

SRI BALAJI VIDYAPEETH (DEEMED TO BE UNIVERSITY) U/S 3 OF UGC ACT 1956 Puducherry–607402 Accredited by NAAC with 'A' Grade

MD PATHOLOGY

POST GRADUATE CURRICULUM

For course conducted in

MAHATMA GANDHI MEDICAL COLLEGE AND

RESEARCH INSTITUTE

And

SRI SATYA SAI MEDICAL COLLEGE AND RESEARCH

INSTITUTE



Preface 1

The promulgation of the much-awaited Competency Based Medical Education (CBME) for post graduate programs by the National Medical Council is a welcome move. Sri Balaji Vidyapeeth (SBV), Puducherry, deemed to be University, declared u/s 3 of the UGC Act. and accredited by the NAAC with A grade, takes immense privilege in preparing such an unique document in a comprehensive manner and most importantly the onus is on the Indian setting for the first time, with regard to the competency based medical education for post graduate programs that are being offered in the broad specialty departments. SBV is committed to making cardinal contributions that would be realised by exploring newer vistas. Thus, post graduate medical education in the country could be made to scale greater heights and SBV is poised to show the way in this direction.

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Preface 2

The National Medical Council has laid down the PG curricula in their website https://www.nmc.org.in/information-desk/for-colleges/pg-curricula-2 that is listing the syllabus course wise, listing competency to some extent, teaching learning methods and the assessment methods as well. The document describes competencies in three domains (knowledge, skill, and attitude). However, the most significant problem in competency-based training is the development of appropriate assessment tools.

The salient feature of this document is defining the program educational objectives (PEO) for its postgraduate program as a whole, defining program outcomes (PO) based on the competencies to be practiced by the specialist, course outcomes (CO) and program specific sub-competencies and their progression in the form of milestones. The compilation of the milestone description leads to the formation of the required syllabus. This allows the mentors to monitor the progress in sub-competency milestone levels. It also defines milestone in five levels, for each sub-competency. Although NMC has described three domains of competencies, the domain 'Attitude' is elaborated into 4 more competencies for ease of assessment. The six competency model (ACGME) for residency education: Medical Knowledge, Patient Care, Practice Based Learning and Improvement, Systems Based Practice, Professionalism, Inter personal and Communication Skills gives better clarity and in-depth explanation and is used in this document. The sub-competency and their milestone levels are mapped into the entrustable professional activities(EPA) that are specific to the individual postgraduate program. While doing all this, the syllabus prescribed by NMC is fully incorporated into the curriculum. To make the program more relevant, PEO, PO, CO and EPAs are mapped with each other. EPAs which are activity based are used for formative assessment and graded. EPA assessment is based on workplace based assessment (WPBA), multisource feedback (MSF) and eportfolio. A great emphasis is given on monitoring the progress in acquisition of knowledge, skill and attitude through various appraisal forms including e-portfolios during three years of residency period.

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Foreword

The Postgraduate curriculum in Pathology has evolved into a Competency Based Medical Education which provides an objective mode of teaching and evaluation. Based on the MCI guidelines the curriculum is modified to accommodate program outcomes, course outcomes and thereby achieving the Program Educational Objectives.

The Entrustable Professional Activities are defined with the levels of achievement for individual postgraduate student for each phase of their program, thereby enabling self assessment and improvement as well. The 360 degree feedback and e- portfolio will assist in learning and evaluation of all the three domains – cognitive, psychomotor and affective.

The new curriculum encompasses recent concepts and their applications in the core, which will provide exposure to the practical skills to be acquired.

Overall the postgraduates will get an opportunity to learn, practice, analyse and teach the basics of Pathology, during their tenure.

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1 List of Abbreviations and Acronyms

PEO	Programme Educational Objective
РО	Programme Outcome
CO	Course outcome
EPA	Entrustable Professional Activity
МК	Medical Knowledge
PC	Patient Care
SBP	System Based Practice
PBLI	Practice Based Learning and Improvement
IPCS	Interpersonal Communication Skills
Р	Professionalism

Sri Balaji Vidyapeeth

Post - Graduate Programme, MD Pathology

2 **Preamble**

The competency based curriculum should take into account the needs of the society, both local and global. It needs to outline the demand for the present day as well as future. The curriculum needs to be reviewed at least every five years to address the trending needs, as new knowledge is evolving and communication of the same is seamless. Accordingly the competencies need to meet the societal needs detailing the cognitive, psychomotor and affective domain development for attaining these competencies.

The curriculum indicates to the candidate the knowledge, basic skills and attitudes required to become a Pathologist. It disciplines the thinking habits for problem solving and discovery of new knowledge in the field of Insert speciality. It defines the Teaching - Learning methods adopted for the resident to achieve the goals and the methods of assessment performed throughout the training period and at the completion of training. The purpose of this document is to provide teachers and learners illustrative guidelines to achieve defined outcomes through learning and assessment.

3 Programme Educational Objective (PEO)

Programme Educational Objectives are broad statements that describe what graduates are expected to attain within few years of completing their programme. These are based on the needs of the society as analysed and outlined by the regulatory body. So as defined by NMC, the PEO for MD Pathology are as follows:

- PEO1: Pathologist who can provide all diagnostic information in the fraternity of laboratory medicine especially in Histopathology,Cytopathology, Clinical Pathology, Hematologyand Transfusion Medicine with a knowledge of general principles and methodology with interpretation of values.
- **PEO2:** Understands health care system and be a leader and as a part of a team, to develop an attitude of cooperation with colleagues, and interact with the patient and the clinician or other colleagues to provide the best possible diagnosis or opinion.
- **PEO3:** Communicator(Affective domain) Interpret and correlate clinical and laboratory data so that clinical manifestations of diseases can be explained.
- PEO4: Life long learner keen on updating oneself regarding the advancement in diagnostic field and capable to teach Pathology to undergraduates, postgraduates, nurses and paramedical staff including laboratory personnel.
 PEO5: Professional who understands and follows the principle of bio-ethics / ethics related to health care system and biomedical waste disposal, quality system and ensures good clinical laboratory practices

4 Programme Outcome (PO)

POs represent broad statements that incorporate many areas of inter - related knowledge and skills developed over the duration of the programme through a wide range of courses and experiences. They represent the big picture and describe broad aspects of knowledge, skill and attitude development. They encompass multiple learning experiences.

After a period of 3 years, the resident should be able to attain the following PO's:

- **PO1:** Provide a definitive diagnosis in Histopathology, Cytology, Clinical Pathology and Hematology.
- **PO2:** Practice safe blood transfusion practices with management of its adverse reactions.
- **PO3:** Interpret all routine diagnostic tests and handle in trouble shootings.
- **PO 4:** Identify patient safety and system approach to different errors in laboratory in phases like pre-analytical, analytical and post-analytical.
- **PO5:** Communicate with colleagues, faculty and stake holders of the health care system.
- **PO6:** Update with recent advances and Develop communication skills to word reports and professional opinion as well as to educate students and interact with patients, peers and paramedical staff, and for effective teaching.
- **PO7:** Maintain proper etiquette in dealings with patients, relatives and other health personnel and to respect the rights of the patient including the right to information and second opinion.
- **PO8:** Perform Self Directed Learning and Critical appraisal of medical literature.
- **PO9:** Develop & execute a protocol for a scientific research project, collect and analyze the data and scientifically communicate to the others.

5 Course and Course Outcomes (CO)

CO's describe the learning that will take place across the curriculum through concise statements, made in specific and measurable terms, of what students will know and /or be able to do after successful completion of each course.

There are four courses for MD Pathology:

- Course 1 (C1) : General Pathology, Pathophysiology, Immunopathology and Cytopathology
- 2. Course 2 (C2): Systemic Pathology.
- Course 3 (C3): Haematology, Transfusion Medicine (Blood Banking) and Laboratory Medicine.
- 4. Course 4 (C4): Recent advances and Applied aspects.

5.1 Course 1 (C1) (General Pathology, Pathophysiology,

ImmunoPathology and Cytology)

C1.1.Aquire Basic knowledge about the General Pathology.

C1.2 Be Familiar with the concepts of Pathophysiology and Immunopathology.

C1.3 Possess background knowledge about Cytopathology and able to perform its techniques.

C1.4 Undergo Basic Course in Biomedical Research, Data collection and analysis, Scientific communication.

5.2 Course 2 (C2) (Systemic Pathology)

C2.1 Capable of performing a systematic gross examination of the tissues and demonstrate the orientation of tissues in paraffin blocks.

C2.2 Identify and systematically and accurately describe the chief histomorphological alterations in the tissue received in the surgical pathology service.

C2.3 Conversant with automatic tissue processing machine and the principles of its running.

C2.4 Capable of Tissue processing, make a paraffin block and cut sections of good quality on a rotary microtome and also able to stain the tissue with various stains.

C2.5 Capable to do frozen section using cryostat, to stain and interpret the slide in correlation with the clinical data provided.

C2.6 Able to understand the utility of various immunohistochemical stains especially in the diagnosis of tumour subtypes.

C2.7 Capable of performing the Adult and Fetal autopsy.

5.3 Course 3 (C3) (Hematology, Transfusion medicine,

Laboratory medicine)

C3.1 Capable of performing various routine and special haematological test and also molecular investigation related to hematology.

C3.2 Capable of performing various clinical pathology investigations like urine analysis, fluid examination and semen analysis.

C3.3 Capable of performing various laboratory medicine investigation and good clinical laboratory practice (GCLP).

C3.4 Capable of Donor selection and drawing blood from donor, Prepration of components and blood grouping.

C3.5 Capable of performing Antenatal and Neonatal work up and other work up related to transfusion services.

5.4 Course 4 (C4): Recent Advances and Applied aspects

C4.1 Demonstrate familiarity with the principles and techniques of various advances in Pathology like Immunohistochemistry, Immunofluorescence and Electron microscopy.

C4.2 Possess the knowledge of various molecular techniques like PCR and FISH.

C4.3 Demonstrate familiarity with the principles of Biostatistics.

C4.4 Possess knowledge about Biomedical waste management.

5.5 Mapping of PEO, PO and CO

Programme mapping facilitates the alignment of course - level outcomes with programme outcomes. It allows faculty to create a visual map of a programme. It is also used to explore how students are meeting program - level outcomes at the course level. Outcomes mapping focuses on student learning also.

	PE	01	PEO1PEO3	PEO1	PEO2 PEO3	PEO2 PEO3 PEO4 PEO5	PEO2 PEO4	PEO4 PEO5	PEO5
	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9
C1	Y		Y	Y	Y		Y	Y	Y
C2	Y		Y	Y	Y	Y	Y	Y	Y
C3	Y	Y	Y	Y	Y	Y	Y	Y	Y
C4					Y	Y		Y	Y

Table1. Mapping of PEO, PO and CO

All courses run concurrently for 3 years, with a summative assessment at the end.

6 Competencies, Sub - competencies and milestones

The post graduate programme is competency based, consisting of six domains of competency. Sub - competencies under these domains, specific to the speciality, have been mentioned in general terms. The progression through the curriculum is detailed in sub - competency milestone levels, that direct the prescribed syllabus. These sub - competency milestones are mapped to the Entrustable Professional Activities (EPAs), identified as essential for a specialist. Formative assessment includes EPA assessment, and is carried out every quarter using appropriate tools, for identifying eligibility for transfer of trust, to the resident.

6.1 **Domain of Competencies**

- 1. **Medical Knowledge** (**MK**)–Acquiring Knowledge of established and evolving biomedical, clinical, epidemiological, and social behavioural sciences and the application of this knowledge to patient care.
- 2. **Patient Care/Procedural Skill (PC/PS)**–Demonstrate ability to provide patient centred care/demonstrate skills required for teaching and conducting research.
- 3. **System Based Practise (SBP)** Demonstrate the ability to follow the standard operating procedures relevant to practices of the organisations for patient care, inculcating quality and economical practices.
- 4. **Practice Based Learning and improvement (PBLI)** Demonstrate the commitment to learn by literature search, feedback, practice and improve upon their ability.
- 5. **Interpersonal Communication skills (IPCS)** Demonstrate behaviour and skills that result in the effective communication, exchange of information and cooperation with patients, their families, and health professionals
- 6. **Professionalism** (**P**) Demonstrate a commitment to carrying out professional responsibilities and an adherence to ethical principles.

6.2 Sub - competencies

6.2.1 Medical Knowledge (MK)

Mk 1: knowledge specific to General pathology and Pathophysiology.

Mk 2: Knowledge required for Immunopathology.

Mk 3: Knowledge specific to Diagnostic Cyto-Pathology and Cytological techniques.

Mk 4: Knowledge required for Systemic Pathology.

Mk 5: Knowledge specific to Surgical Pathology.

MK6: Knowledge required for Autopsy Pathology.

Mk 7: Knowledge required for Haematology.

Mk 8: Knowledge specific to Blood Banking (Transfusion Medicine).

Mk 9: Knowledge required for principles and practices of Laboratory Medicine.

Mk 10: Knowledge specific to recent advances and applied aspects.

Mk 11: Knowledge required for presenting Seminars and Journal club.

