for different values of the parameters of standard distributions. Generations of random samples from standard distributions.	<ul> <li>Understanding of basic concepts of probability theory and random variables.</li> <li>Understanding different probability distributions commonly used in statistical analysis.</li> <li>Demonstrate the ability to plot cumulative distribution functions (CDFs) and probability density functions (PDFs) for different parameter values of standard distributions using R.</li> </ul>
Module IV Descriptive statistics (15 Hours)	The Descriptive statistics, the comparison of means, ANOVA, non-parametric tests, correlation and regression procedures.	<ul> <li>Demonstrate mastery in descriptive statistics, including measures of central tendency, dispersion, and distributional shape.</li> <li>Able to conduct and interpret various tests for comparing means, including independent samples t-tests, paired samples t-tests, etc.</li> </ul>

	Text Books						
	1. Purohit, S. G, Ghore, S. D and Deshmukh, S. R. (2004): Statistics						
D - f	Using R. Narosa.						
References	Reference Books						
	1. Dalgaard, P. (2008): Introductory Statistics with R, (Second						
	Edition), Springer.						
Course	After successful completion of this course, student will be able to:						
Outcomes	1. Understand various built-in functions in R programming for						
	statistical data analysis.						
	2. Understand different functions in R programming for writing						
	computer programmes and develop computer programmes for						
	different problems.						
	3. Understand different statistical test using R software						

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous E	valuation
Tests	16
Assignment	08
Seminar	16
Total	40

#### Sample Questions to Test Outcomes:

1. How will you install R in your computer?

- 2. How will you save, store and retrieve workspace in R?
- 3. Explain the seq() and rep() in R with examples.
- 4. How will you import data from excel to R?
- 5. What are the advantages of R over other statistical softwares?
- 6. Explain different types of arithmetic operators and assignment operators in R. Give examples.
- 7. Explain different ways of defining matrices in R.
- 8. Explain the different forms of sequence function in R. Give examples.
- 9. Expalin the built-in functions in R with examples.
- 10. Describe low level plotting functions in R.



	FOURTH SEMESTER								
Sl No	Course Code	Title of Paper		Contact Hours/Week		Marks			
			L	T/S	Р	ESE	CE	Total	Credits
4.1	MSDST04DSC12	Project/Dissertation and	2			60	40	100	12
	WSD5104D5C12	Subject Viva	20	2	1	2			
		18/ 1		102	1	60	40	100	12
DISCIPLINE SPECIFIC ELECTIVES (DSE)									
		Elective-I (DSE) (Practical)							
4.2	MSBST04DSExx	(One course has to be chosen	3	2		60	40	100	3
		from <b>Pool H</b> )							
		Elective-II (DSE) (One		1					
4.3	MSBST04DSExx	course has to be chosen from	3	2		60	40	100	3
		Pool I)							
Total Credits						18			

L=Lecture, T/S=Tutorials/Seminar, P/I=Practical/Internship, CE =Continuous Evaluation, ESE = End

Semester Evaluation

POOL H:- List of Courses for Elective (Practical)									
51110	DISCIPLINE SPECIFIC ELECTIVES (DSE)								
4.2.1	MSBST04DSE13	Biostatistical Computing Using SAS-III ( <b>Practical</b> )	7	2	6	60	40	100	3
4.2.2	MSBST04DSE14	Biostatistical Computing Using R-III ( <b>Practical</b> )		2	6	60	40	100	3
	POOL I:- List of Courses for Elective								
<b>DISCIPLINE SPECIFIC ELECTIVES (DSE)</b>									
4.3.1	MSBST04DSE15	Categorical Data Analysis	4	1			1		3
4.3.2	MSBST04DSE16	Advanced Time Series Analysis	4	1					3
4.3.3	MSBST04DSE17	Actuarial Statistics	4	1					3
4.3.4	MSBST04DSE18	Statistical Quality Control	4	1					3

4.3.5	MSBST04DSE19	Advanced Bayesian Computing with R	4	1			3
4.3.6	MSBST04DSE20	Demographic Studies	4	1			3
4.3.7	MSBST04DSE21	Analysis of Longitudinal Data	4	1	1		3

### POOL H: DISCIPLINE SPECIFIC ELECTIVE COURSE

Course Code &	MSBST04DSE13-BIOSTATISTICAL	COMPUT	FING USING				
Title	SAS - III (PRACTICAL)						
Programme	M.Sc. Biostatistics	Semester	IV				
Course	• To introduce some advanced statistic	cal computir	ng techniques in				
Objectives	biostatistics to extract information and visualization thereby						
	enabling them to perform data analysis effectively and efficiently						
	in SAS programming.						
	• Illustrate different statistical techniqu	es based on	all the elective				
	course in fourth semester.						

Modules	Content	Module Outcome					
	Biostatistical Computing IV is a	• Describe different					
5	practical course. The practical is based	statistical technique to					
- Α.	on all the elective courses in the	solve problems coming					
	fourth semester.	under all the elective					
2-1	N.	courses in fourth					
	WUD WIER	semester.					
Course	After successful completion of this course, student will be able to:						
Outcomes	1. Equipped with different theoretical methods in biostatistics to						
	achieve the objectives.						
	2. Enhanced with the basic concepts of statistical theories besides						
	developing their ability to handle real world problems with large						
	scale data.						

