

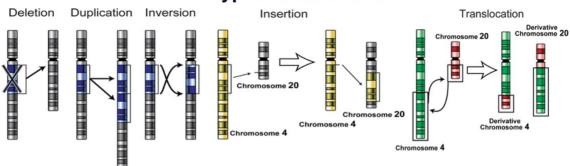
STUDY GUIDE

GENETICS-II MODULE

FOURTH YEAR MBBS

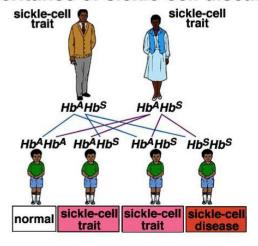
13th Feb - 29th Feb 2020 Duration: 2.5 weeks

Types of Mutations



Chromosome Nucleosome Nucleotide base pairs: Guanine Cytosine Adenine Thymine

Inheritance of sickle-cell disease







STUDY GUIDE FOR GENETICS-II MODULE

S.No	CONTENTS	Page No.
1	Overview	3
2	Introduction to Study Guide	4
3	Learning Methodologies	5
4	Module 3: GENETICS-II	7
4.1	Introduction	7
4.2	Objectives and Learning Strategies	8
5	Learning Resources	10
5.1	Additional Learning Resources	10
6	Assessment Methods	11
7	LNMC Examination Rules and Regulations	12
8	Schedule	13

Module name: Genetics-II Year: Four Duration: 2.5 weeks (Feb 2020)

Timetable hours: Interactive Lectures, Case-Based Discussion (CBD), Clinical Rotations,

Presentations, Demonstrations, Skills, Self-Study

MODULE INTEGRATED COMMITTEE

MODULE COORDINATOR:	Dr. Kashif Nisar (Biochemistry)
CO-COORDINATOR:	Dr. M. Suleman Sadiq (DHCE)

DEPARTMENTS' & RESOURCE PERSONS' FACILITATING LEARNING

BASIC HEALTH SCIENCES	CLINICAL AND ANCILLARY DEPARTMENTS
BIOCHEMISTRY	MOLECULAR PATHOLOGY
• Dr. Kashif Nisar	 Dr. Muhammad Israr Nasir
	Dr. Sobia Rafiq
PATHOLOGY	
 Prof. Naveen Faridi 	
PHARMACOLOGY	
 Prof. Nazir Solangi 	
 Prof. Tabassum Zehra 	

DEPARTMENT OF HEALTH PROFESSIONS EDUCATION

- Professor Nighat Huda
- Dr. Sobia Ali
- Dr. Afifa Tabassum

- Dr. M. Suleman Sadiq
- Dr. Mehnaz Umair

LNH&MC MANAGEMENT

- Professor Karimullah Makki, Principal, LNH&MC
- Dr. Shaheena Akbani, Director A.A & R.T LNH&MC

STUDY GUIDE COMPILED BY: Department of Health Professions Education

INTRODUCTION

WHAT IS A STUDY GUIDE?

It is an aid to:

- Inform students how student learning program of the module has been organized
- Help students organize and manage their studies throughout the module
- Guide students on assessment methods, rules and regulations

THE STUDY GUIDE:

- Communicates information on organization and management of the module.
 This will help the student to contact the right person in case of any difficulty.
- Defines the objectives which are expected to be achieved at the end of the module.
- Identifies the learning strategies such as lectures, small group teachings, clinical skills, demonstration, tutorial and case based learning that will be implemented to achieve the module objectives.
- Provides a list of learning resources such as books, computer assisted learning programs, weblinks, journals, for students to consult in order to maximize their learning.
- Highlights information on the contribution of continuous on the student's overall performance.
- Includes information on the assessment methods that will be held to determine every student's achievement of objectives.
- Focuses on information pertaining to examination policy, rules and regulations.

CURRICULUM FRAMEWORK

Students will experience integrated curriculum similar to previous modules.

INTEGRATED CURRICULUM comprises system-based modules such as Eye/ENT, dermatology, genetics, rehabilitation and neurosciences-II & psychiatry modules which link basic science knowledge to clinical problems. Integrated teaching means that subjects are presented as a meaningful whole. Students will be able to have better understanding of basic sciences when they repeatedly learn in relation to clinical examples.

LEARNING EXPERIENCES: Case based integrated discussions, Task oriented learning followed by task presentation, skills acquisition in skills lab, computer-based assignments, learning experiences in clinics, wards.