6.2.2 Patient Care/ Procedural skill (PC/PS)

PS 1: Perform gross examination of surgical pathology specimens

PS2: Perform grossing in grossing station.

PS 3: Interpretation of Microscopic findings in tissue.

PS 4: Perform Histotechniques in Histopathology laboratory

PS 5: Perform And Interpretation of FNAC and other Cytological smears (PAP smear,

Sputum, Bronchial washings, Serous effusions, etc.

PS 6: Interpretation of Frozen sections

PS 7: Selection, performance and interpretation of appropriate Immuno-histochemical (IHC) markers.

PS 8: Interpretation of Bone Marrow Smears

PS 9: Perform and Interpretation of routine haematological investigations like

haemoglobin, TLC, DLC, ESR PCV, Blood indices and peripheral smear.

PS 10: Perform and Interpretation of special investigations like Reticulocyte count,

Sickling test, Osmotic Fragility Test, Haemoglobin Electrophoresis, Fetal Haemoglobin, etc.

PS 11: Planning investigations for a Clinical case

PS 12: Perform and Interpret Urine Examination, Body fluids and semen analysis

PS 13 : Interpretation of ancillary techniques like Immunofluorescence, Karyotyping, FISH, PCR and Electron Microscopy.

PS 14 : Demonstration of familiarity within laboratory investigations in Microbiology and Biochemistry

PS 15 : Perform and interpret Blood banking techniques (Blood grouping, Rh typing, Cross-matching and Coomb's test, ELISA for infectious disease)

PS 16 : Selection of blood donors and Management of adverse donor reactions.

PS 17 : Perform investigation for a case of mismatched blood transfusion.

PS 18 : Participation and Presentation in multidisciplinary meetings like tumor

boards, CPCs, Dermato-Pathological conferences.

PS 19 : Teaching pathology to undergraduates (MBBS), and allied health sciences like BDS, BSc (Nursing), BSc (MLT), BSc (Radiology), etc.

6.2.3 System Based Practice

IPCS 1: Communicate the reports effectively with clinician, patients and families.

IPCS 2: Communication with Colleagues and Technical staffs

IPCS 3: Teaching to Undergraduates students and other colleagues

6.2.4 Practice based learning and improvement

PBLI 1: Self-directed Learning/Critical Appraisal of Medical Literature.

6.2.5 Interpersonal communication skills

SBP 1: Laboratory safety and Systems. Approach to Laboratory Errors.

6.2.6 Professionalism

P 1: Compassion, Integrity, and Respect for Others

P 2: Feedback

6.3 Milestone Levels for Sub–competencies.

6.3.1 Medical Knowledge

MK 1: Knowledge specific to General pathology and Pathophysiology.

	Knowledge speci	fic to General pathology and P	athophysiology.	
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate a	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone
knowledge of normal	1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate an in-	level 4 Demonstrate the
anatomy, histology and	about inflammation and	about various genetic	depth knowledge regarding	knowledge regarding
functions of various	repair.	disorders and	environmental and	recent advances in
organs		pathophysiology of various	childhood disease.	Pathophysiology of
	Also demonstrate the	genetic diseases and		diseases.
Demonstrate the	knowledge about the	autoimmunity		
Knowledge of basic	mechanism of			
pathophysiology, cellular	Hypersensitivity and	Also demonstrate the		
adapataions and cell	Neoplasia.	knowledge about the various		
injury		infectious diseases		

MK 2: Knowledge required for Immunopathology.

	Knowledge required for Immunopathology.								
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5					
Demonstrate knowledge	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone					
of normal structure and	1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate in-	level 4 Demonstrate the					
function of immune	about classification and	about various autoimmune	depth knowledge regarding	knowledge regarding					
system.	mechanism of various	diseases.	interpretation of	recent advances in					
	Immunological diseases.		1.Immuno- electrophoresis,	Immunopathology.					
		Also demonstrate the	2 .Immunofluorescence						
	Also demonstrate the	knowledge about scope and	techniques especially on						
	knowledge about the	principles of various	kidney and skin biopsies,						
	mechanism of	immunological test like (a)	3.Anti-nuclear antibody and						
	Hypersensitivity and	ELISA techniques (b)	4.Anti-neutrophil						
	Neoplasia.	Radioimmunoassay	cytoplasmic antibody.						
		(c) HLA typing							

MK 3: Knowledge specific to Diagnostic Cyto-Pathology and Cytological techniques.

	Knowledge specific to Diagnostic Cyto-Pathology and Cytological techniques.								
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5					
Demonstrate the	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone					
knowledge of cytological	1	2 demonstrate the knowledge	level 3 should possess in-	level 4 Demonstrate the					
techniques and basic	Demonstrate the knowledge	about malignant cytology of	depth knowledge regarding	knowledge regarding					
cytology of various	about basic cyto-pathology of	various organs.	cytology of rare lesions and	recent advances in Cyto-					
organs.	reactive and benign lesion of		able to discuss the	Pathology.					
	various organs.	Demonstrate the familiarity	differential diagnosis with						
Demonstrate the		of knowledge related to	cytological findings.						
knowledge about FNAC	Also exhibit the knowledge	diagnosis of Bronchoalveolar	-Demonstrate the familiarity						
procedure and stains used	of fluid cytology like Pleural	lavage, Urine cytology, etc.	of knowledge related to						
in Cytopathology.	fluid, CSF, etc.		procedure and diagnosis of						
			cell block technique.						

MK 4: Knowledge required for Systemic Pathology

	Knowledge required for Systemic Pathology							
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5				
Demonstrate the	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone				
knowledge of normal	1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 exhibit the	level 4 Demonstrate the				
anatomy, embryology,	about aetiopathogenesis,	about aetiopathogenesis,	knowledge about	knowledge regarding				
histology, physiology of	gross and microscopic	gross and microscopic	aetiopathogenesis,	recent advances in				
various systems.	alterations of common	alterations of various tumors	morphology of rare diseases	aetiopathogenesis and				
	diseases in various system	in different system.	and complex syndromes.	morphology of various				
	like Diabetes Mellitus,			organs.				
	Pneumonia, Rhematic heart							
	disease, etc.							

MK 5: Knowledge specific to Surgical Pathology

Knowledge specific to Surgical Pathology							
Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5				
In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone				
1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate an in-	level 4 Demonstrate the				
about grossing and	about grossing and	depth knowledge about	knowledge regarding				
microscopy of small	microscopy of specimens	grossing and microscopy of	recent advances in				
specimen like	like Hysterectomy,	bigger specimens like	surgical pathology.				
appendicectomy,	Parotidectomy,etc	mastectomy, colectomy,					
cholecystectomy,etc		gastrectomy, etc without					
	In addition to milestone level	assistance.					
Also demonstrate the	3 Demonstrate an in-depth						
knowledge about various	knowledge about	Also demonstrate the					
special stains.	grossing and microscopy of	knowledge about the					
	bigger specimens like	various					
	mastectomy, colectomy,	immunohistochemical					
	gastrectomy,etc with	markers					
	assistance.						
	Milestone Level 2In addition to milestone level1 Demonstrate the knowledgeabout grossing andmicroscopy of smallspecimen likeappendicectomy,cholecystectomy,etcAlso demonstrate theknowledge about various	Milestone Level 2Milestone Level 3In addition to milestone levelIn addition to milestone level1 Demonstrate the knowledge2 demonstrate the knowledgeabout grossing andabout grossing andmicroscopy of smallmicroscopy of specimensspecimen likelike Hysterectomy,appendicectomy,Parotidectomy,etccholecystectomy,etcIn addition to milestone levelAlso demonstrate the3 Demonstrate an in-depthknowledge about variousgrossing and microscopy ofspecial stains.grossing and microscopy ofbigger specimens likemastectomy, colectomy,gastrectomy, etc withhile	Milestone Level 2Milestone Level 3Milestone Level 4In addition to milestone levelIn addition to milestone levelIn addition to milestone1 Demonstrate the knowledge about grossing and microscopy of small2 demonstrate the knowledge about grossing and microscopy of specimenslevel 3 Demonstrate an in- depth knowledge about grossing and microscopy of bigger specimens like mastectomy, ecloctomy, cholecystectomy,etcgrossing and microscopy of specimens bigger specimens like mastectomy, colectomy, gastrectomy,etc without assistance.Also demonstrate the knowledge about various special stains.In addition to milestone level 3 Demonstrate an in-depth knowledge about grossing and microscopy of bigger specimens like mastectomy, colectomy, gastrectomy, colectomy, immunohistochemical markers				

MK 6: Knowledge required for Autopsy Pathology

	Knowledge required for Autopsy Pathology							
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5				
Demonstrate knowledge	In addition to milestone level	In addition to milestone level 2	In addition to	In addition to				
about basic autopsy	1Demonstrate the knowledge	demonstrate the knowledge about	milestone level 3	milestone level 4				
techniques.	of different dissection	performing fetal autopsy with	Demonstrate an in-	Demonstrate the				
	methods of various organs	assistance.	depth knowledge	knowledge regarding				
Also possess the	and to give representative		regarding microscopic	recent advances in				
knowledge about the	sections from the organs.	Also exhibit the knowledge about	examination of various	autopsy pathology.				
prerequisites for autopsy.		identification of various pathological	disease.					
	Also possess knowledge	changes during autopsy.						
	about collection of samples		Also demonstrate					
	in different cases for		the knowledge about					
	biochemical and		performing fetal					
	microbiological		autopsy without					
	investigation.		assistance.					
	Also exhibit the knowledge							
	about identification of							
	various pathological changes							
	during autopsy.							

MK 7: Knowledge required for Hematology

	Knowledge required for Hematology							
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5				
Demonstrate a	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone				
knowledge of basic	1Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate an in-	level 4 Demonstrate the				
haematological	common haematological	about haematological	depth knowledge regarding	knowledge regarding				
techniques.	disorders like Iron deficiency	neoplasm like leukemia,	rare haematological disease	recent advances in				
	anemia, thalassemia, etc	lymphoma, etc.	like Congenital	haematological disorders				
Demonstrate the			dyserythropoeitic anemia,					
knowledge about	Able to demonstrate the		histiocytic disorders, etc.					
hematopoiesis and	knowledge about various	Possess the knowledge about						
morphology of each	haematological techniques.	special haematological tests,	Should possess knowledge					
hematopoietic cells.		cytochemical stains and basic	regarding approach to					
		flowcytometry.	Anemia, Leukemia,					
			Bleeding disorders, etc.					
				·				

	Knowledge specific to Transfusion Medicine (Blood Banking)					
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5		
Demonstrate the	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone		
knowledge about basic	1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate an in-	level 4 Demonstrate the		
Immunology and Blood	about clinical significance of	about rationale of pre-	depth knowledge Blood	knowledge regarding		
grouping system.	blood groups.	transfusion testing.	component therapy.	recent advances in		
			In addition to milestone	Transfusion medicine.		
			level 3 Demonstrate an in-	Also demonstrate the		
			depth knowledge about	knowledge about Quality		
			blood transfusion reaction	control in blood bank.		
			and its management.			
		26				

MK 8: Knowledge specific to Transfusion Medicine (Blood Banking)

MK 9: Knowledge required for Principles and practice of Laboratory Medicine

Knowledge required for Principles and practice of Laboratory Medicine					
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5	
Demonstrate knowledge	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone	
of the normal range of	1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate an in-	level 4 Demonstrate the	
values of the biochemical	about basic principles like	about	depth knowledge the	knowledge the principles	
content of body fluids,	Renal function test, Liver	the principles, advantages and	principles and methodology	good clinical laboratory	
significance of the	function test, Pancreatic	disadvantages, scope and	of quality control in the	practices (GCLP).	
altered values and its	function test,etc.	limitation of automation in	laboratory.		
interpretation.		the laboratory.			
		27			

Knowledge specific to Recent Advances and applied aspects.				
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate knowledge	In addition to milestone level	In addition to milestone level	In addition to milestone	In addition to milestone
of importance of	1 Demonstrate the knowledge	2 demonstrate the knowledge	level 3 Demonstrate an in-	level 4 Demonstrate the
statistical methods	about the principles and	about Recognise the	depth knowledge regarding	knowledge regarding
inassessing data from	techniques of	appearance of the normal	recent advances in	recent advances
patient material and	electronmicroscopy and the	subcellular organelles and	hematology.	Histopathology and
experimental studies.	working of an electron	their common abnormalities		Cytopathology.
	microscope	(when provided with		
Demonstrate familiarity	(includingTransmission and	appropriate photographs).		
with Biomedical Waste	Scanning Electron			
management disposal as	microscope: TEM and SEM).			
per Biomedical Waste				
Management				
amendment.				
	1	1	1	1

MK 10: Knowledge specific to Recent Advances and applied aspects.