### **TEACHING LEARNING STRATEGIES**

• Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION

• Lecture, Seminar, Hands on training

# ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation	60
Continuous Evaluation	
Practical Tests	32
Record	8
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### POOL H: DISCIPLINE SPECIFIC ELECTIVE COURSE

Course Code &	MSBST04DSE14-BIOSTATISTICAL COMPUTING USING R -		
Title	III (PRACTICAL)		
Programme	M.Sc. Biostatistics Semester IV		
Course	• To introduce some advanced biostatistical computing techniques		
Objectives	in applied statistics to extract information and visualization		
	thereby enabling them to perform data analysis effectively and		
	efficiently in R programming.		
5	• Illustrate different biostatistical techniques based on all the		
— Д.	elective course in fourth semester.		
	7		

Modules	Content	Module Outcome
	Biostatistical Computing IV is a practical course. The practical is based on all the elective courses in	• Describe different statistical technique to solve problems coming
	the fourth semester.	under all the elective courses in fourth semester.
Course	After successful completion of this course,	student will be able to:

Outcomes	1. Equipped with different theoretical methods in biostatistics to	
	achieve the objectives.	
	2. Enhanced with the basic concepts of biostatistical theories besides	
	developing their ability to handle real world problems with large	
	scale data.	

- Practical sessions through computers, statistical computations, Team Learning MODE OF TRANSACTION
  - Lecture, Seminar, Hands on training

#### **ASSESSMENT RUBRICS**

Components	Weightage	
End Semester Evaluation	60	
Continuous Evaluation		
Practical Tests	32	
Record	08	
Total	40	

### **POOL I: DISCIPLINE SPECIFIC ELECTIVE COURSE**

Course Code & Title	MSBST04DSE15- CATEGORICAL DATA ANALYSIS		
Programme	M.Sc. Biostatistics Semester IV		
Course Objectives	<ul> <li>Understand the funda their implications in r</li> <li>Study various sampling random sampling, stra apply them appropria</li> <li>Learn about exact tes traditional statistical to</li> </ul>	mentals of statistical measurement scales and research design and analysis. ng frameworks and techniques, including atified sampling, and cluster sampling, and tely in research settings. ts and their significance in situations where tests may not be applicable.	

Modules	Content	Module Outcome
<b>Module I</b> Overview of analysis strategies ( <b>15 Hours</b> )	Scale of Measurements, sampling frameworks, overview of analysis strategies, Chi-square statistic, exact tests, difference in proportions, odds ratio and relative risk, sensitivity and specificity, McNemar's test.	<ul> <li>Understand the differences between nominal, ordinal, interval, and ratio scales of measurement.</li> <li>Apply appropriate measurement scales in research design and data analysis.</li> <li>Identify the implications of different measurement scales on statistical analysis techniques.</li> </ul>
Module II Measure of association and contingency tables (15 Hours)	Mantel-Haenszel test, Measure of association, sets of 2 x r tables, sets of s x 2 tables, relationship between sets of tables.	<ul> <li>Demonstrate proficiency in selecting and implementing appropriate sampling frameworks for different research scenarios.</li> <li>Understand the strengths and limitations of various sampling methods.</li> </ul>
Module III Mantel- Haenszel methodology and application (15 Hours)	Association, exact tests for association, Measure of association, observer agreement, test for ordered differences, General Mantel-Haenszel methodology, Mantel- Haenszel applications.	<ul> <li>Gain familiarity with descriptive statistics, hypothesis testing, and regression analysis.</li> <li>Identify appropriate analysis strategies based on research objectives and data characteristics.</li> <li>Evaluate the assumptions and validity of different analysis techniques.</li> </ul>
Module IV Advanced topics	Advanced topics: application to repeated measures, Wilcoxon-Mann- Whiteney test, Kruskal-Wallis test,	• Navigate and manipulate data tables efficiently.

(15 Hours)	Friedman's Chi-square test, Aligned • Perform basic to advanced data		
	rank test for randomised complete transformations.		
	blocks, Durbin's test for balanced • Generate informative tables and		
	incomplete blocks, Rank analysis of graphical displays to		
	covariance. summarize data effectively.		
Y	Text Books		
	1. Stokes, M. E., Davis, C. S., & Koch, G. G. (2012). Categorical data		
	analysis using SAS. SAS institute.		
	2. Agresti, A. (2012). Categorical data analysis (Vol. 792). John Wiley &		
	Sons.		
References	Reference Books		
	1. Powers, D., & Xie, Y. (2008). Statistical methods for categorical data		
	analysis. Emerald Group Publishing.		
	<ol> <li>Sloane, D., &amp; Morgan, S. P. (1996). An introduction to categorical data analysis. <i>Annual review of sociology</i>, 22(1), 351-375.</li> <li>Lawal, B., &amp; Lawal, H. B. (2003). <i>Categorical data analysis with SAS</i> <i>and SPSS applications</i>. Psychology Press.</li> </ol>		
	After successful completion of this course, student will be able to:		
	1. Demonstrate a thorough understanding of various statistical		
	methods.		
- A	2. Proficiency in working with data tables, conducting data		
Course	manipulation, and performing statistical analysis, enhancing their		
Outcomes	ability to manage and analyze large datasets in real-world research		
-1/	settings.		
	3. Cultivate critical thinking skills by critically evaluating research		
	designs, selecting appropriate sampling frameworks.		
	4. Communicate statistical findings clearly and effectively through		
	written reports, presentations, and graphical representations.		

• Lecturing, Visualization, Team Learning

### **MODE OF TRANSACTION**

# • Lecture, Seminar, Discussion, Questioning and Answering ASSESSMENT RUBRICS

Components	Weightage
End Semester Evaluation	60
Continuous Eva	luation
Tests	16
Assignment	08
Seminar	16
Total	40

#### Sample Questions to test Outcomes:

- Explain the different scales of measurements and how they impact statistical analysis. Provide examples for each scale.
- 2. Discuss the importance of sampling frameworks in research design. How do different sampling techniques affect the validity of study findings?
- 3. Give an overview of analysis strategies commonly used in research. Compare and contrast their strengths and limitations.
- 4. Demonstrate how to work with tables in the SAS system for data analysis and interpretation.
- 5. Explain the Chi-square statistic and its significance in hypothesis testing. Provide a real-world example illustrating its application.
- 6. Discuss the concept of odds ratio and relative risk in epidemiological studies. How are they calculated and interpreted?
- 7. Define sensitivity and specificity in diagnostic testing. How do these measures inform the accuracy of a diagnostic test?
- 8. Explain McNemar's test and its application in paired nominal data analysis. Provide a step-by-step example to illustrate its usage.
- 9. Describe the Mantel-Haenszel test and its role in analyzing categorical data. Provide examples of situations where this test would be appropriate.