PHARMACOLOGY GENETICS-II MOLECULAR PATHOLOGY PATHOLOGY

INTEGRATING DISCIPLINES OF GENETICS-II MODULE

LEARNING METHODOLOGIES

The following teaching / learning methods are used to promote better understanding:

- Interactive Lectures
- Small Group Discussion
- Case- Based Discussion (CBD)
- Clinical Experiences
 - o Clinical Rotations
- Skills session

INTERACTIVE LECTURES: In large group, the lecturer introduces a topic or common clinical conditions and explains the underlying phenomena through questions, pictures, videos of patients' interviews, exercises, etc. Students are actively involved in the learning process.

SMALL GROUP SESSION: This format helps students to clarify concepts, acquire skills or desired attitudes. Sessions are structured with the help of specific exercises such as patient case, interviews or discussion topics. Students exchange opinions and apply knowledge gained from lectures, tutorials and self study. The facilitator role is to ask probing questions, summarize, or rephrase to help clarify concepts.

CASE-BASED DISUCSSION (CBD): A small group discussion format where learning is focused around a series of questions based on a clinical scenario. Students' discuss and answer the questions applying relevant knowledge gained previously in clinical and basic health sciences during the module and construct new knowledge. The CBD will be provided by the concern department.

CLINICAL LEARNING EXPERIENCES: In small groups, students observe patients with signs and symptoms in hospital wards, clinics and outreach centers. This helps students to relate knowledge of basic and clinical sciences of the module and prepare for future practice.

CLINICAL ROTATIONS: In small groups, students rotate in different wards like Medicine, Pediatrics, Surgery, Obs & Gyne, ENT, Eye, Family Medicine clinics, outreach centers & Community Medicine experiences. Here students observe patients, take histories and perform supervised clinical examinations in outpatient and inpatient settings. They also get an opportunity to observe medical personnel working as a team. These rotations help students relate basic medical and clinical knowledge in diverse clinical areas.

SKILLS SESSION: Skills relevant to respective module are observed and practiced where applicable in skills laboratory.

SELF-DIRECTED STUDY: Students' assume responsibilities of their own learning through individual study, sharing and discussing with peers, seeking information from Learning Resource Center, teachers and resource persons within and outside the college. Students can utilize the time within the college scheduled hours of self-study.

MODULE 3: GENETICS II

INTRODUCTION

A genetic disorder is a disease that is caused by a change, or mutation, in an individual's DNA sequence. These mutations can be due to an error in DNA replication or due to environmental factors, such as cigarette smoke and exposure to radiation, which cause changes in the DNA sequence.

Internationally, 3-5% of all births result in congenital malformations; 20-30% of all infant deaths are due to genetic disorders; 30-50% of post-neonatal deaths are due to congenital malformations. 11.1% of pediatric hospital admissions are for children with genetic disorders and 18.5% are children with other congenital malformations; 12% of adult hospital admissions are for genetic causes. 50% of mental retardation, a common global occurrence, has a genetic basis. Cancers are one of the most dreaded conditions; 15% of all cancers have an inherited susceptibility whereas 10% of chronic diseases (heart, diabetes arthritis) which occur in the adult population have a significant genetic component.

Pakistan has a high frequency of marriages among close cousins, i.e. consanguineous marriages. This percentage is 62.70 — by far the highest among countries in the 'consanguinity belt' which includes countries of the Middle East and the Sub continent. It is estimated that about 29 million people out of Pakistan's 200 million population suffer from genetic defects attributable to close or first-cousin marriages.

Hence, it becomes imperative for medical graduates in Pakistan to understand how such conditions occur and how they can be managed and prevented. Since genetics, and the pathophysiology of genetic disorders, is a complex process, this topic was initially dealt with in the first spiral of the curriculum at a simpler level and is now being re-visited at a more advanced level. In the first year, normal genetic processes were described so that learners get a clear understanding of how chromosomes function.

In this 2nd spiral of Genetics, students will learn about the process of mutations, single gene defects and techniques used to diagnose such disorders. You may experience various genetic disorders during the clinical rotation and clerkship.



Reference: JSMU Study guide dated May 10th, 2018

COURSE OBJECTIVES AND STRATEGIES

By the end of Genetics-II module students should be able to:

BIOCHEMISTRY 1. Describe the process of DNA Replication and repair 2. Explain the mechanism of Transcription and Post Transcriptional modification 3. Discuss the process of Translation and Post Translational modification MOLECULAR PATHOLOGY 1. Discuss the basic concepts of genetics including DNA and RNA structure, Mendel's Laws of inheritance and Pedigree Chart Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis 14. Justify indications for and utility of prenatal diagnosis tests		OBJECTIVES	TEACHING STRATEGIES
2. Explain the mechanism of Transcription and Post Transcriptional modification 3. Discuss the process of Translation and Post Translational modification MOLECULAR PATHOLOGY 1. Discuss the basic concepts of genetics including DNA and RNA structure, Mendel's Laws of inheritance and Pedigree Chart Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List the advantages and disadvantages of prenatal diagnosis Interactive Lectures			
2. Explain the mechanism of Transcription and Post Transcriptional modification 3. Discuss the process of Translation and Post Translational modification MOLECULAR PATHOLOGY 1. Discuss the basic concepts of genetics including DNA and RNA structure, Mendel's Laws of inheritance and Pedigree Chart Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List the advantages and disadvantages of prenatal diagnosis Interactive Lectures Interactive Lectures/Sma Group Discussion/Interactive Lectures Interactive Lectures/Sma Group Discussion/Interactive Lectures Interactive Lectures Small Group Discussion/Interactive Lectures	1.	Describe the process of DNA Replication and repair	
modification 3. Discuss the basic concepts of genetics including DNA and RNA structure, Mendel's Laws of inheritance and Pedigree Chart Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis Interactive Lectures Interactive Lectures and Therefore Company of the protein it encodes Single gene disorders Small Group Discussion/Interactive Lectures Interactive Lectures Interactive Lectures Interactive Lectures Interactive Lectures Interactive Lectures Interactive Lectures	2.		
3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis Interactive Lectures		·	<u>-</u>
modification MOLECULAR PATHOLOGY 1. Discuss the basic concepts of genetics including DNA and RNA structure, Mendel's Laws of inheritance and Pedigree Chart Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	3.	Discuss the process of Translation and Post Translational	Group Discussion
1. Discuss the basic concepts of genetics including DNA and RNA structure, Mendel's Laws of inheritance and Pedigree Chart Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures		·	
Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List the advantages and disadvantages of prenatal diagnosis Interactive Lectures		MOLECULAR PATHOLOGY	
Mutations 2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Small Group Discussion/Interactive Lectures Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures	1.	Discuss the basic concepts of genetics including DNA and RNA structure,	
2. List the different types of mutations in the coding and non-coding regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures			
regions of genes 3. Distinguish between the different types of mutations in the coding and non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	Mu		
non-coding regions of genes that result in phenotypic change 4. Differentiate between spontaneous and induced mutations 5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	2.		
5. Explain how a point mutations or frameshift mutation in a gene may alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	3.		
alter the activity of the protein it encodes Single Gene Mutation 6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	4.	Differentiate between spontaneous and induced mutations	
6. Define single gene disorders 7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	5.	· · · · · · · · · · · · · · · · · · ·	Interactive Lectures
7. List different types of single gene disorders 8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	Sing	le Gene Mutation	
8. List characteristics of single gene which gives variation in expression of diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	6.	Define single gene disorders	
diseases 9. Describe genetic changes which occur in these disorders Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis Interactive Lectures Interactive Lectures	7.	List different types of single gene disorders	
Genetic Technique 10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis Interactive Lectures	8.		
10. Discuss the basic principles of recombinant genetic techniques and their applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis Interactive Lectures	9.	Describe genetic changes which occur in these disorders	
applications in the detection of genetic diseases which includes PCR, FISH, RFLP, BLOTTING Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis Interactive Lectures	Gen	etic Technique	
Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis	10.	Discuss the basic principles of recombinant genetic techniques and their	Small Group
Prenatal Diagnosis 11. Define prenatal diagnosis 12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis		applications in the detection of genetic diseases which includes PCR,	Discussion/Interactive
Define prenatal diagnosis List different techniques used for prenatal diagnosis Interactive Lectures State of the advantages and disadvantages of prenatal diagnosis		FISH, RFLP, BLOTTING	Lectures
12. List different techniques used for prenatal diagnosis 13. List the advantages and disadvantages of prenatal diagnosis	Pre	natal Diagnosis	
13. List the advantages and disadvantages of prenatal diagnosis	11.	Define prenatal diagnosis	
	12.	List different techniques used for prenatal diagnosis	Interactive Lectures
14. Justify indications for and utility of prenatal diagnostic tests	13.	List the advantages and disadvantages of prenatal diagnosis	
	14.	Justify indications for and utility of prenatal diagnostic tests	

OBJECTIVES	TEACHING STRATEGIES	
Gene Therapy & Counseling		
15. Describe how gene therapy works		
16. Describe how gene therapy works	Interactive Lectures	
17. Explain the tools and techniques used to deliver gene therapies, to the		
disease candidates for gene therapy and associated risks and challenges		
PATHOLOGY		
Pathophysiology of Inheritance		
18. Explain the pathophysiology of classical and non-classical mode of		
inheritance of genetic diseases	Interactive Lectures	
19. Discuss the clinical features of important genetic disorders which		
includes Down's syndrome, Turner's syndrome, Cystic Fibrosis, Sickle Ce	Small Crave Discussion	
Anemia, Thalassemia	Small Group Discussion	
PHARMACOLOGY		
 Identify the different components of medical prescription and able to write the medical prescription. 	Small Group Discussion	