MK11: Knowledge required for presenting seminar and journal club

Knowledge required for presenting seminar and journal club				
Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5	
Knowledge of analysing	Understands how to critically	Understands the direction of	Updates the knowledge	
Journals, articles,	analyse and compare articles	growth of the various Sub-	in recent advances in	
methodology and statistics	relevant to topic/practise	speciality in the field of	various branches of	
Knowledge of gathering	Able to form concepts on the	Pathology.	Pathology.	
relevant information from	subject and critically evaluate			
various sources and cites the	the limitation of the article.			
references.				
	Milestone Level 2Knowledge of analysingJournals, articles,methodology and statisticsKnowledge of gatheringrelevant information fromvarious sources and cites the	Milestone Level 2Milestone Level 3Knowledge of analysing Journals, articles, methodology and statisticsUnderstands how to critically analyse and compare articles relevant to topic/practiseKnowledge of gathering relevant information from various sources and cites theAble to form concepts on the subject and critically evaluate the limitation of the article.	Milestone Level 2Milestone Level 3Milestone Level 4Knowledge of analysing Journals, articles, methodology and statisticsUnderstands how to critically analyse and compare articles relevant to topic/practiseUnderstands the direction of growth of the various Sub- speciality in the field of Pathology.Knowledge of gathering relevant information from various sources and cites theAble to form concepts on the subject and critically evaluate the limitation of the article.Image: Compare articles growth of the various Sub- speciality in the field of Pathology.	

6.3.2 Patient Care/Procedural Skill – PC/PS

PC/PS.1. Perform gross examination of surgical pathology specimens

	Perform gross exa	amination of surgical pa	thology specimens	
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate a knowledge	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone
of normal anatomy,	Demonstrate the ability to	level 2 Identifies the gross	level 3 Ability to give	level 4 Demonstrate the
histology and functions of	identify the normal tissue	morphological changes in	various differential	knowledge regarding
various organs	and diseased tissue during	organs.	diagnoses by identifying	grossing rare diseases and
	grossing.		the different gross	Large cancer specimens
Demonstrate the ability to		Understand the grossing	morphology.	
attained proper sample		protocol for various		
collection and sample		organs.		
receiving				

	Perform grossing in grossing station.				
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5	
	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone	
Demonstrate the	Demonstrate the ability to	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the	
ability to attained	assists in grossing the seniors	ability to perform grossing	ability to perform grossing	ability to perform grossing	
proper sample	and colleagues.	of small specimen like	of large specimens	of large specimens	
identification,		Appendicectomy,	Mastectomy, Gastrectomy	Mastectomy, Gastrectomy	
labelling and sample	Ability to enter and correct the	Cholecystectomy, etc.	etc with assistance.	etc without assistance.	
receiving	final report in Hospital information system.		In addition to milestone		
		In addition to milestone	level 3 Demonstrate the		
	In addition to milestone level 2	level 2 demonstrate the	ability to perform grossing		
	demonstrate the ability to ink	ability to differentiate	of tumor area, lymph node		
	the specimen and fixation of	between the neoplastic and	dissection and margins		
	large specimens.	non neoplastic lesions	status.		

PC/PS.2. **Perform grossing in grossing station.**

	Interpretat	ion of Microscopic findi	ngs in tissue.	
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the ability to	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone
identify the normal	Demonstrate the ability to	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the
histology of various	identify the basic pattern and	ability to perform	ability to perform	ability to perform
tissues and organs.	reporting protocol in	histopathology reporting	histopathology reporting	histopathology reporting
	Histopathology.	of small specimens like	of large specimens like	of large specimens like
		Appendicectomy,	Mastectomy, Gastrectomy	Mastectomy, Gastrectomy
	Able to approve the report in	Cholecystectomy, etc.	etc with assistance.	etc without assistance.
	hospital information system			
	with assistance.	Able to write the	Able to write the	Able to give the final
		miscroscopic description	microscopic description	microscopic impression of
		and impression of non	for tumors and comment	tumor specimens with
		neoplastic specimens.	about the margin status	staging.
			and lymph node	
			metastasis.	Demonstrate the
				knowledge in recent
				advances.
L			1	1

PC/PS.3. Interpretation of Microscopic findings in tissue.

Perform histotechniques in Histopathology laboratory					
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5	
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone	
knowledge about basics in	Demonstrate the ability to	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the	
histotechniques and	observe the various	ability to tissue processing	ability to perform	ability to perform	
staining.	techniques in tissue	and staining.	microtomy without	special stains without	
	processing, section cutting		assistance.	assistance.	
	and staining.	Demonstrate the ability to			
		perform	Demonstrate the ability to	Possess the knowledge	
		Microtomy with	perform special stains	about the Internal and	
		assistance.	with assistance.	External quality control in	
				Histopathology laboratory.	
		33			

PC/PS.4. **Perform histotechniques in Histopathology laboratory**

PC/PS.5. Perform And Interpretation of FNAC and other Cytological smears (Pap smear, Sputum, Bronchial washings, Serous effusions, etc.

Perform And Interpre	etation of FNAC and other Cy	vtological smears (Pap smea	r, Sputum, Bronchial washi	ings, Serous effusions, etc.
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone
knowledge about FNAC	Demonstrate the ability to	level 2 Demonstrate the	level 3	level 4 Demonstrate the
procedure	observe the various	ability to sample	Demonstrate the ability to	ability to perform
Demonstrate the	techniques in cytology and	processing and staining in	describe the microscopic	Special stains and
knowledge about Sample	to perform	cytology.	findings of neoplastic	additional test in cytology.
collection, Identification,	FNAC procedure with	Demonstrate the ability to	lesion in FNAC, NGC and	
Receiving.	assistance.	perform FNAC procedure	cell block techniques	Demonstrate the ability to
		without assistance.		perform reporting various
To know the basics in			Ability to identify LGSIL,	cytological smears without
normal cytology reporting.	Able to approve the report in	Able to describe the	HGSIL and Invasive	assistance.
	hospital information system.	microscopic findings of	carcinoma in Pap smear.	
		non neoplastic lesion in		Demonstrate the
		FNAC and NGC		knowledge about recent
				advances in Cytology.
		Ability to identify the		
		reactive and inflammatory		
		lesions in Pap smear.		

PC/PS.6. Interpretation of Frozen sections

	Into	unnotation of Englan soatio	na	
	Inte	rpretation of Frozen sectio	ns	
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to milestone 1	In addition to milestone level	In addition to milestone	In addition to
knowledge about sample	Demonstrate the ability to	2 Demonstrate the ability to	level 3 Demonstrate the	milestone level 4
collection, receiving for	observe the various	do quick stain for frozen	ability to perform reporting	Demonstrate the
Frozen section	techniques in cytology.	sections.	various frozen section	knowledge about
			without assistance.	recent advances in
Demonstrate the	To know the basics in	Demonstrate the ability to		frozen section.
knowledge about	normal frozen section	perform reporting various	Ability to differentiate	
functioning of frozen	histology.	frozen section with assistance.	between benign and	
section microtome.			malignant tumors in frozen	
		Ability to identify the lymph	section.	
		node metastasis and margins		
		status in tumor cases.		
			1	I

Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to milestone 1	In addition to milestone level	In addition to milestone level	In addition to
knowledge about Basic	Demonstrate the ability to	2 Demonstrate the knowledge	3 Demonstrate the ability to	milestone level 4
requirement for IHC and	observe the various	about IHC markers panel for	perform IHC reporting using	Demonstrate the
IHC procedure	procedures in IHC.	various tumors to differentiate	various markers of each	ability to perform
		between epithelial or	system.(Lymphoma panel,	reporting various
Demonstrate the	To know the basics in	mesenchymal origin.	Small blue round cell tumor	IHC smears without
knowledge about Sample	normal IHC pattern.		Panel, etc)	assistance.
collection, Identification,		Demonstrate the ability to		
Receiving.	Demonstrate the ability to	perform IHC reporting using	Demonstrate the ability to	Demonstrate the
	perform	CK 7 and CK 20 to identify	perform IHC procedure	knowledge about
	IHC procedure with	the native of origin of	without assistance.	recent advances in
	assistance.	metastatic tumors.		IHC.
			1	1

PC/PS.7. Selection, performance and interpretation of appropriate Immuno-histochemical (IHC) markers.

PC/PS.8. Interpretation of Bone Marrow Smears

	Interp	retation of Bone Marrow Sn	nears	
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone
knowledge about Basics in	Demonstrate the ability to	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the
Bone marrow Aspiration	observe the Bone marrow	knowledge about Bone	ability to perform	ability to perform bone
procedure	aspiration procedure.	marrow reporting for	reporting of chronic	marrow smears of rare
		various acute leukemia	leukemia, Non-Hodgkins	lesions.
Demonstrate the	Demonstrate the knowledge	and Myelodysplastic	lymphoma, plasma cell	
knowledge about Sample	about Bone marrow	syndrome, etc.	disorder and metastasis.	Demonstrate the
collection, Identification,	reporting for various anemia			knowledge about recent
Receiving.	like Nutritional anemia and	Demonstrate the ability to		advances in Bone marrow
	haemolytic anemia.	perform		pathology.
To know the basics in		Bone marrow aspiration		
normal Bone marrow		procedure and staining		
histology.		without assistance.		

PC/PS.9. Perform and Interpretation of routine haematological investigations such as haemoglobin, TLC, DLC,

ESR PCV, Blood indices and peripheral smear.

Perform and Inter	Perform and Interpretation of routine haematological investigations such as haemoglobin, TLC, DLC, ESR PCV, Blood indices and peripheral smear.					
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5		
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone		
knowledge about Sample	Demonstrate the ability to	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the		
collection, Identification,	observe the various routing	knowledge about	ability to report various	ability to perform internal		
Receiving.	haematological procedure.	peripheral smear findings	peripheral smear of	and external quality		
		for acute leukemia,	chronic leukemia, Non-	control in hematology		
Demonstrate the ability to	Demonstrate the ability to	Neonatal jaundice,	Hodgkins lymphoma,	laboratory		
perform phlebotomy.	perform	Haemolytic anemia, etc.	plasma cell disorder and			
	Peripheral smear and	Demonstrate the working	metastasis	Demonstrate the		
Demonstrate the ability to	staining.	principle of Automated		knowledge about recent		
report the emergency		hematology analyzer.	Demonstrate the ability to	advances in hematology.		
haematological	Demonstrate the ability to		Handle trouble shooting in			
investigation with	report the nutritional anemia,	Demonstrate the ability to	Automated hematology	Demonstrate the		
assistance during stay	causes for leukocytosis,etc	report the emergency	analysers.	knowledge about NABL,		
duties		haematological		NABH and Biomedical		
		investigation without		Waste Management.		
		assistance during stay				
		duties.				

PC/PS.10. Perform and Interpretation of special investigations like Reticulocyte count, Sickling test, Osmotic Fragility

Test, Haemoglobin Electrophoresis, Fetal Haemoglobin, etc.

Perform and Interpretation of special investigations like Reticulocyte count, Sickling test, Osmotic Fragility Test, Haemoglobin Electrophoresis, Fetal Haemoglobin, etc.

Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone level 4
Demonstrate the	Demonstrate the ability to	level 2 Demonstrate the	level 3 Demonstrate the	Demonstrate the ability to perform
knowledge about	observe the various	ability to report the	ability to report the	internal and external quality control
Sample collection,	special haematological	special haematological	special haematological	in hematology laboratory
Identification,	procedures in new born	investigation for	investigation for	
Receiving.	smear.	Hereditary spherocytosis,	Dyserthyropoeitic	Demonstrate the knowledge about
		Thalassemia, Sickle cell	anemia, Pancytopenia,	recent advances in hematology.
	Demonstrate the ability to	anemia, etc	etc.	
	perform			Demonstrate the knowledge about
	Knowledge about various			NABL, NABH and Biomedical
	Special haematological			Waste Management.
	test.			
	1		1	L

Planning investigations for a Clinical case				
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone
knowledge about Basic	Demonstrate the knowledge	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the
pathology of various	about investigation for	ability to correlate	ability to interact with	knowledge about recent
diseases.	various diseases.	Clinical, Radiological and	clinicians and plan a	advances in Investigations
		other findings	investigation for a	for various diseases.
			particular disease.	
		40		

PC/PS.11. Planning investigations for a Clinical case

Perform and Interpret Urine Examination, Body fluids and semen analysis					
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5	
	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone	
Demonstrate the	Demonstrate the ability to	level 2 Demonstrate the	level 3 Should posses the	level 4 Demonstrate the	
knowledge about Sample	observe the various clinical	ability to report the Urine	knowledge about	ability to perform internal	
collection, Identification,	pathological procedure and	microscopy like	automations in Urine	and external quality	
Receiving.	semen analysis	sediments, cast and	analysis, Computer	control in clinical	
		crystals.	Assisted Semen Analysis	pathology laboratory	
	Demonstrate the ability to		(CASA).		
	report the emergency fluid	Demonstrate the ability to		Demonstrate the	
	counts and urine ketone	report the	Demonstrate the ability to	knowledge about recent	
	investigation during stay	Reactive non-neoplastic	report the	advances in clinical	
	duties.	lesions in various Body	Malignant lesions like	pathology	
		fluids.	Adenocarcinoma and		
	Ability to perform test for		Squamous cell carcinoma	Demonstrate the	
	Bence zones proteins.	Demonstrate the ability to	in various Body fluids	knowledge about NABL,	
		report the		NABH and Biomedical	
		Oligospermia and		Waste Management.	
		azoospermia in semen			
		analysis.			
		1	1	1	

PC/PS.12. **Perform and Interpret Urine Examination, Body fluids and semen analysis**

PC/PS.13. Interpretation of ancillary techniques like Immunofluorescence, Karyotyping, FISH, PCR and Electron Microscopy.