10. Discuss advanced topics such as repeated measures analysis and non-parametric tests like Wilcoxon-Mann-Whitney, Kruskal-Wallis, and Friedman's Chi-square tests.

Course Code & Title	MSBST04DSE16-ADVANCED TIME SERIES ANALYSIS			
Programme	M.Sc. Biostatistics Semester IV			
Course Objectives	<ul> <li>Understand the fundamental concepts of stochastic processes and their applications in modeling time series data, providing a solid foundation for further analysis.</li> <li>Learn to analyze auto-covariance, auto-correlation, and spectral density properties of time series data, enabling students to characterize and interpret temporal patterns effectively.</li> <li>Gain in-depth knowledge of autoregressive (AR), moving average (MA), autoregressive moving average (ARMA), and autoregressive integrated moving average (ARIMA) models, enabling students to select and apply appropriate models for different time series data sets.</li> <li>Explore spectral analysis, periodgrams, correlograms, and diagnostic checks for model validation, equipping students with advanced analytical tools for interpreting and analyzing time series data effectively.</li> </ul>			
	7			

#### POOL I: DISCIPLINE SPECIFIC ELECTIVE COURSE

Modules	Content	Module Outcome
	Motivation, Time series as a discrete	• Understanding Stochastic
Madula I	parameter stochastic process, Auto-	Processes.
Noulle I Devisit to	Covariance, Auto-Correlation and	• Applying Stochastic Processes
foundations of	spectral density and their properties.	to Time Series Analysis.
time agrice	Exploratory time series analysis,	• Interpreting Stochastic Process
(15 Hours)	Exponential and moving average	Properties.
(15 Hours)	smoothing, Holt-Winter smoothing,	• Applying Stochastic Processes
	forecasting based on smoothing,	in Forecasting:
	Adaptive smoothing.	
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Module II Detailed study of the stationary ARMA models (15 Hours)	Detailed study of the stationary process: Autoregressive, Moving Average, Autoregressive Moving Average and Autoregressive Integrated Moving Average Models. Choice of AR/MA periods.	<ul> <li>Understanding exploratory Analysis Techniques.</li> <li>Understanding Smoothing Methods.</li> <li>Characterizing Stationary Processes.</li> <li>Interpreting Time Series Properties.</li> </ul>
Module III Estimation and forecasting of ARIMA models (15 Hours)	Estimation of ARMA models: Yule- Walker estimation for AR Processes, Maximum likelihood and least square estimation for ARMA Processes, Discussion (without proof) of estimation of mean, Auto- covariance and autocorrelation function under large samples theory, Residual analysis and diagnostic checking. Forecasting using ARIMA models.	<ul> <li>Understanding forecasting Techniques.</li> <li>Identifying model Selection and Estimation.</li> <li>Validation and Diagnostic Checks.</li> <li>Utilize advanced analytical tools such as spectral analysis, periodograms, and correlograms to analyze and interpret time series data effectively, enhancing their ability to derive insights and make informed decisions.</li> </ul>
Module IV Spectral and seasonal analysis (15 Hours)	Spectral analysis of weakly stationary process. Periodogram and correlogram analysis. Seasonal ARIMA models (Basic concepts only), ARCH and GARCH models (Basic concepts only)	<ul> <li>Understand seasonal and non- seasonal models such as seasonal ARIMA, ARCH, and GARCH models, enabling them to capture and forecast complex temporal patterns within data sets.</li> <li>Apply advanced analytical techniques including spectral</li> </ul>

	analysis, periodograms, and		
	correlograms to analyze and		
	interpret time series data		
	effectively.		
	Text Books		
	1. Box G.E.P and Jenkins G.M. (1970). Time Series Analysis, Forecasting		
	and Control. Holden -Day.		
	2. Brockwell P.J and Davis R.A. (1987). Time Series: Theory and		
	Methods, Springer Verlag.		
Keterences	Reference Books		
	1. Abraham B and Ledolter J.C. (1983). Statistical Methods for		
	Forecasting, Wiely		
	2. Anderson T.W. (1971). Statistical Analysis of Time Series, Wiely.		
	3. Fuller W.A. (1978). Introduction to Statistical Time Series, John Wiley.		
	After successful completion of this course, student will be able to:		
Course	1. Understand exploratory time series analysis and its real		
Outcomes	data application.		
	2. Understand autoregressive models and their estimation methods.		
	3. Understand non-linear time series models and their		
	estimation methods.		
	4 Apply statistical techniques to time series data and make		
	+. Apply statistical techniques to time series data and make		

• Lecturing, Visualization, Team Learning

## **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

# ASSESSMENT RUBRICS

Components	Weightage	
End Semester	60	
Evaluation(ESE)		

Continuous Evaluation	
Tests	16
Assignment	08
Seminar/Viva	16
Total 40	

- 1. What is the significance of time series analysis in understanding data trends and making predictions? Discuss the motivation behind studying time series data.
- 2. Define a discrete parameter stochastic process and explain how time series can be viewed as such a process. How does this perspective help in analyzing time series data?
- 3. Explain the concepts of auto-covariance, auto-correlation, and spectral density in the context of time series analysis. Discuss their properties and their roles in characterizing time series behavior.
- 4. Describe the techniques involved in exploratory time series analysis. How do tests for trend and seasonality contribute to understanding time series patterns?
- Compare and contrast exponential smoothing, moving average smoothing, and Holt-Winter smoothing methods for time series forecasting. Provide examples illustrating their application.
- 6. Provide a detailed study of stationary processes, including autoregressive (AR), moving average (MA), autoregressive moving average (ARMA), and autoregressive integrated moving average (ARIMA) models. Discuss the choice of AR/MA periods in model selection.
- 7. Discuss different estimation methods for ARMA models, including Yule-Walker estimation for AR processes and maximum likelihood/least square estimation for ARMA processes. Explain the concept of residual analysis and diagnostic checking.
- 8. Explain the concept of spectral analysis for weakly stationary processes. Discuss the use of periodograms and correlograms in analyzing spectral density and autocorrelation functions.