Apart from attending daily scheduled sessions, students too should engage in self-study to ensure that all the objectives are covered



LEARNING RESOURCES

SUBJECT	RESOURCES
BIOCHEMISTRY	1. Harper's Illustrated Biochemistry 2. Lehninger Principle of Biochemistry 3. Biochemistry by Devlin
PATHOLOGY	TEXT BOOKS 1. Robbins & Cotran, Pathologic Basis of Disease, 9th edition. 2. Rapid Review Pathology, 4th edition by Edward F. Goljan MD
	WEBSITES: 1. http://library.med.utah.edu/WebPath/webpath.html 2. http://www.pathologyatlas.ro/
MOLECULAR PATHOLOGY	REFERENCE BOOK: 1. Thompson & Thompson Genetics in Medicine 8th Edition

ADDITIONAL LEARNING RESOURCES

Hands-on Activities/ Practical	Students will be involved in Practical sessions and hands-on activities that link with the Genetics-II Module to enhance learning.
<u>Museum</u>	Models available in the museum are a rich learning resource for quick review of anatomy and related educational activities
<u>Skills Lab</u>	Skills acquisition in a simulated environment in the skills lab involving experiential learning will ensure patient safety and will also help to build confidence in approaching the patients
<u>Videos/Podcasts</u>	Videos and podcasts will familiarize the student with the procedures and protocol which they can watch and listen to at any time and wherever they are, as part of task oriented learning
Internet Resources	Students will use easily accessible internet resources with added time flexibility to enrich and update their knowledge and its application

ASSESSMENT METHODS:

- Best Choice Questions(BCQs) also known as MCQs (Multiple Choice Questions)
- Objective Structured Practical/Clinical Examination (OSPE or OSCE)

BCQs:

- A BCQ has a statement or clinical scenario of four options (likely answers).
- Correct answer carries one mark, and incorrect 'zero mark'. There is NO negative marking.
- Students mark their responses on specified computer-based sheet designed for LNHMC.

OSCE:

- All students rotate through the same series of stations in the same allocated time.
- At each station, a brief written statement includes the task. Student completes the given task at one
 given station in a specified time.
- Stations are observed, unobserved, interactive or rest stations.
- In unobserved stations, flowcharts, models, slide identification, lab reports, case scenarios may be used to cover knowledge component of the content.
- Observed station: Performance of skills /procedures is observed by assessor
- Interactive: Examiner/s ask questions related to the task within the time allocated.
- In Rest station, students in the given time not given any specific task but wait to move to the following station.

Internal Evaluation

- Students will be assessed comprehensively through multiple methods.
- 20% marks of internal evaluation will be added to JSMU final exam. That 20% may include class tests, assignment, practicals and the internal exam which will all have specific marks allocation.

Formative Assessment

Individual department may hold quiz or short answer questions to help students assess their own learning. The marks obtained are not included in the internal evaluation

For JSMU Examination Policy, please consult JSMU website!

More than 75% attendance is needed to sit for the internal and final examinations

LNH&MC EXAMINATION RULES & REGULATIONS

- Student must report to examination hall/venue, 30 minutes before the exam.
- Exam will begin sharp at the given time.
- No student will be allowed to enter the examination hall after 15 minutes of scheduled examination time.
- Students must sit according to their roll numbers mentioned on the seats.
- Cell phones are strictly not allowed in examination hall.
- If any student is found with cell phone in any mode (silent, switched off or on) he/she will be not be allowed to continue their exam.
- No students will be allowed to sit in exam without University Admit Card, LNMC College ID Card and Lab Coat
- Student must bring the following stationary items for the exam: Pen, Pencil, Eraser, and Sharpener.
- Indiscipline in the exam hall/venue is not acceptable. Students must not possess any written material or communicate with their fellow students.

SCHEDULE:

WEEKS	4TH YEAR	MONTH
WEEK 1 -8	ORTHOPEDICS & REHABILITATION	2 nd Dec 2019
		25 th Jan 2020
WEEK 1	DEDIALTOLOGY	27 th Jan 2020
WEEK 2.5	DERMATOLOGY	12 th Feb 2020
WEEK 1	GENETICS-II	13 th Feb 2020
WEEK 2.5	GENETICS-II	29 th Feb 2020
1 WEEK	Extracurricular Activities	2 nd March 2020 - 7 th March 2020
1 WEEK	Revision Classes (Earlier Modules)	9 th March 2020 – 14 th March 2020*
	PREPARATORY LEAVE	16 th March 2020 – 19 th March 2020*
	MID TERM EXAM	20 th March 2020 – 21 st March 2020*

^{*}Final dates will be announced later