Interpretation of	of ancillary techniques like Im	munofluorescence, Karyoty	ping, FISH, PCR and Elect	tron Microscopy.
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the Basic	In addition to milestone 1	In addition to milestone	. In addition to milestone	In addition to milestone
knowledge of various	Demonstrate the knowledge	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the
ancillary techniques.	of principle of various	knowledge about	knowledge about	knowledge about recent
	ancillary techniques.	procedures of various	interpretation of various	advances in various
Demonstrate the		ancillary techniques	ancillary techniques	ancillary techniques
knowledge about Sample				
collection, Identification,				
Receiving for various				
ancillary techniques				
		42		

Ι	Demonstration of familiarity wit	hin laboratory investigations	in Microbiology and Biocher	nistry
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the Basic	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone
knowledge of various	Demonstrate the knowledge	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the
Microbiology and	of principle of various	knowledge about	knowledge about	knowledge about recent
Biochemical investigation	Microbiology and	procedures of various	interpretation of various	advances in various
	Biochemical investigation	Microbiology and	Microbiology and	Microbiology and
Demonstrate the		Biochemical investigation	Biochemical investigation	Biochemical investigation
knowledge about Sample				
collection, Identification,				
Receiving for various				
ancillary techniques				

PC/PS.14. Demonstration of familiarity within laboratory investigations in Microbiology and Biochemistry

PS/PC 15 : Perform and interpret Blood banking techniques (Blood grouping, Rh typing, Cross-matching and Coomb's test, ELISA for infectious disease)

Perform and interpret Blood banking techniques (Blood grouping, Rh typing, Cross-matching and Coomb's test, ELISA for infectious disease)

Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to	In addition to milestone level	In addition to milestone	In addition to milestone level 4
knowledge about	milestone 1	2 Demonstrate the	level 3 Demonstrate the	Demonstrate the ability to
Sample collection,	Demonstrate the	knowledge about Principle,	ability to report	perform internal and external
Identification,	ability to observe the	Procedure and interpretation	Blood grouping using gel	quality control in Blood Bank.
Receiving.	various blood	of Blood grouping, Cross	card method, Cross	
	banking techniques.	matching and Coomb test.	matching and Coomb test.	Demonstrate the knowledge
Demonstrate the ability				about recent advances in Blood
to perform phlebotomy.		Demonstrate the Knowledge	Demonstrate the ability to	banking techniques.
		about interpretation of	Handle trouble shooting in	
		ELISA.	Cross matching and	Demonstrate the knowledge
			Coombs test (False positive	about FDA regulations and
			and False Negative cases).	Biomedical Waste Management.
	L4			

PC/PS. 16 : Selection of blood donors and Management of adverse donor reactions.

Milestone Level 1 Demonstrate the	Selection of blood done Milestone Level 2	C	erse donor reactions.						
	Milestone Level 2		Selection of blood donors and Management of adverse donor reactions.						
Demonstrate the		Milestone Level 3	Milestone Level 4	Milestone Level 5					
	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone					
knowledge about criteria	Observe and know the	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the					
for selection of blood	procedure and criteria for	knowledge about various	ability to recognise and	knowledge about recent					
donors.	Donor selection.	local and systemic adverse	management of Adverse	advances in Blood donors					
		blood donor reactions.	blood donor reaction.	and management of					
Demonstrate the	Ability to know the			Adverse donor reactions.					
knowledge about Adverse	procedure for Blood								
donor reaction.	collection from donors.								
		45							

PS /PC-17 : Perform investigation for a case of mismatched blood transfusion.

	Porform investigation	n for a case of mis	matched blood transfusi	0 n
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5
Demonstrate the	In addition to milestone 1	In addition to	In addition to milestone	In addition to milestone level 4
knowledge about routine	Demonstrate the knowledge	milestone level 2	level 3 Demonstrate the	Demonstrate the ability to perform
blood transfusion and its	about various causes for	Demonstrate the	ability to perform the	Internal and External quality
protocol.	mismatch blood transfusion.	ability to perform	Root Cause Analysis of	control in Blood Bank.
		the Root Cause	mismatch blood	
		Analysis of	transfusion without	Demonstrate the knowledge about
		mismatch blood	assistance.	recent advances in Mismatched
		transfusion with		blood transfusion.
		assistance.		
				Demonstrate the knowledge about
				FDA regulations and Biomedical
				Waste Management.
		46		

PC/PS 18 : Participation and Presentation in multidisciplinary meetings like tumor boards, CPCs, Dermato-Pathological conferences.

Participation an	Participation and Presentation in multidisciplinary meetings like tumor boards, CPCs, Dermato-Pathological conferences.					
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5		
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone		
knowledge about various	Demonstrate the knowledge	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the		
pathological diseases.	about reports, its Clinical	ability to perform clinic-	ability to do presentation	ability to help in guiding		
	findings and radiological	Radio-Pathological	in multidisciplinary	the clinician for further		
	investigations.	correlation.	meetings.	management to investigate		
				a particular case.		
		Demonstrate the ability to				
		participate in				
		multidisciplinary				
		meetings.				
		47				

PS/PC 19 : Teaching pathology to undergraduates (MBBS), and allied health sciences like BDS, BSc (Nursing), BSc (MLT), BSc (Radiology), etc.

Teaching pathology	v to undergraduates (MBBS), an	nd allied health sciences like etc.	BDS, BSc (Nursing), BSc	(MLT), BSc (Radiology)		
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5		
Demonstrate the	In addition to milestone 1	In addition to milestone	In addition to milestone	In addition to milestone		
knowledge about basic	Demonstrate the knowledge	level 2 Demonstrate the	level 3 Demonstrate the	level 4 Demonstrate the		
concepts in pathology.	to perform practical teaching	ability to perform theory	ability to perform theory	ability to teach the		
	for undergraduates.	teaching for BSC Nursing	teaching and small group	colleagues and Junior		
		and Allied Health	discussions for BDS and	Resident in the		
		Sciences.	MBBS.	Department.		
			Demonstrate the ability to			
			teach the Technical staffs			
			in the Department.			
		48				

6.3.3 System based practice

SBP 1. Laboratory safety and Systems Approach to Laboratory Errors.

Laboratory safety	y and Systems. Approach to L	aboratory Errors.						
Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5					
In addition to milestone	In addition to milestone	In addition to milestone	In addition to milestone					
level 1 Demonstrate	level 2 Participate in	level 3 Report errors and	level 2 Organize and					
knowledge of	laboratory safety reporting	near- misses to the	leads institutional					
institutional surveillance	and analyzing systems	institutional surveillance	QI/Laboratory safety					
systems to monitor for		system and superiors	projects.					
patient safety (e.g.,	Demonstrate knowledge	Participate in quality						
Laboratory error	national laboratory safety	improvement	Recognize when root					
reporting)	standards, as well as their	(QI)/Laboratory safety	cause analysis is					
	use/application in the	practices.	necessary, and is					
Demonstrate knowledge	institution		capable of participating					
of the epidemiology of			in root cause analysis					
laboratory errors and the								
differences between Pre-								
analytical, Analytical and								
Post-analytical errors.								
	Milestone Level 2 In addition to milestone level 1 Demonstrate knowledge of institutional surveillance systems to monitor for patient safety (e.g., Laboratory error reporting) Demonstrate knowledge of the epidemiology of laboratory errors and the differences between Pre- analytical, Analytical and	Milestone Level 2Milestone Level 3In addition to milestoneIn addition to milestonelevel 1 Demonstratelevel 2 Participate inknowledge oflaboratory safety reportinginstitutional surveillanceand analyzing systemssystems to monitor forDemonstrate knowledgepatient safety (e.g.,Demonstrate knowledgeLaboratory errornational laboratory safetyreporting)standards, as well as theiruse/application in theinstitutionof the epidemiology ofinstitutionlaboratory errors and thedifferences between Pre-analytical, Analytical andInstitution	Milestone Level 2Milestone Level 3Milestone Level 4In addition to milestoneIn addition to milestoneIn addition to milestonelevel 1 Demonstratelevel 2 Participate inlevel 3 Report errors andknowledge oflaboratory safety reportingnear- misses to theinstitutional surveillanceand analyzing systemsinstitutional surveillancesystems to monitor forDemonstrate knowledgeParticipate in qualitypatient safety (e.g.,Demonstrate knowledgeParticipate in qualitytaboratory errornational laboratory safetyimprovementreporting)standards, as well as their(QI)/Laboratory safetypatient safety (e.g.,institutionpractices.laboratory errornational laboratory safetyimprovementreporting)institutioninstitutionof the epidemiology ofinstitutioninstitutionlaboratory errors and theinstitutioninstitutiondifferences between Pre-analytical, Analytical andinstitution					

6.3.4 Practice based learning and improvement

PBLI 1: Self-directed Learning/Critical Appraisal of Medical Literature.

	Self-directed Learning/Critical Appraisal of Medical Literature.									
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5						
Demonstrate an	In addition to milestone	In addition to milestone	In addition to milestone	In addition to milestone						
understanding of	level 1 Identify resources	level 2 Critically review and	level 3 Tailor evidence-	level 4 Design a						
critical appraisal of the	(e.g., texts, search engines)	interprets the literature with	based practice based on	hypothesis-driven or						
literature	to answer questions while	the ability to identify study	the values and preferences	hypothesis-generating						
	providing patient care	aims, hypotheses, design,	of each cases	study Contribute to						
Demonstrate	Recognize limits of	and biases	Reads and assess strength	peer-reviewed medical						
responsiveness to	knowledge, expertise, and		of evidence in current	literature						
constructive feedback	technical skills.		literature and applies it to							
			one's own practice							
	Describe commonly used									
	study designs (e.g.,									
	randomized controlled trial									
	[RCT], cohort; case-									
	control, cross-sectional)									

6.3.5 Interpersonal communication skills

IPCS 1: Communicate the reports effectively with clinician, patients and families

	Communicate the reports effectively with clinician, patients and families									
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5						
Demonstrate the ability	In addition to milestone level	In addition to milestone	In addition to milestone	In addition to milestone						
of basic communication	1 Maintain communication	level 2 Communicate	level 3 Participate in	level 4 Capable of						
skills.	with clinicians and surgeons	effectively the reports to	various inter and	presentation in various						
	regarding the status of the	clinicians and patient	Intradepartmental	forums about the case						
	pathology report.	relatives	meetings like CPC, etc	report.						

IPCS 2: Communication with Colleagues and Technical staffs

Communication with Colleagues and Technical staffs									
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5					
Understand the	In addition to milestone level	In addition to milestone	In addition to milestone	In addition to milestone					
importance of	1 Demonstrate an	level 2 Communicate	level 3 Participate in	level 4 Role model for					
relationship	understanding of the roles of	effectively with	various Departmental	effective					
development,	health care team members,	colleagues within	meetings.	communication to					
information gathering	and communicates effectively	department and with		junior colleagues					
and sharing, and	within the team	technical staffs.							

teamwork		
	Able to communicate the	
	technical orders to the	
	technical staffs regarding	
	laboratory investigation	

IPCS 3: Teaching to Undergraduates students and other colleagues

Teaching to Undergraduates students and other collaegues									
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5					
Understand the	In addition to milestone level	In addition to milestone	In addition to milestone	In addition to milestone					
importance of concepts	1 Assist in teaching the	level 2 Participate in	level 3 Participate in	level 4 Model and					
in pathology and able to	Undergraduate students.	teaching undergraduates	multidisciplinary	coach shared decision					
communicates to		both theory and practical.	family/patient/team	making in complex and					
juniors.	Ť		member conferences for	highly stressful					
			informed consent and	situations					
			shared decision making.						
			•	·					

6.3.6 Professionalism

P.1- Compassion, Integrity, and Respect for Others.

Compassion, Integrity, and Respect for Others									
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5					
Understand the	In addition to milestone	In addition to milestone	In addition to milestone	In addition to milestone					
importance of	level 1Consistently show	level 2 Modify one's own	level 3 Consistently	level 4 Assume long-term					
compassion, integrity,	compassion, integrity, and	behaviour based on	model compassion,	or leadership role in					
and respect for others	respect in typical	feedback to improve his or	integrity, and respect for	community outreach					
Demonstrate sensitivity	situations with patients,	her ability to demonstrate	others.	activities to improve the					
and responsiveness to	peers, and members of the	compassion, integrity, and		health of vulnerable					
patients	health care team Consistently demonstrate sensitivity and responsiveness to the diagnostic services.	respect for others	Coach others to improve compassion, integrity, and respect to diagnostic services	populations					

P.2- Feedback.