9. Introduce the basic concepts of seasonal ARIMA models, ARCH models, and GARCH models in time series analysis. Discuss their relevance and applications in modeling time series data.

Course Code & Title	MSBST04DSE17- ACTUARIAL STATISTICS		
Programme	M.Sc. Biostatistics Semester IV		
Course Objectives	<ul> <li>Develop a greater understand application in actuarial statist</li> <li>Describe the core areas of a areas actuarial principles, the</li> <li>Describe estimation procedur</li> <li>Explain the concept of surviv</li> <li>Understand the application environment.</li> <li>Describe Net premiums and i</li> <li>Expand their applied knowle actuarial studies and statistics</li> </ul>	ing of statistical pri ics. ctuarial practice an ories and models. es for lifetime distri al models. of knowledge of th ts various types. edge in various spe	inciples and their ad relate to those ibutions. he life insurance

Modules	Content	Module Outcome
Module I Insurance and utility theory (15 Hours)	Insurance and utility theory, models for individual claims and their sums, survival function, curtate future lifetime, force of mortality. Life tables and its relation with survival function, examples, assumptions for fractional ages, some analytical laws of mortality, select and ultimate tables.	<ul> <li>Explains the utility theory and insurance.</li> <li>Explain survival function and application.</li> <li>Examine the properties of force of mortality.</li> <li>Define Life tables and its relation with survival function, examples.</li> </ul>
Module II	Multiple life functions, joint life and	• Explain Multiple life functions

Multiple life	last survivor status, insurance and	and its properties.	
functions	annuity benefits through multiple life	• Articulate the insurance and	
(15 Hours)	functions evaluation for special	annuity benefits through	
	mortality laws. Multiple decrement	multiple life functions	
	tables, central rates of multiples	evaluation for special mortality	
	decrement, net single premiums and	laws.	
	their numerical evaluations.	• Explains the Multiple	
	Ver NIII	decrement tables.	
		• Describe net single premiums	
		and their numerical evaluations.	
	Compound Poisson distribution and	Define Distribution of	
Module III	its applications. Principles of	aggregate claims.	
Compound	compound interest: Nominal and	• Derive the compound Poisson	
Poisson	effective rates of interest and	distribution and explain its	
distribution and	discount, force of interest and	applications.	
its applications	discount, compound interest,	• Explain Principles of	
(15 Hours)	accumulation factor, continuous	compound interest and its	
	compounding.	attributes.	
	Insurance payable at the moment of		
	death and at the end of the year of		
	death-level benefit insurance,	- C	
A.	endowment insurance, differed	• Explain the Life insurance and	
Module IV	insurance and varying benefit	its types.	
Different types	insurance, recursions, commutation	• Describe Insurance payable at	
of insurance	functions. Life annuities: Single	the moment of death and at the	
and amenities	payment, continuous life annuities,	end of the year of death-level	
(15 Hours)	discrete life annuities, life annuities	benefit insurance	
(13 110013)	with monthly payments,	• Explain the Life annuities and	
	commutation functions, varying	its types.	
	annuities, recursions, complete		
	annuities immediate and apportion		
	able annuities-due.		

References	Text Books		
	1. Beard, R.E., Penlikainen, T. and Pesonnen, E (1984): Risk Theory: The		
	Stochastic Basis of Insurance, 3rd Edition, Chapman and Hall, Londan.		
	Reference Books		
	1. Bowers, N.L., Gerber, H.U., Hickman, J.E., Jones, D.A. and Nesbitt.		
-	C.J. (1997): Actuarial Mathematics', Society of Actuarias, Ithaca,		
	Illiois, U.S.A., second Edition.		
	2. Neill, A. (1977): Life Contingencies, Heineman.		
Course	After successful completion of this course, student will be able to:		
Outcomes	1. Understand the principles of insurance and utility theory, and apply		
	them to analyze individual claims and their aggregate sums.		
	2. Analyze the relationship between life tables and survival functions,		
	and apply this knowledge to evaluate insurance and annuity		
	benefits.		
	3. Demonstrate proficiency in using multiple decrement tables to		
	assess insurance risks.		
	4. Apply the Compound Poisson distribution and principles of		
	compound interest to model insurance-related phenomena.		
	5. Develop skills in evaluating and designing life annuities, including		
	single payment, continuous, discrete, and monthly payment		
5	annuities.		

• Lecturing, Visualization, Team Learning

# MODE OF TRANSACTION

• Lecture, Seminar, Discussion, Questioning and Answering

# ASSESSMENT RUBRICS

Components	Weightage	
End Semester	60	
Evaluation(ESE)	00	
<b>Continuous Evaluation</b>		

Total	40
Seminar/Viva	16
Assignment	08
Tests	16

- 1. Define insurance and utility theory. How does utility theory influence decisionmaking in insurance?
- 2. Discuss models for individual claims and their sums in the context of insurance. How are these models used to assess risk and determine premiums?
- 3. Explain the concepts of survival function, curtate future lifetime, and force of mortality in the context of life insurance. How are these concepts related?
- 4. Describe life tables and their relationship with the survival function. Provide examples to illustrate the use of life tables in actuarial calculations.
- 5. What assumptions are made for fractional ages in life tables? How do these assumptions impact the accuracy of actuarial calculations?
- 6. Discuss some analytical laws of mortality commonly used in actuarial science. What are select and ultimate tables, and how are they applied?
- 7. Explain multiple life functions and their significance in joint life and last survivor status insurance policies. How are insurance and annuity benefits evaluated using multiple life functions?
- 8. Define multiple decrement tables and central rates of multiple decrements. How are net single premiums calculated using multiple decrement tables?
- 9. Describe the compound Poisson distribution and its applications in insurance. How is it used to model the frequency and severity of insurance claims?
- 10. Discuss the principles of compound interest, including nominal and effective rates of interest, force of interest and discount, and continuous compounding. How are these principles applied in actuarial calculations for insurance products like endowment and annuity policies?

Course Code & Title	MSBST04DSE18-STATISTICAL QUALITY CONTROL			
Programme	M.Sc. Biostatistics	Semester	IV	
Course Objectives	<ul> <li>Understand the principles and theory control (SPC) and the significance management.</li> <li>Develop proficiency in constructing control charts for variables (R, s charcharts), as well as modified control cliptication of the definition of the performance operating characteristic (OC) and curves.</li> <li>Learn about advanced control chart to average control charts, exponentiall (EWMA) charts, and cumulative sum</li> <li>Master various sampling plans such sampling, multiple sampling, and including rectifying inspection plans.</li> <li>Analyze the performance of sampling average total inspection (ATI) curves und average total inspection (ATI) curves average with double specification limits</li> </ul>	ry behind st of control of g and interp rts) and attributes average run techniques, in y weighted (CUSUM) of as single s sequential g plans using erage sample rves. lans for sing ariance, as v	tatistical process charts in quality oreting Shewhart butes (p, np, c, u charts through n length (ARL) ncluding moving moving average charts. sampling, double sampling plans, metrics like OC, e number (ASN), gle specification vell as sampling	

Modules	Content	Module Outcome
	Introduction to quality and quality	• Understand the theory behind
Module I	assurance, total quality	control charts and their role in
Introduction to	management, quality control,	SPC.
quality and	Statistical process control, theory of	• Demonstrate proficiency in
quality	control charts, Shewhart control	constructing and interpreting

assurance	charts for variables, R,s charts, p,	Shewhart control charts for
(15 Hours)	np, c, u charts, modified control	variables (R, s charts) and
	charts.	attributes (p, np, c, u charts).
		• Explore modified control charts
	1.0	and their applications in various
	0000 cm100	industries.
Module II Control charts and process capability indices (15 Hours)	O.C and ARL curves of control charts, moving average control charts, EWMA charts, CUSUM charts, process capability analysis, process capability indices.	<ul> <li>Analyze operating characteristic (OC) and average run length (ARL) curves of control charts.</li> <li>Apply moving average control charts, exponentially weighted moving average (EWMA) charts, and cumulative sum (CUSUM) charts to monitor process variability.</li> <li>Conduct process capability analysis and calculate process capability indices to assess the performance of a process.</li> </ul>
<b>Module III</b> Sampling Plans ( <b>15 Hours</b> )	Single sampling, double sampling, multiple sampling and sequential sampling plans, rectifying inspection plans, measuring performance of the sampling plans - OC,AOQ,ASN, ATI curves.	<ul> <li>Differentiate between single sampling, double sampling, multiple sampling, and sequential sampling plans.</li> <li>Develop rectifying inspection plans to improve quality control processes.</li> <li>Measure the performance of sampling plans using metrics such as OC, average outgoing quality (AOQ), average sample number (ASN), and average total inspection (ATI) curves.</li> </ul>

Module IV Sampling plans with double specification limits (15 Hours)	Sampling plans for single specification limit with known and unknown and unknown variance. Sampling plans with double specification limits, comparison of sampling plans by variables and attributes, Continuous sampling plans I, II and III.	<ul> <li>Design sampling plans for single specification limits with known and unknown variance.</li> <li>Implement sampling plans with double specification limits and compare them based on variables and attributes.</li> <li>Explore continuous sampling plans (I, II, and III) and understand their applications in industries with continuous production processes.</li> </ul>		
	Text Books	actiontoStatisticalQualityControl 5thEd		
	1. Montgomory, D.C. (2005), Introduction to Statistical QualityControl. 5thEd ition. Wiley, New-York.			
	2. Gerant, E.L. and Leaven Worth, R.S. (1980). Statistical Quality Control. Mc			
	GrawHill			
Reference Books				
References	1. Duncan, A.J.(1986).Quality Control and Industrial Statistics.			
	2. Mittage, H.J. and Rinne, H. (1993). Statistical Methods for Quality			
	Assurance. Chapmanand Hall.			
A	1990).Statistical Process Control. East-			
- 73	west Press. A Schilling F G (1982) Acceptant	esamplinginQualityControl MarcelDe		
41	kker.	esampning inquanty control. Marcenze		
	After successful completion of this of	course, student will be able to:		
	1. Understand the construction v	arious control charts and their real		
Course	data applications.			
Outcomes	2. Understand various process ca	pability indices and their applications.		
	<ol> <li>Understand various different a and variables.</li> </ol>	acceptance sampling plans for attributes		

## • Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

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# • Lecture, Seminar, Discussion, Questioning and Answering ASSESSMENT RUBRICS

	Components	Weightage		
2	End Semester Evaluation(ESE)	60		
×,	Continuous Evaluation			
T A S	Tests	16		
	Assignment	08		
	Seminar/Viva	16		
	Total	40		

- 1. Define Statistical Process Control (SPC) and discuss its importance in quality management. What are the key objectives of implementing SPC in manufacturing processes?
- 2. Explain the theory of control charts in SPC. What are the fundamental principles behind control charts, and how do they help in monitoring and controlling process variability?
- 3. Describe Shewhart control charts for variables, including R charts and s charts. How are these charts constructed, and what do they indicate about process stability and variation?
- 4. Discuss p, np, c, and u charts in SPC. What types of processes are these charts suitable for, and how are they interpreted in terms of process control?
- 5. Explain modified control charts and their applications in SPC. What modifications can be made to traditional control charts, and under what circumstances are these modifications necessary?
- 6. Define Operating Characteristic (OC) and Average Run Length (ARL) curves of control charts. How are these curves used to evaluate the performance of control charts?