Feedback.								
Milestone Level 1	Milestone Level 2	Milestone Level 3	Milestone Level 4	Milestone Level 5				
Seeks constructive	Accepts feedback from	Correlates feedback with	Provides constructive	Effectively seeks and				
feedback from faculty	faculty members and	self-reflection and	feedback to juniors in a	provides constructive				
members and	incorporates suggestions	incorporates it into	tactful and supportive	feedback in challenging				
colleagues.	into practice.	lifelong learning to	way to enhance patient	situations.				
		enhance patient care.	care.					

7 Syllabus

7.1 Course 1 General Pathology, Pathophysiology, Immunopathology and Cytopathology

7.1.1 General Pathology and pathophysiology

- Normal cell and tissue structure and function.
- The changes in cellular structure and function in disease.
- Causes of disease and its pathogenesis.
- Reaction of cells, tissues, organ systems and the body as a whole to various sublethal and lethal injuries.
- Basic Course in Biomedical Research, Data collection and analysis, Scientific communication.

7.1.2 Immunopathology

- Demonstrate familiarity with the current concepts of structure and function of the immune system, its aberrations and mechanisms thereof.
- Demonstrate familiarity with the scope, principles, limitations and

interpretations of the results of the following procedures employed in clinical and experimental studies relating to immunology.

- (a) ELISA techniques
- (b) Radioimmunoassay
- (c) HLA typing
- Interpret simple immunological tests used in diagnosis of diseases and in research procedures.
 - (i) Immunoelectrophoresis
 - (ii) Immunofluorescence techniques especially on kidney and skin

biopsies

- (iii) Anti-nuclear antibody (ANA)
- (iv) Anti-neutrophil cytoplasmic antibody (ANCA)

7.1.3 Cytopathology

• Should possess the background necessary for the evaluation and reporting of cytopathology specimens.

• Demonstrate familiarity with the following, keeping in mind the indication for the test.

(i) Choice of site from which smears may be taken

(ii) Type of samples

(iii) Method of obtaining various specimens (urine sample, gastric

smear, colonic lavage etc.)

(iv) Be conversant with the principles and preparation of solutions of

stains

7.2 Course 2: Systemic Pathology:

7.2.1 Systemic Pathology

- Normal structure and function of various organ systems.
- Aetiopathogenesis, gross and microscopic alterations of structure of these organ systems in disease and functional correlation with clinical features.

7.2.2 Surgical Pathology

- Histogenetic and patho-physiologic processes associated with various lesions.
- Problems in the laboratory and offer viable solutions.

7.2.3 Autopsy Pathology

- Technique of autopsy.
- Various disease processes so that a meaningful clinico-pathological correlation can be made.



7.3 Course 3:Haematology, Transfusion Medicine (Blood Banking) and Laboratory Medicine

7.3.1 Hematology

- Should demonstrate the capability of utilising the principles of the practice of Haematology for the planning of tests, interpretation and diagnosis of diseases of the blood and bone marrow.
- Should be conversant with various equipment used in the Haematologylaboratory.
- Should have knowledge of automation and quality assurance in Haematology.
- Correctly plan a strategy of investigating at least 90% of the cases referred forspecial investigations in the Hematology Clinic and give ample justification for each step in consideration of the relevant clinical data provided.
- The student is expected to acquire a general acquaintance of techniques and principles and to interpret data in the following fields.
 - a) Immunopathology
 - b) Electron microscopy
 - c) Histochemistry
 - d) Immunohistochemistry
 - e) Cytogenetics
 - f) Molecular Biology
 - g) Maintenance of records

h) Information retrieval, use of Computer and Internet in medicine.

i) Quality control, waste disposal

7.3.2 Transfusion Medicine (Blood Banking)

- Basic immunology.
- ABO and Rh groups.
- Clinical significance of other blood groups.
- Transfusion therapy including the use of whole blood and RBC concentrates.
- Blood component therapy.
- Rationale of pre-transfusion testing.

- Infections transmitted in blood.
- Adverse reactions to transfusion of blood and components.
- Quality control in blood bank.

7.3.3 Laboratory Medicine

- Possess knowledge of the normal range of values of the biochemical content of body fluids, significance of the altered values and its interpretation.
- Possess knowledge of the principles of following specialized organ function tests and the relative utility and limitations of each and significance of the altered values.
 - (i) Renal function tests
 - (ii) Liver function tests
 - (iii) Pancreatic function tests
 - (iv) Endocrine function tests
 - (v) Tests for malabsorption
- Know the principles, advantages and disadvantages, scope and limitation of automation in the laboratory.
- Know the principles and methodology of quality control in the laboratory.
- Know the principles good clinical laboratory practices(GCLP)

7.4 Course 4: Recent Advances and Applied aspects

7.4.1 Recent advances:

- Recent advances in Histopathology and Cytology
- Recent advances in Hematology and Transfusion medicines.

7.4.2 Applied aspects:

- Demonstrate familiarity with the principles and techniques of electronmicroscopy and the working of an electron microscope (includingTransmission and Scanning Electron microscope: TEM and SEM).
- Recognise the appearance of the normal subcellular organelles and their common abnormalities (when provided with appropriate photographs).
- Should be familiar with the principles, use and interpretation of common enzyme histochemical procedures (Alkaline Phosphatase, Acid Phosphatase, Glucose-6-Phosphate Dehydrogenase, Chloroacetate Esterase).
- Demonstrate familiarity with the principles and exact procedures of various immunohistochemical stains using both PAP (Peroxidase-antiperoxidase)and AP-AAP (Alk. Phosphatase-anti-Alk. Phosphatase) ABC(Avidin-Biotin Conjugate) systems; employing monoclonal and polyclonalantibodies.
- Be aware of the limitations of immuno-histochemistry.
- Should understand the principles of molecular biology especially related to the understanding of disease processes and its use in various diagnostic tests.
 Should be conversant with the principle and steps and interpretation of Polymerase Chain Reaction (PCR), Western Blot, Southern Blot, Northern Blot and Hybridisation) procedures.
- Demonstrate familiarity with methods of Karyotyping and Fluorescent in-situ Hybridisation (FISH).
- Demonstrate familiarity with methods of tissue culture.
- Demonstrate familiarity with importance of statistical methods in assessing data from patient material and experimental studies.
- Demonstrate familiarity with Biomedical Waste management disposal as per Biomedical Waste Management amendment. (2016 guidelines)

8 Teaching and Learning Method

The trainee will undergo a graded training over a period of three years.

o <u>Orientation</u>

At the beginning of the course each resident should be given an orientation to the department and subject. The candidate shall be assigned dissertation guides so as to help them prepare protocols

8.1 Theory (Knowledge/ Cognitive Domain)

The teaching learning methods does not totally depend on didactic lectures. Only the introductory lectures by faculty are in this format.

8.1.1 Introductory lectures

8.1.2 Teaching programme

- Lectures, seminars, symposia, Inter- and intra- departmental meetings (clinicpathological, Tumor board, Derm path, OG- Path, Ortho-Path meet), maternal morbidity/mortality meetings and journal club. Records of these are to be maintained by the department.
- By encouraging and allowing the students to attend and actively participate in CMEs, Conferences by presenting papers.

• Maintenance of log book:E-portfolio:- It is an electronic portfolio to be maintained by the resident to record their activities under the section:

- EPA,
- Daily log
- Diagnostic work
- Procedure
- Dissertation
- Academic activities(Seminar, symposium, case presentation, journal club)
- Co-curricular activities (Conference, CME, Workshop),
- Teaching Assignments,
- Awards and achievements
- Outreach activities.

- **E-portfolio** shall be checked and assessed periodically by the faculty members. This will enable to monitor progress of the resident, his level of attainment of milestone and impart the training accordingly
- Writing thesis following appropriate research methodology, ethical clearance and good clinical practice guidelines.
- The postgraduate students shall be required to participate in the teaching and training programme of undergraduate students and interns.
- A postgraduate student of a postgraduate degree course in broad specialities/super specialities would be required to present one poster presentation, to read one paper at a national/state conference and to present one research paper which should be published/accepted for publication/sent for publication during the period of his postgraduate studies so as to make him eligible to appear at the postgraduate degree examination.
- Department should encourage e-learning activities.

8.1.3 Structured Graded Training-Year wise Knowledge / cognitive

domain

8.2 Practical skills training (psychomotor domain)

8.2.1 Resident Rotations

	1^{st}	2^{nd}	3 rd	4^{th}	5^{th}	6^{th}	7^{th}	8^{th}	9 th	10^{th}	11 th	12 th
	Mo	Mo	Mo	Mo	Mo	Mo	Mo	Mo	Mo	Mo	Mo	Mo
	n	n	n	n	n	n	n	n	n	n	n	n
1 st yea r	TE C	HM	HM	HM	С	С	С	Н	Н	Н	BB	BB
2 nd yea r	HM	HM	HM	С	C	C	Н	Н	Н	Н	MT	AP*
3 rd yea r	НМ	НМ	НМ	С	C	Н	Н	Н	MP	MP	Н	Н

• Details of 3 years posting in the PG programme (6 terms of 6 months each)

TEC– Lab techniques, HM- Hematology, C-Cytopathology,H- Histopathology, BB-Blood banking, MT- Museum Technique, MP- Molecular Pathology, AP-Allied post *Allied posts should be done during the course – for 4 weeks

• Biochemistry- 2 weeks

• Microbiology - 2 weeks

Section/Subject

months

Duration in

•	(i) Surgical Pathology and Autopsy and Pathology Techniques	12
•	(ii) Haematology and Laboratory Medicine	10
•	(iii) Cytopathology	08
•	(iv) Transfusion Medicine/Blood Bank	02
•	(v) Museum techniques and record management	01
•	(vi) Basic Sciences including Immunopathology,	
•	Electron microscopy, Molecular Biology,	
•	Research Techniques and cytogenetics etc	02

Total 35

Details of training in the subject during resident posting. The student should attend to the duties (Routine and emergency).

8.2.2 Structured Graded Training –Year - wise Practical training objectives

8.2.3 Practical Training

- Collection of specimens including Fine Needle Aspiration of lumps.
- Grossing of specimens.
- Performing autopsies.
- Processing and Block making
- H&E staining
- PAP smear staining
- Peripheral smear staining.
- Bone marrow aspiration.
- Discussion during routine activities such as during signing out of cases.
- Presentation and work-up of cases including the identification of special stains and ancillary procedures needed.
- Research Presentation and review of research work.
- Laboratory work of haematological test.
- Selection and bleeding of donors
- Blood grouping and cross matching
- Coombs test
- Use and maintenance of equipment.
- Maintenance of records.

E - portfolio

It is an electronic portfolio to be maintained by the resident to record their day to day academic and patient care activities under the following sections:

- Entrustable Professional Activity assessment
- Daily log
- Patient care
- Procedure

- Dissertation
- Academic activities(Seminar, symposium, case presentation, journal club)
- Co curricular activities (Conference, CME, Workshop),
- Teaching Assignments,
- Awards and achievements
- Outreach activities.

E - portfolio will be monitored and endorsed periodically by the faculty supervisors. This will enable faculty to monitor residents progress, attainment of milestones and impart the training accordingly.

9 Assessment

Assessment will have 2 components Formative and Summative

9.1 Formative assessment

9.1.1 Cognitive Assessment

- Assessment in Cognitive Domain
- Schedule of theory tests
 - 1st year 2 papers consisting of syllabus from Course 1
 - \circ 2nd year 2 papers consisting of syllabus from Course 2 and 3
 - \circ 3rd year one paper consisting of syllabus from Course 4
 - 3rd year Mock exams one month prior to University examination,

consisting of 4 papers, including syllabus from all the four courses.

9.1.2 EPA Assessment

Assessment of Entrustable Professional Activities (EPA) done during the OT posting by the consultant in - charge. EPA assessment will be done once by the end of the 1st week of the posting and then again at the end of the posting, for monitoring of resident progress.

EPA	
No.	Entrustable Professional Activity
	Perform gross examination of surgical pathology specimens and Interpretation of
1.	Microscopic findings in tissue.
2	Perform histotechniques (Tissue processing, embedding, microtomy, Staining and
2	special staining)
3.	Perform And Interpretation of FNAC and other Cytological smears (Sputum, Bronchial
5.	washings, Serous effusions, etc.
4.	Interpretation of Frozen sections
5.	Selection, performance and interpretation of appropriate Immunohistochemical markers.
6.	Interpretation of Bone Marrow Smears
7.	Perform and Interpretation of routine haematological investigations like haemoglobin,
7.	TLC, DLC, ESR PCV, Blood indices and peripheral smear.
8.	Perform and Interpretation of special investigations like Reticulocyte count, Sickling
0.	test, Osmotic Fragility Test, Haemoglobin Electrophoresis, Fetal Haemoglobin, etc.
9.	Planning investigations for a Clinical case
10.	Perform and Interpret Urine Examination, Body fluids and semen analysis.
11	Interpretation of ancillary techniques like Immunofluorescence, Karyotyping, FISH,
	PCR and Electron Microscopy.
12	Demonstration of familiarity within laboratory investigations in Microbiology and
	Biochemistry
13	Perform and interpret Blood banking techniques (Blood grouping, Rh typing, Cross-
	matching and Coomb's test, ELISA for infectious disease like HIV, HBsAg).
14	Selection of blood donors and Management of adverse donor reactions. Perform
	investigation for a case of mismatched blood transfusion.
15	Participation and Presentation in multidisciplinary meetings like tumor boards, CPCs,
15	Dermato-Pathological conferences. Teaching pathology to undergraduates (MBBS), and allied health aciences like RDS. RSe (Nursing) RSe (MLT) RSe (Bediology) ato
	allied health sciences like BDS, BSc (Nursing), BSc (MLT), BSc (Radiology), etc.