- 7. Describe moving average control charts, exponentially weighted moving average (EWMA) charts, and cumulative sum (CUSUM) charts. What advantages do these charts offer over traditional control charts?
- 8. Explain process capability analysis and process capability indices. How are these measures used to assess the ability of a process to meet specified quality requirements?
- 9. Discuss single sampling, double sampling, multiple sampling, and sequential sampling plans. What are the differences between these sampling plans, and how are they applied in quality inspection?
- 10. Describe sampling plans for single specification limits and double specification limits. How do these sampling plans vary based on known and unknown variance, and how are they evaluated using OC, AOQ, ASN, and ATI curves?

Course Code & Title	MSBST04DSE19- ADVANCED BAYESIAN COMPUTING WITH R				
Programme	M.Sc. Biostatistics Semester IV				
	• Understand the fundamental principles of statistical decision-				
	making, including randomized decision rules and standard loss				
	functions.				
	• Explore the concept of prior information and its incorporation into				
Course	decision-making.				
Objectives	• Master the application of Bayes' theorem for inference, including				
	the estimation of prior and posterior densities, and gain				
<1/	proficiency in analyzing parametric families and likelihoods, such				
1	as the exponential family.				
	• Apply learned concepts and techniques to practical examples and				
	real-world problems, using software packages such as Learn				
	Bayes and Win-BUGS.				

Modules	Content	Module Outcome

Module I Fundamentals of Bayesian concepts (15 Hours)	Statistical decision problem, randomized decision rule, decision principle, standard loss functions, Prior information, subjective determination of prior density, non- informative priors, maximum entropy priors, conjugate priors, discrete prior. Parametric family and likelihood, exponential family, Bayes' theorem for inference, prior and posterior densities.	· · · / 3 · · · · ·	Understand and explain the concept of statistical decision- making. Apply decision principles effectively in various decision-making scenarios. Evaluate and compare different decision strategies based on their performance in minimizing expected loss and achieving desired outcomes.
Module II Bayes models and Learn Bayes package (15 Hours)	single parameter models, normal distribution with known variance and unknown mean, normal with known mean and unknown variance, Poisson model, normal distribution with both parameters unknown, multinomial model, Dirichlet prior, Bioassay experiment, comparing two proportions, predictive distribution, beta-binomial distribution, multivariate normal distribution, Introduction to Learn Bayes package, Examples using Learn Bayes package.		Demonstrate proficiency in formulating and specifying prior. Apply Bayes' theorem for inference tasks and analyze the impact of prior specification on posterior inference. Evaluate the suitability of different prior distributions for specific modeling scenarios.
Module III	Computing integrals using Monte-	•	Implement computational
Introduction to	Carlo simulation, approximation based		methods such as Monte Carlo
Markov Chain	on posterior mode, importance		simulation, importance
Monte Carlo	sampling, Markov Chain Monte Carlo		sampling, and Markov Chain
methods	methods, Metropolis-Hastings		Monte Carlo (MCMC).
(15 Hours)	algorithm, random walk, Gibbs	•	Evaluate the performance and
	sampling.		efficiency of different

| Scheme and Syllabus of M Sc. Biostatistics- 2023 Admission onwards- Kannur University

	computational techniques in		
	generating posterior samples		
	and estimating posterior		
	distributions.		
Module IV Hierarchical models (15 Hours)	<ul> <li>Hierarchical models, shrinkage estimators, posterior predictive model checking, comparison of hypotheses, Bayes factor, one sided test for normal mean, two-sided test for normal mean, two-sided test for normal mean, normal linear regression model, prediction of future observations, examples and R codes, introduction to Win-BUGS package.</li> <li>Understand the concept of hierarchical models and their application in modeling complex data.</li> <li>Perform posterior predictive model checking to assess the adequacy of hierarchical models and identify potential model misspecifications.</li> </ul>		
	Text Books		
	1. Jim Albert (2007). Bayesian Computation with R, New York: Springer		
	Verlag.		
	2. Berger, O.J. (1985). Statistical decision Theory and Bayesian Analysis, Seco		
	ndEdition,SpringerVerlag.		
	3. Bensal, A. K.(2008).Bayesian Parametric Inference, New Age, Delhi.		
	Reference Books		
References	1. Ferguson, T.S. (1967). Mathematical Statistics: ADecision Theoretic Appr		
- A.	oach,AcademicPress,New-York.		
	2. Bolstad, W.(2004). Introduction to Bayesian Statistics, Hoboken, NJ:		
<1/	John Wiley.		
1	3. Gelman, A., Carlin, J., Stern, H. and Rubin, D. (2003). Bayesian Data Analys		
	is,NewYork:ChapmanandHall.		
	4. Gilks, W. R., Richardson, S and Spiegelhalter, D.J.(1996). Markov		
	Chain Monte Carlo in Practice. Chapman & Hall/ CRC, New York.		
Course	After successful completion of this course, student will be able to:		
Outcomes	1. Understand the advantageous Bayes estimation over that based		
	on frequentist approach.		
	2. Understand the LearnBayes package for various Bayesian		

computations

- 3. Understand MCMC methods in various situations in which the exact computation is difficult.
- 4. Understand Gibbs sampling to generate random samples from a multivariate distribution.

## **TEACHING LEARNING STRATEGIES**

• Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

## **ASSESSMENT RUBRICS**

Components	Weightage	
End Semester Evaluation(ESE)	60	
Continuous Ev	aluation	
Tests	16	
Assignment	08	
Seminar/Viva	16	

#### Sample Questions to Test Outcomes:

1. Define a randomized decision rule and explain its significance in statistical decisionmaking.

2. Discuss the role of prior information in Bayesian inference and explain how it is incorporated into the decision-making process.