Table 3: List the of Entrustable Professional Activity

EPA Descriptions (Enter all the EPA and their descriptions)

9.1.3 Table : 4. EPA description

9.1.4 EPA 1: Perform gross examination of surgical pathology specimens and Interpretation of Microscopic findings in tissue.

EF	EPA 1: Perform gross examination of surgical pathology specimens and Interpretation of						
Mi	croscopic findings in tissue.						
1.	Description of the activity:	Residents should be	able to identify v	arious path	ological		
		changes grossly and	take section for f	further mici	roscopic		
		examination. Able to interpret the microscopic findings of					
		various tissues.					
2.	Most relevant domains of competence:	MK, PS, ICS, P					
3.	Sub-Competencies within each	ch MK1,4,5,6					
	domain:	PS 1,2,3					
		ICS1&2					
		P1					
		SBP 1					
		PBLI 1					
4.	Methods of assessment	1. MK and PS with a la					
			gy postings, Writ n skills and profe				
			ltisource feedbac		a 4		
		5. Assessment do	ne during the end	i posting te	st.		
5.	Resident will be entrustable when this subcompetency	МК	1 L4	4 L3	5&6 L4		
	milestones levels are attained.	PS	1 L4	2 L3	3 L4		
		ICS	1 L3	2 L2			
		Р	1 L3				
		SBP	1 L3				
		PBLI	1 L3				

EPA 2: Perform histotechniques (Tissue processing, embedding, microtomy, Staining and special stainning)

EPA :					
1.Description of the	Residents should be able to perform various				
activity:	histotechniques (7	histotechniques (Tissue processing, embedding,			
	microtomy, Staini	microtomy, Staining and special staining). Also to			
	address any techn	ical issues in the	histopathology lab		
2. Most relevant domains of competence:	MK, PS, ICS, P	MK, PS, ICS, P			
3. Sub-Competencies	MK.5	MK.5			
within each domain:	PS 4				
	ICS 2				
	P1				
	SBP 1				
	PBLI 1				
4. Methods of assessment	4. MK and PS will be assessed by faculty during grossing in histopathology postings, written				
	exam & epor	tings, written			
	5. Communication skills and professionalism will				
	be assessed by multisource feedback.6. Assessment done during the end posting test.				
	0. 14556551101110	ione during the e	nd posting tost.		
5. Resident will be	МК	5 L3			
entrustable when this subcompetency	PS	4 L4			
milestones levels are	ICS	2 L2			
attained.	Р	1 L3			
	SBP	1 L2			
	PBLI	1 L3			

EPA 3: Perform And Interpretation of FNAC and other Cytological
smears (Sputum, Bronchial washings, Serous effusions, etc.)

EPA :				
1. Description of the	Residents should be able to perform And			
activity:	Interpret FNAC and other Cytological smears			
	(Sputum, Bronchial washings, Serous effusions			
	Pap smear, etc.)			
2. Most relevant				
domains of	MK, PS, ICS, P			
competence:				
3. Sub-Competencies	MK3			
within each domain:	PS 5			
	ICS1&2			
	P1			
	SBP 1			
	PBLI 1			
4. Methods of	MK and PS will be assessed by faculty			
assessment	during grossing in histopathology postings,			
	Written exam & eportfolio.			
	Communication skills and professionalism will be assessed by multisource feedback.			
	Assessment done during the end posting			
	test.			
5. Resident will be	MK 3 L4			
entrustable when this	PS 5 L4			
subcompetency	ICS 1 L2 2 L2			
milestones levels are	P 1 L3			
attained.				
	SBP 1 L3			
	PBLI 1 L3			

EPA 4: Interpretation of Frozen sections

1.Description of the activity:	Residents should know the process and				
	interpretation of f	frozen section.			
2. Most relevant domains of competence:	MK, PS, ICS, P	MK, PS, ICS, P			
3. Sub-Competencies within	MK 5				
each domain:	PS 1&6				
	ICS1				
	P1				
	SBP 1				
	PBLI 1				
4. Methods of assessment	MK and PS will be assessed by faculty				
	during grossing in histopathology				
	postings, Written exam & eportfolio.				
	Communication skills and professionalism				
	will be assessed by multisource feedback				
		lone during the			
	test.		r - St	D	
5. Resident will be	МК	5 L3			
entrustable when this subcompetency milestones	PS	1 L 3	6 L3		
levels are attained.	ICS	1 L2			
	Р	1 L1			
	SBP	1 L2			
	PBLI	1 L1			

EPA 5: Selection, performance and interpretation of appropriate

Immunohistochemical markers.

EPA :				
1.Description of the	Residents should know the techniques of			
activity:	imunohistochemistry. Residents should			ıld
	know the IHC panel for various tumors a			ors and
	its interpretation.			
2 Most relevant				
domains of	MK, PS, ICS, P			
competence:				
3. Sub-Competencies	MK 5			
within each domain:	PS 7			
	ICS 1&2			
	P1			
	SBP 1			
	PBLI 1			
4. Methods of	- MK and PS will be assessed by			у
assessment	faculty during grossing in			
	 histopathology postings, Written ex & eportfolio. Communication skills and professionalism will be assessed by multisource feedback. Assessment done during the end 			en exam
				ed by
				J
				nd
	posting te		[1
5. Resident will be	MK	5 L4		
entrustable when this	PS	7 L3		
subcompetency	ICS	1L2	2 L2	
milestones levels	Р	1L3		
are attained.	_			
	SBP	1L2		
	PBLI	1L3		

EPA 6: Interpretation of Bone Marrow Smears.

1.Description of the	Resident should be able to do interpretation of bone marrow smears of various anemia, Leukemia, etc.			
activity:				
2 Most relevant domains of competence:	MK, PS, ICS, P			
3 Sub-Competencies	MK 7			
within each domain:	PS 8			
	ICS1&2			
	P1			
	SBP 1			
	PBLI 1			
4 Methods of		will be assesse	•	• •
assessment	grossing in histopathology postings, V exam & eportfolio.			
	 Communication skills and professionalism wi 			
	be assessed by multisource feedback.			
	- Assessment	t done during th	ne end pos	ting test.
			1	1
	MK	7 L3		
5 Resident will be				
entrustable when	PS	8 L 4		
entrustable when this subcompetency milestones levels are		8 L 4 1 L2	2 L2	
entrustable when this subcompetency	PS		2 L2	
entrustable when this subcompetency milestones levels are	PS ICS	1 L2	2 L2	

EPA 7: Perform and Interpretation of routine haematological investigations like haemoglobin, TLC, DLC, ESR PCV, Blood indices and peripheral smear

EPA :					
1.Description of the	Resident should be able to Perform and				
activity:	Interpretation of routine haematological				
	investigations like haemoglobin, TLC, DLC,				
	ESR PCV, Bloo				
2. Most relevant		d marces and	periprici	ai silicai.	
domains of	MK, PS, ICS, P				
competence:					
3. Sub-	MK 7				
Competencies	PS 9				
within each	ICS 2				
domain:	P1				
	SBP 1				
	PBLI 1				
4. Methods of	- MK and P	S will be asso	essed by	faculty	
assessment	during gro	ossing in histo	patholog	y	
	postings, Written exam & eportfolio.				
	- Communication skills and				
	professionalism will be assessed by				
	multisource feedback.				
	- Assessment done during the end posting				
5. Resident will be	test. MK	7 L4			
entrustable when	WIK	/ L4			
this sub	PS	9 L3			
competency	ICS	2			
milestones levels		L2			
are attained.	Р	1			
		L2			
	SBP	1			
		L3			
	PBLI	1			
	I DLI	L3			
		LJ			

EPA :				
1.Description of the activity:	Resident should be able to Perform and			
	Interpretation of	special invest	stigations lik	se
	Reticulocyte co	unt, Sickling	test, Osmot	ic
	Fragility Test, H	laemoglobin		
	Electrophoresis,			
2. Most relevant				
domains of	MK, PS, ICS, P			
competence:				
3. Sub-Competencies	MK 7			
within each	PS 10			
domain:	ICS2			
	P1			
	SBP 1			
	PBLI 1			
4. Methods of	- MK and I	PS will be ass	essed by	
assessment	-	ring grossing		
	-	ology posting	s, Written	
	exam & e	-	and	
	- Communication skills and professionalism will be assessed by			
	-	ce feedback.		у
		nt done durin	ig the end	
	posting te		~	
5. Resident will be	MK 7	L3		
entrustable when	PS 10	L4		
this				
subcompetency milestones levels	ICS2	L2		
are attained.	P1	L2		
	SBP1	L3		
	PBLI 1	L3		

EPA 8: Perform and Interpretation of special investigations like Reticulocyte count, Sickling test, Osmotic Fragility Test, Haemoglobin Electrophoresis,etc

EPA 9:	Planning investigations for a Clinical case
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EPA :			
1.Description of the activity:	Resident should a particular cas	d be able to plan for e	investigating
2. Most relevant domains of competence:	MK, PS, ICS, I	2	
3. Sub- Competencies within each domain:	MK 9 PS 11 ICS 2 P1		
	SBP 1 PBLI 1		
4. Methods of assessment	during g Written o - Commun will be a	PS will be assessed rossing in histopatho exam & eportfolio. nication skills and pr ssessed by multisou ent done during the	ology postings, rofessionalism rce feedback.
5. Resident will be entrustable	MK	9 L3	
when this sub competency	PS ICS	11 L4 2 L2	
milestones levels are	Р	1 L3	
attained.	SBP PBLI	1 L3 1 L3	

EPA 10: Perform and Interpret Urine Examination, Body fluids and semen analysis.

PA:						
1.Description of the activity:	Resident should be able to perform and Interpret					
	Urine Examination	ation, Body fluid	ds and seme	n		
	analysis.	4				
2. Most relevant domains of competence:	MK, PS, ICS,	Р				
3. Sub-Competencies						
within each domain:	PS 12					
	ICS1&2					
	P1 SBP 1					
	PBLI 1					
4. Methods of assessment	during g Written - Commu will be a	I PS will be asse grossing in histo exam & eportfo inication skills a assessed by mult nent done during	pathology p lio. nd profession tisource feed	ostings onalism dback.		
5. Resident will be	МК	9 L3				
entrustable when this subcompetency	PS	12 L 4				
milestones levels are	ICS	1 L2	2 L2			
attained.	Р	1 L2				
attained.	P SBP	1 L2 1 L3				

EPA 11: Interpretation of ancillary techniques like Immunoflourescence, Karyotyping, FISH, PCR and ElectronMicroscopy.

PA:					
1.Description of the	Resident should know the techniques and				
activity:	Interpretation of ancillary techniques like				
	Immunoflourescence, Karyotyping, FISH, PCR ar	nd			
	ElectronMicroscopy.				
2. Most relevant domains of competence:	MK, PS, ICS, P				
3. Sub-Competencies	MK 10				
within each domain:	PS 13				
uomam.	ICS 2				
	P1				
	SBP 1				
	PBLI 1				
4. Methods of	- MK and PS will be assessed by faculty duri	_			
assessment	grossing in histopathology postings, Writter exam & eportfolio.	1			
	 Communication skills and professionalism 	wi			
	be assessed by multisource feedback.				
	- Assessment done during the end posting tes	t.			
5. Resident will be	MK 10 L3				
entrustable when this	PS 13 L4				
subcompetency	ICS 2 L2				
milestones levels are attained.	P 1 L2				
તા દ તાપતાાણા.	SBP 1 L3				
	PBLI 1 L3				

EPA 12: Demonstration of familiarity within laboratory investigations in Microbiology and Biochemistry

1.Description of the activity:	Resident should	be able to de	emonstrate the	
	familiarity with	in laboratory	investigations	
	in Microbiology	and Biocher	mistry.	
2. Most relevant domains of	MK, PS, ICS, P			
competence:				
3. Sub-Competencies	MK 9			
within each	PS 14			
domain:	ICS 2			
	P1			
	SBP 1			
	PBLI 1			
4. Methods of	- MK and PS will be assessed by			
assessment	faculty during grossing in			
	-	ology posting	gs, Written	
	exam & eportfolio. - Communication skills and			
	professionalism will be assessed by			
	multisource feedback.			
	- Assessme	ent done duri	ng the end	
	posting te		1	
5. Resident will be	MK	9		
entrustable when		L3		
this subcompetency milestones levels	PS	14		
are attained.		L3		
	ICS	2		
		L2		
	Р	1		
		L2		
	SBP	1		
		L3		
	PBLI	1		
		L3		

EPA 13: Perform and interpret Blood banking techniques (Blood grouping, Rh typing, Cross-matching and Coomb's test, ELISA for infectious disease)

EPA:						
1.Description of the activity:	Residents should be able Perform and interpret Blood banking techniques (Blood grouping, Rh typing, Cross- matching and Coomb's test, ELISA for infectious disease)					
2. Most relevant domains of competence:	MK, PS, ICS, P					
3. Sub-Competencies within each domain:	MK 8 PS 15 ICS1&2 P1 SBP 1 PBLI 1					
4. Methods of assessment	 MK and PS will be assessed by faculty during grossing in histopathology postings, Written exam & eportfolio. Communication skills and professionalism will be assessed by multisource feedback. Assessment done during the end posting test. 					
5. Resident will be entrustable when this subcompetency milestones levels	MK PS ICS P	8 L3 15 L4 1 L3 1 L2	2 L3			
are attained.	SBP PBLI	1 L2 1 L3 1 L3				

EPA 14: Selection of blood donors and Management of adverse donor reactions. Perform investigation for a case of mismatched blood transfusion.