3. Describe the concept of maximum entropy priors and explain when they are useful in Bayesian analysis.

4. Compare and contrast Monte Carlo simulation and importance sampling methods for estimating posterior distributions.

5. Explain the Metropolis-Hastings algorithm and discuss its advantages and limitations in Markov Chain Monte Carlo (MCMC) sampling.

6. Provide an example of a hierarchical model and explain how it can be used to analyze data with nested levels of variability.

7. Discuss the concept of shrinkage estimators and explain how they address overfitting in hierarchical modeling.

8. Explain the process of posterior predictive model checking and discuss its importance in assessing model adequacy.

9. Compute the Bayes factor for two competing hypotheses and interpret the results in the context of model comparison.

10. Implement the Win-BUGS software package to perform Bayesian analysis on a given dataset, and interpret the results obtained.

Course Code &	MSBST04DSE20- DEMOGRAPHIC STUDIES				
Title					
Programme	M.Sc. Biostatistics Semester IV				
Course	• To introduce students to key concepts and t	heories in			
Objectives	demography.				
	• To provide students with an understanding	of demograp	hic data		
	sources and measurement techniques.				
	• To familiarize students with demographic methods for analyzing				
5	population dynamics.				
- A.	• To demonstrate the applications of demographic analysis in				
1	various fields, including public health, economics, and social				
<1/	policy.				
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Modules	Content	Module Outcome

	Definition and scope of demography,	•	Gain a comprehensive
	Historical development of		understanding of demographic
	demography, Importance and		concepts such as population
	applications of demographic research,		size, composition, fertility,
Module I	Population Size and Composition,		mortality, migration, and
Scope of	Measures of population size:	8.	population aging.
demography	population counts, estimates, and	•	Learn to interpret and analyze
(15 Hours)	projections, Population composition:		demographic measures
	age structure, sex ratio, and		including birth rates, death
	demographic characteristics,		rates, life expectancy,
	Interpretation of population pyramids		migration flows, and
			population pyramids.
	Fertility: Measures of fertility: birth		
	rates, total fertility rates, age-specific	•	Acquire proficiency in
	fertility rates, Determinants of	6	identifying and utilizing
Module II	fertility: socioeconomic, cultural, and		various sources of
Measures of	policy factors, Trends and patterns in		demographic data, including
mortality and	fertility, Mortality: Measures of		censuses, surveys, and vital
fertility	mortality: death rates, life		registration systems.
(15 Hours)	expectancy, age-specific mortality	•	Learn data quality assessment
	rates, Causes of mortality: infectious		techniques and sampling
S.	diseases, chronic diseases, external		methods for demographic
- A	causes, Epidemiological transition	7	research
	theory	1	
<1/	Migration: Types of migration:	1	Develop skills in demographic
dia	internal, international, refugee	2	analysis techniques, including
Module III	movements, Measures of migration:	/	standardization methods and
Theories of	net migration rates, migration flows,		demographic modeling
migration	migration stocks, Theories of		approaches such as cohort-
(15 Hours)	migration: push and pull factors,		component projection and
	network theory, migration systems		survival analysis.
	Population Aging, Concepts and	•	Apply statistical methods to
	measures of population aging,		analyze demographic trends

	Causes and consequences of and patterns, and interpret
	population aging, Challenges and findings accurately.
	opportunities of an aging population
	Demographic Data Sources and
	Methods; Sources of demographic • Explore historical trends in
	data: censuses, surveys, vital population dynamics and
Module IV	registration systems, Data quality understand the underlying
Demographic	issues and sampling techniques causes and consequences of
Analysis	Demographic Analysis Techniques demographic changes.
Techniques	Standardization techniques, • Analyze contemporary
	Demographic modeling: cohort- population trends, including
(15 Hours)	component projection method, fertility, mortality, migration,
	population momentum, Survival and population aging, and
	analysis techniques for life tables and assess their implications for
	mortality data, Population Policy and society.
	Planning.
	Text Book
	1. Poston Jr, D. L., & Bouvier, L. F. (2010). Population and society: An
	introduction to demography. Cambridge University Press.
	2. Cox PR (1957). Demography. Cambridge University Press
	Reference Book
References	1. Croxton F E and Crowder D J (1967) Applied General statistics, Prentice
- A	- Hall India.
	2.Bogue, Donald J: Principles of Demography, John Wiley and Sons, New
<1/	York,1969
1	3. Shrivastava O S: A Text Book of Demography with Economics of Man
	Power Supply and Manpower Demand, Vikas, New Delhi, 1983
Course	After successful completion of this course, student will be able to:
Outcomes	1. Define and explain fundamental demographic concepts such as
	population size, composition, fertility, mortality, migration, and
	population aging.
	2. Identify and utilize various sources of demographic data, including
	censuses, surveys, and vital registration systems, and assess data quality.

3. Apply demographic analysis techniques, including standardization
methods and demographic modeling approaches, to analyze population
dynamics and trends.
4. Interpret demographic measures and trends accurately, and assess their
implications for social, economic, and political contexts.

• Lecturing, Visualization, Team Learning

#### **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

#### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous Ev	aluation
Tests	16
Assignment	08
Seminar/Viva	16

- 1. Define demography and explain its significance in the field of statistics.
  - 2. Describe the measures used to quantify population size and composition. How are these measures calculated?
  - 3. Discuss the factors influencing fertility rates and their variations across different populations.
  - 4. Compare and contrast crude birth rate, total fertility rate, and age-specific fertility rates. Provide examples illustrating their use.
  - 5. Explain the concept of the epidemiological transition and its implications for population health.

- 6. Discuss the main determinants of mortality rates and how they have changed over time.
- Describe the different types of migration and their impact on population dynamics. Provide examples of push and pull factors influencing migration.
- 8. Define population aging and discuss its causes and consequences for societies.
- 9. Explain the cohort-component projection method and its application in population forecasting.
- 10. Discuss the role of demography in informing public policy decisions related to healthcare, labor markets, and environmental planning.