EPA:						
1.Description of the	Resident should be able to do selection of blood					
activity:	donors and Management of adverse donor reactions.					
	Perform investigation for a case of mismatched bloo					
	transfusion					
2.Most relevant domains of competence:	MK, PS, ICS, P					
3.Sub-Competencies	MK 8					
within each domain:	PS 16 & 17					
	ICS1&2 P1					
	SBP 1					
	PBLI 1					
4.Methods of	- MK and PS will be assessed by faculty during					
assessment	grossing in histopa	athology postin	igs, Writte	n exam &		
	eportfolio.					
	- Communication	skills and profe	essionalisr	n will be		
	assessed by multis	source feedbacl	κ.			
	- Assessment done	e during the end	d posting t	est.		
5.Resident will be	МК	8 L4				
entrustable when this	PS	16 L3	17 L4			
subcompetency	ICS	1 L2	2 L2			
milestones levels are	Р	1 L3				
attained.	SBP	1 L3				
	PBLI	1 L3				

EPA 15: Participation and Presentation in multidisciplinary meetings like tumor boards, CPCs, Dermato-Pathological conferences. Teaching pathology to undergraduates (MBBS), and allied health sciences like BDS, BSc (Nursing), BSc (MLT), BSc (Radiology), etc.

1.Description of the	Resident should be able to participation and					
activity:	Presentation in m	ultidisciplinar	y meeting	s like		
	tumor boards, CPCs, Dermato-Pathological					
	conferences. Teaching pathology to undergraduate					
	(MBBS), and allie	ed health scien	ices like B	BDS, BSG		
	(Nursing), BSc (N	ALT), BSc (Ra	adiology),	etc.		
2. Most relevant						
domains of	MK, PS, ICS, P					
competence:						
2 Sel Courseter in	MIZ 1.0.5					
3. Sub-Competencies within each	MK 1&5					
domain:	PS 18& 19					
uomum	ICS 2&3					
	P1					
	SBP 1					
	PBLI 1		11 0	1,		
4. Methods of assessment		S will be assess using in histopa	•	•		
assessment		am & eportfoli		osungs,		
		ation skills and		onalism		
		essed by multis	-			
		t done during	1	sting test		
5. Resident will be	МК	1 L3	5 L3			
entrustable when this subcompotency	PS	18 L3	19 L3			
this subcompetency milestones levels	ICS	2 L2	3 L2			
are attained.	P P	1 L				
	Г	1 L 3				
	SBP	1 L				
	SDF	1 L 3				
	PBLI	1 L 3				
		3				

9.1.5

9.1.6 Mapping of EPA to Programme Outcomes (PO)

Table 4 showing mapping of the EPA's to the Programme outcomes (*Tick the boxes as appropriate*)

	PO1.	PO2.	PO3.	PO4.	PO5.	PO6.	PO7.	PO8.	PO9.
EPA1.	~			~		\neg			~
EPA2.	~			~					~
EPA3.	\checkmark			✓					~
EPA4.	~			\checkmark					~
EPA5.	~			✓		~		~	~
EPA6.	✓			~					~
EPA7.	~			\checkmark					~
EPA8.	~			\checkmark					~
EPA9.	~		~						~
EPA10.	~								~
EPA11.	\checkmark		V	~		×		~	~
EPA12.			~		~				~
EPA13.		~	✓	×	~				~
EPA14.		✓	~	\checkmark		~			~
EPA15.							\checkmark	\checkmark	~

9.2 Summative assessment

9.2.1 Dissertation

Objectives

- 1. The student should be able to demonstrate capability in research by planning and conducting systematic scientific inquiry & data analysis and deriving conclusion.
- 2. Communicate scientific information for health planning.

Guide for dissertation

- 1. Chief guide will be allocated from the Department of Anesthesiology.
- Co guides can be selected from within the department or from other disciplines related to the dissertation topic.

Submission of dissertation protocol

It should be submitted at the end of six months after admission in the course, in the format prescribed by the institute:

- 1. Protocol in essence should consist of:
 - a) Introduction and objectives of the research project.
 - b) Brief review of literature
 - c) Suggested materials and methods, and (scheme of work)
 - d) Statistician should be consulted at the time of selection of groups, number of cases and method of study. He should also be consulted during the study.
 - e) Bibliography
- 2. The protocol must be presented in the Department of Anesthesiology before being forwarded to the Institutional Research Committee (IRC) for review.
- 3. Protocol must be approved by the research committee, which is appointed by the Dean / Principal to scrutinize the dissertation protocol in references to its feasibility, statistical validity, ethical aspects, etc.
- 4. Once approved by the IRC, the protocol will be forwarded to the Institutional Human Ethics Committee (IHEC) for review.
- After presentation and approval of the protocol by the IHEC, the dissertation must be registered in the Clinical Trial Registry of India - <u>http://ctri.nic.in</u>, following which data collection may be initiated.

Submission of dissertation

- 1. The dissertation shall relate to the candidates own work on a specific research problem or a series of clinical case studies in accordance with the approved plan.
- 2. The dissertation shall be written in English, printed or typed double line spacing, on white bond paper 22x28 cm with a margin of 3.5 cm, bearing the

matter on one side of paper only and neatly bound with the title, the name of the College and University printed on the front cover.

- 3. The dissertation shall contain: Introduction, review of literature, material and methods, observations, discussion, conclusion and summary and reference as per index medicus.
- 4. Each candidate shall submit to the Dean four copies of dissertation, through their respective Heads of the Department not later than six months prior to the date of commencement of theory examination in the subject.

Evaluation of Dissertation:

- The dissertation shall be referred by the University for Evaluation, to External Examiners appointed by the University. The examiners will evaluate and report independently to the Controller of Examinations using Proforma for Dissertation Evaluation Form and recommend whether the dissertation
 - a. Accepted as submitted
 - b. Accepted pending modification as suggested
 - c. Not Accepted for reasons specified
- 2. The dissertation shall be deemed to be accepted when it has been approved by at least two external examiners, who will allocate marks from which an average will be taken.
- 3. If the dissertation is rejected by one of the external examiners it shall be referred to another external examiner (other than the one appointed for initial evaluation) whose judgment shall be final for purposes of acceptance or otherwise of the dissertation.
- 4. Where improvements have been suggested by the external examiners, the candidate shall be required to re submit the dissertation, after making the required improvements for evaluation.
- 5. When a dissertation is rejected by the examiners, it shall be returned to the candidate who shall have to rewrite it. The second version of the dissertation, as and when submitted shall be treated as a fresh dissertation and processed.
- Acceptance of dissertation submitted by the candidate is a pre condition for his / her admission to the written, oral and practical / clinical part of the examination.

- a. Provided that under special circumstances if the report from one or more examiners is not received by the time the Post - Graduate examination is due, the candidate may be permitted provisionally to sit for the examination but the result be withheld till the receipt of the report, subject to the condition that if the dissertation is rejected then the candidate in addition to writing a fresh dissertation, shall have to reappear for the examination.
- 7. A candidate whose dissertation stands approved by the examiners but fails in the examination, shall not be required to submit a fresh one if he/she appears in the examination in the same branch on a subsequent occasion.



9.2.2 Eligibility Criteria

- Candidates will be eligible to appear for the university examinations after completion of 3 years and when following criteria are fulfilled:
 - 1. Attendance of 80%
 - 2. Submission of dissertation and acceptance by external examiner
 - 3. One research Publication based on the Dissertation
 - 4. One poster and one Podium presentation at National or Regional conferences, recognised by Theory (Subject contents already outlined in syllabus)

9.2.3 Theory

- Final Theory Papers: 4 papers
- All papers should have 10 short answer questions.
- Question papers are prepared based on the prescribed blueprint described later (see blueprint section)
- Model question paper is attached for ready reference.

9.2.4 Practical

Each student will be evaluated with all the components of Practical and viva-voce

• Practical(300)

1. Clinical Pathology: Discussion of a clinical case history, plan relevant investigations of the above case, perform them..

2. Haematology: Discuss haematology cases given the relevant history Plan relevant investigations Perform atleast two tests: one routine and one special exercise. Identify

3. Electrophoresis strips, osmotic fragility chart etc. Examine report and discuss ten cases given the history and relevant blood smears and/or bone marrow aspirate smears.

4. Transfusion (Medicine): Perform blood grouping or cross matching or Direct coomb's test

5. Histopathology (Cytopathology): Examine, report and discuss fifteen histopathology and five cytopathology cases given the relevant history and slides. Perform a Haematoxylin and Eosin stain and any special stain on a paraffin section Report on a frozen section

6. Autopsy: Given a case history and relevant organs (with or without slides) give a list of anatomical diagnosis in a autopsy case.

7. Gross Pathology : Describe findings of at least 10 gross specimens, give diagnosis and identify the sections to be processed

8. Basic Sciences : Identify electronmicrographs, Identify gels, results of PCR, immunological tests including staining for direct/indirect immunofluorescence, Identify histochemical and immunohistochemistry stains

9.1.3. Viva-voce is expected to be conducted at every stage of the practical examination. Additionally a formal "grand" viva-voce may be held at the end of the practical examination. Questions on the thesis/dissertation may be asked at this time.

Duration: 2 Days

1. Histopathology & Cytology
2. Autopsy
3. Surgical Pathology
4.Hematology case discussion and exercised including i) Hb/TLC/DLC/PBS
ii) Special Haematology Exercise
5. Transfusion Medicine (Blood Group / DCT/ cross matching)
6. Haematology slides
7. Clinical Pathology case discussion and Exercise (Urine examination)20 marks
8. Histotechnique –20 marks
Microtomy10 marks
H & E stain05 marks
Special stain05 marks
Total 200 marks
VIVA 100 marks
GRAND TOTAL 300 marks

- Recommendations for passing:
- The candidate will be required to secure minimum 50% marks in theory and 50% marks in Practical and viva - voce separately, which is mandatory for passing the whole examination.
- 2. There will be enough gap between theory and practical examination as recommended by NMC rules.
- 3. There university practical examination will be conducted by 2 external and 2 internal examiners.

10 Blueprint of Theory exam paper

Paper I: General Pathology, Pathophysiology, Immunopathology and Cytopathology

		b	<i></i>		M :	
:MĐ	b m	D	fft fft		ttı	m
1	General	Cell injury & Cellular				
	Pathology	adaptation,				
		Inflammation, Tissue	29	29	2	
		repair, Hemodynamic				
		disorders, Neoplasia,				
		Genetics disorder,				
		infectious disease,				
		childhood and				
		environmental disease.				
2	Pathophysiology	Cell injury & Cellular				
		adaptation,				
		Inflammation, Tissue	29	29	2	
		repair, Hemodynamic				
		disorders, Neoplasia,				
		Genetics disorder,				
		infectious disease,				
		childhood and				
		environmental disease.				
3	Immunopathology	Immunological disorder,				
		Autoimmune disorders,	19	19	1	
		Immunodeficiency				
		disorder and AIDS				
4	Cytopathology	Cytology of various				
		organs, Body fluids for	19	19	1	
		malignancy, Various				
		cytotechnique and				
		cytostains. Quality				
		control in cytology				

Paper II Systemic pathology

SI.NO	b m	b	fft fft		M: tti	m
		Pathogenesis,				
1	Systemic	Morphology of	500/	50	-	
1	pathology	diseases of	50%		5	
		various systems,				
		Morphology				
		reporting criteria,				
		staging and				
		grading of				
		various diseases,				
2	Surgical pathology	Histopathology	50%	50	5	
		techniques,				
		Automation and				
		quality control in				
		histopathology				
		lab.				