Course Code & Title	MSBST04DSE21- ANALYSIS OF LONGITUDINAL DATA
Programme	M.Sc. Biostatistics Semester IV
Course	Master advanced statistical modeling techniques for longitudinal
Objectives	data.
	• Develop proficiency in estimating model parameters using
	maximum likelihood (ML), restricted maximum likelihood
	(REML).
	• Gain a comprehensive understanding of missing data mechanisms
	and strategies for handling missing values in longitudinal studies.
5	• Learn to address challenges such as time-dependent covariates,
- A.	intermittent missing values, and dropout processes in longitudinal
>	data analysis.

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Modules	Content	Module Outcome
	General Linear Model for	• Explore maximum likelihood
Module I	Longitudinal Data. ML and	(ML) and restricted maximum
General Linear	REML estimation, EM algorithm:	likelihood (REML) estimation
Model for	General linear mixed-effects	methods.
Longitudinal	model, Inference for; the random	• Understand the expectation-

Data	effects, BLUPs, Empirical Bayes,	maximization (EM) algorithm.
(15 Hours)	Bayes, Shrinkage Model building	• Perform inference for random
	and diagnostic, relaxing parametric	effects.
	assumptions: generalized additive	
	mixed model.	
Module II Random effects models for binary and count data (15 Hours)	Generalized Linear Model for Longitudinal Data, Marginal models, for binary, ordinal, and count data: Random effects models for binary and count data: Transition models: Likelihood- based models for categorical data; GEE; Models for mixed discrete and continuous responses.	<ul> <li>Extend the framework to handle generalized linear models (GLMs) for longitudinal data.</li> <li>Study marginal models for binary, ordinal, and count data, as well as random effects models.</li> <li>Explore transition models and likelihood-based approaches for categorical data</li> </ul>
Module III Modeling the dropout process (15 Hours)	Classification missing data mechanism; Intermittent missing values and dropouts; Weighted estimating equations; Modeling the dropout process (Selection and pattern mixture models).	<ul> <li>Investigate classification of missing data mechanisms and strategies for addressing intermittent missing values and dropouts.</li> <li>Learn about weighted estimating equations and modeling the dropout process.</li> </ul>
Module IV Multivariate longitudinal data (15 Hours)	Dangers of time dependent covariates, Lagged covariates; Marginal Structural models; Joint models for longitudinal and survival data; Multivariate longitudinal data; Design of randomized and observational longitudinal studies.	<ul> <li>Address challenges associated with time-dependent covariates and lagged covariates.</li> <li>Explore marginal structural models and joint models for longitudinal data.</li> <li>Discuss strategies for analyzing multivariate</li> </ul>

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	longitudinal data.	
	Text books	
	1. Diggle, P. J., Heagerty, P., Liang, K. Yand Zeger.S. L (2003	3). Analysis
	of Longitudinal Data, 2 <sup>nd</sup> Edn. Oxford University press, New	w York.
	2. Fitzmaurice, G.M., Laird, N.M and Ware, J.H. (2004). Appli	ed
	Longitudinal Analysis, John Wiley & Sons, New York.	
	Reference Books	
	1. Crowder, M. J. and Hand, D. J. (1990). Analysis of	of Repeated
	Measures. Chapman and Hall/CRC Press, London.	
Defense	2. Davidian, M. and Giltinan, D. M. (1995). Nonlinear	Models for
Kelefences	Repeated Measurement Data. Chapmanand Hall/CRC Pres	s, London.
	3. Hand, D and Crowder, M. (1996). Practical Longitu	ıdinal Data
	Analysis. Chapman and Hall/CRC Press, New York.	
	4. Little, R. J. A and Rubin, O. B.(2002). Statistical A	nalysis with
	Missing Data, 2 <sup>nd</sup> Edition, Wiley, New York.	
	5. McCullagh, P. and Nelder. J. A. (1989). Generalized Lin	ear Models.
	2nd Edition, Chapman and Hall/CRC Press, London.	
	6. Weiss, R. E. (2005). Modeling Longitudinal Data. Springer	r, New York
~	After successful completion of this course, student will be a 1. Conduct analysis of longitudinal data.	to:
- A		
Course	2. Apply statistical techniques to model longitudinal data	and
Outcomes	make predictions.	
outcomes	3. Understand analysis of longitudinal data with missing of	lata.
100	<ol> <li>Understand analysis of longitudinal data with time-dependent covariates.</li> </ol>	endent

• Lecturing, Visualization, Team Learning

## **MODE OF TRANSACTION**

• Lecture, Seminar, Discussion, Questioning and Answering

#### **ASSESSMENT RUBRICS**

Components	Weightage
End Semester Evaluation(ESE)	60
Continuous E	valuation
Tests	16
Assignment	08
Seminar/Viva	16
Total	40

- 1. What distinguishes maximum likelihood (ML) from restricted maximum likelihood (REML) estimation in the context of longitudinal data analysis?
- 2. Explain the concept of the expectation-maximization (EM) algorithm and how it is utilized in fitting general linear mixed-effects models for longitudinal data.
- 3. How would you handle intermittent missing values in a longitudinal dataset? Describe the strategies and techniques you would employ.
- 4. Discuss the advantages and limitations of using generalized estimating equations (GEE) for analyzing longitudinal data compared to mixed-effects models.
- 5. What are the key differences between marginal models and random effects models for handling binary data in longitudinal studies?
- How do you assess the impact of time-dependent covariates on longitudinal outcomes?
   Describe the statistical methods used for this analysis.
- 7. Explain the concept of best linear unbiased predictions (BLUPs) and their relevance in estimating random effects in longitudinal models.
- 8. Describe the process of building a generalized additive mixed model (GAMM) for longitudinal data and discuss its advantages over parametric approaches.
- 9. What are selection models and pattern mixture models, and how are they used to address the issue of dropout in longitudinal studies?

10. Can you outline the steps involved in designing a longitudinal study, including considerations for handling missing data and analyzing multivariate longitudinal outcomes?



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