Paper III Hematology, Transfusion Medicine and Laboratory Medicine

SI.NO	b m	b	fft fft		M: tti	m
1	Hematology	RBC disorders, WBC disorders, Platelet and coagulation disorders, Hematological techniques	59	59	5	
2	Transfusion medicine	Basic immunology, ABO and Rh groups, Clinical significance of other blood groups, Transfusion therapy including the use of whole blood and RBC concentrates, Blood component therapy, Rationale of pre-transfusion testing, Infections	19	19	1	

	transmitted in			
	blood, Adverse			
	reactions to			
	transfusion of			
	blood and			
	components			
	and Quality			
	control in			
	blood bank.			
	Body fluids			
	interpretation,			
	Renal function			
	tests, Liver			
	function test,			
	Pancreatic			
	function test,			
	Endocrine			
Laboratory medicine	function test,			
	Tests for			
	malabsorption,			
Ť	automation in			
	the laboratory			
	and Quality			
	control in the			
	laboratory.	19	19	1
	Laboratory medicine	blood, Adverse reactions to transfusion of blood and components and Quality control in blood bank. Body fluids interpretation, Renal function tests, Liver function test, Pancreatic function test, Endocrine function test, Endocrine function test, Tests for malabsorption, automation in the laboratory and Quality control in the	blood, Adverse reactions to transfusion of blood and components and Quality control in blood bank. Body fluids interpretation, Renal function tests, Liver function test, Pancreatic function test, Pancreatic function test, Function test, audomation in interpretation, automation in the laboratory and Quality control in the	blood, Adversereactions toreactions totransfusion ofblood andcomponentsand Qualitycontrol inblood bank.Body fluidsinterpretation,Renal functiontests, Liverfunction test,Pancreaticfunction test,Fandocrinefunction test,Tests formalabsorption,automation inthe laboratoryand Qualitycontrol in the

Paper IV: Recent advance and applied aspects

	b m	b	fft	fft		М:	
SI.NO		U				ttı	m
		Recent advances in					
1.	Recent	histopathology,	50%		50	5	
1.	advances	Cytology, Hematology,	50%		50	5	
		Transfusion Medicine					
		Applied aspects in					
		Immunopathology,					
		Electron microscopy,					
		Histochemistry,					
		Immunohistochemistry,					
		Cytogenetics,					
2.	Applied	Molecular Biology,	500/			-	
Ζ.	aspects	Maintenance of	50%		50	5	
		records, Information					
		retrieval, use of					
		Computer and Internet					
		in medicine, Quality					
		control, waste disposal.					

11 Model Question Paper

PAPER I : General Pathology, Pathophysiology, Immunopathology and Cytopathology

3 Hours

(10 x 10 = 100 marks)

ANSWER ALL QUESTIONS

(Draw labelled diagram wherever required)

- Q1. Explain role of apoptosis in Health and Disease
- Q2. Describe chemokines and their role in inflammation
- Q3. Describe Role of Myofibroblast in health and disease
- Q4. 35 years old male presented to ICU with 3 days old perforation. Posted for emergency lapratomy. He had fever, tachypneic and tense abdomen with bilateral crypts. His Pulse- 130/mts and Bp is 80/60 mm of Hg. Total WBC count-19000 cells/cu mm.
 - a. what is the possible diagnosis?
 - b. Describe the etiopathogenesis of this disease?
- Q5. Explain Implications of Genomic imprinting in human disease
- Q6. 40 yrs old female presenting with butterfly rash on her face. She does not use any medication. Also presented with arthralgia, alopecia and weakness but there is no fever.
 - a. what is the possible diagnosis?
 - b. Describe the lab diagnosis of this disease?
- Q7. 35 years old female came with complaints of fever, sweating and dizziness. Also shows nocturnal rise in the temperature. Complete blood count shows thrombocytopenia.
 - a. what are the possible differential diagnosis?
 - b. what the various investigation and diagnostic finding for the disease.
- Q8. Explain Role of angiogenesis in Neoplasia
- Q9. Describe cytofixatives
- Q10. Role of squash cytology in CNS tumors.

PAPER II Systemic Pathology

3 Hours

(10X10=100 marks)

(Draw labelled diagram wherever required)

ANSWER ALL QUESTIONS

Q1. Describe vasculitis syndromes

Q2. Describe Pathology of pneumocystis carinii pneumonia

Q3. Explain role of endoscopic biopsy in diagnosis of gastrointestinal lesions

Q4. Explain Radiological appearances in correlation with pathological changes in various bone tumors

- Q5. A 35 year old daily labourer presents with a history of coughing with expectoration for the past 2 months. He has loss of weight and evening rise of temperature. X-ray reveals a cavitary lesion in the upper lobe apex.
 - a) What is the provisional diagnosis?
 - b) Describe the morphology and classification of the diseases
- Q6 Explain the pathology of neurodegenerative disorders

Q7. Describe Microscopic variants of papillary carcinoma of thyroid.

- Q8. 30 years old male with recent consumption of unknown drug, pain in the loin, passing dark colored urine. On withdrawing blood plasma is pink in color. Hb-6gm/dl, reticulocyte count-12%
 - a) What is the probable diagnosis
 - b) Describe the renal biopsy findings of the diagnosis
- Q9. 30 year old female presented with blister over the face, scalp and upper chest. Lesion also seen in the oral mucous membrane.
 - a) What is the probable diagnosis?

- b) How will you approach for vesiculobullous diseases.
- Q10. Discuss the differential diagnosis of spindle cell sarcoma.

PAPER III Hematology, Transfusion Medicine and Laboratory medicine

3 Hours

(10X10=100 marks)

ANSWER ALL QUESTION

(Draw labelled diagram wherever required)

- Q1. Explain Lab diagnosis of cold agglutinin disease
- Q2. Explain Immunophenotyping and cytogenetics of acute leukemias
- Q3. A 10 years old male child came with complaints of petechiae and purpura. His platelet count is 60000 cells/cu mm.
 - a. What is your probable diagnosis?
 - b. How will you approach to any bleeding disorder?

Q4. Define quality control and quality assurance. Discuss internal and external quality control programmes with specific reference to haematology

Q5. Explain use of Microwave in histopathology

Q6. Explain Diagnostic application of microscopic examination of urine

Q7. Explain role of enzymes in health and disease

- Q8. 50 years old male came with complaints of polyuria and polydipsia. His FBS-200 mg/dl and urine analysis shows positive for microalbumninuria.
 - a. what is glycosylated hemoglobin? Expalin its importance?
 - b. Describe the morphology of renal lesions in this disease?
- Q9. Describe in brief the standard protocol and requirement in establishing modern blood bank
- Q10. 35 years old female develops chest pain, palpitation and circulatory shock following blood transfusion in OG ward.
 - a. What is your probable diagnosis?
 - b. How will you investigate for mismatch transfusion reaction.

Paper IV: Recent advance and Applied aspects

3 Hours

(10X10=100 marks)

ANSWER ALL QUESTIONS

(Draw labelled diagram wherever required)

- Q1. Explain role of matrix metalloproteinases in tumor progression.
- Q2. Describe role of microarray analysis in diagnosis of tumors
- Q3. Describe in detail principle, technique and clinical application of flow cytometry
- Q4. Describe liquid based cytology for cervical screening
- Q5. Explain in brief methods of separation of blood components
- Q6. Discuss role of immunohistochemistry in diagnosis of round cell tumors
- Q7. Describe role of Automation in Histopathology laboratory
- Q8. Explain role of karyotyping in haematological malignancies
- Q9. Explain principle and application of Polymerase Chain reaction
- Q10. Explain role of Telepathology in modern Laboratory practice.

12 Recommended reading

12.1List of recommended books

1	Rosai and Ackerman's Surgical Pathology
2	Atlas and Text of Haematology by Tejinder Singh
3	
_	Orell's Atlas of Aspiration Cytology
4	Lever's Dermatopathology
5	Novak's Gynecologic and Obstetric Pathology with Clinical and Endocrine
5	Relations by Edmund R. Novak
6	Bone Pathology by H. Jaffe
7	MacSween's Pathology of the liver
8	Iochim's Lymph Node Pathology
9	Text Book on Breast Pathology by Tavasoli
10	Text Book on Thyroid Pathology by GeethaJayaram
11	Theory and Practice of Histological Techniques by Bancroft
12	Gray's Diagnostic Cytopathology
13	Sternberg's Diagnostic Surgical Pathology
14	Dacie's Practical Haematology
15	Wintrobe's Haematology
16	Heptinstall's Pathology of the Kidney
17	Enzinger's Soft Tissue Tumours

12.2List of recommended journals

S. No	Name of the Journal			
1	Acta Cytologica			
2	The American Journal of Pathology			
3	The American Journal of Surgical Pathology			
4	The American Journal of Hematology			
5	The American Journal of Clinical Pathology			
6	Archives of Pathology and Laboratory Medicine			
7	British Journal of Haematology			
8	Blood			

9	Diagnostic Cytopathology	
10	Histopathology	
11	Human Pathology	
12	Indian Journal of Cytology	
13	Indian Journal of Pathology and Microbiology	
14	Journal of Pathology	
15	Journal of Clinical Pathology	
16	Laboratory Investigation	
17	Modern Pathology	
18	Pathology	
19	Seminars in Hematology	
20	Seminars in Diagnostic Pathology	
21	Virchows Archives	
22	Recent Advances Series	

13 Annexures - Assessment and Feedback forms

Annexure 1 – Multisource Evaluation sheet MAHATMA GANDHI MEDICAL COLLEGE AND RESEARCH INSTITUTE PILLAIYARKUPPAM, PUDUCHERRY – 607 402 Evaluation sheet for postgraduate clinical work

			Score	
S1.	Criteria to be assessed	Below	At par	Above
No.		par	(1)	par
		(0)		(2)
1	INTERPERSONAL COMMUNCATION SKILLS(IPCS)			
1.	Ability to gather the needed information during History taking and physical examination in a respectful manner.			
2.	Ability to give the necessary information regarding choice of management and guide the patient/attenders to make appropriate decisions.			
3.	Ability to communicate the risks involved for patient care, in an understandable language without making the patient/attenders apprehensive, allowing 2 way communication.			
4.	Ability to be caring and respectful with patients during any procedure.			
5.	Ability to convey the required information clearly to the consultants, peers and other health care workers.			
	PROFESSIONALISM(P)			
1.	Ability to be regular and punctual			
2.	Demonstrate respectfulness and obedience to consultants, peers and other health care workers.			
3.	Ability to accept and follow constructive feedback from consultants, peers and other health care workers.			
4.	Ability to maintain emotional balance during triggering situations, people and environment.			
5.	Makes their presence respectful, with their physical appearance and wearing appropriate attire.			
	IPCS Total score: IPCS Final score= IPCS Total score*10			
	Milestone Level: IPCS=1 0 - 20%, IPCS=2 20 - 40%, IPCS IPCS=5 80 - 100%,	=3 40 - 60%	, IPCS=4	60 - 80%,
	P Total score: P Final score= P Total score*10			
	Milestone Level: 0 - 20%, P=1. 20 - 40%, P=2. 40 - 60% 100%, P=5	, P=3. 60 - 3	80%, P=4.	80 -
	Signature:			

Annexure 2–Seminar

MAHATMA GANDHI MEDICAL COLLEGE AND RESEARCH INSTITUTE PILLAIYARKUPPAM, PUDUCHERRY – 607 402 Evaluation sheet for postgraduate seminar

(To be marked individually by each faculty)

▲ Date:

Name of the Resident: UIN No

Name of the Faculty:

S. No.	Criteria to be assessed	*Score (1 – 10)
1	Introduction of subject and its importance / Objectives	
2	Completeness of presentation	
3	Cogency of presentation	
4	Consulted all relevant literature	
5	Use of audio - visual aids	
6	Understanding of subject	
7	Summary and take home message	
8	Cites appropriate references / suggests further reading	
9	Time management	
10	Overall performance – relevant answers to questions, attitude during presentation and confidence	

*Score interpretation – 1-3->Needs improvement; 4-6->Meets expectations; 7-

9->Exceeds expectation; 10->Outstanding.

General Comments:

Highlights in performance (strengths)

Possible suggested areas for improvement (weakness)

Signature

Annexure 3 – Journal Club

MAHATMA GANDHI MEDICAL COLLEGE AND RESEARCH INSTITUTE PILLAIYARKUPPAM, PUDUCHERRY – 607 402 Evaluation sheet for postgraduate journal club

(To be marked individually by each faculty)

Name of the Resident:

UIN No

Date:

Name of the Faculty:

S. No.	Criteria to be assessed	*Score(1-10)
1	Relevance of article chosen	
2	Identifies the problem addressed in the paper	
3	Completeness of presentation	
4	Analyses and gives comments on methodology and statistics	
5	Brief summary of results	
6	Comparison of work with other published work	
7	Merits and demerits of the paper	
8	Summary and take home message	
9	Time management	
10	Overall performance – relevant answers to questions, attitude	
	during presentation and confidence	

*Score interpretation – 1-3->Needs improvement; 4-6->Meets expectations; 7-

9->Exceeds expectation; 10->Outstanding.

General Comments:

Highlights in performance (strengths)

Possible suggested areas for improvement (weakness)

Signature:

Annexure 4 - Case Presentation

MAHATMA GANDHI MEDICAL COLLEGE AND RESEARCH INSTITUTE PILLAIYARKUPPAM, PUDUCHERRY – 607 402 Evaluation sheet for postgraduate case presentation

(To be marked individually by each faculty)

Name of the Resident:

UIN No

Date:

Name of the Faculty:

S. No.	Criteria to be assessed	*Score (1-10)
1	Logical order in presentation (History taking)	
2	Cogency of presentation	
3	Accuracy and completeness of general and local physical examination	
4	Description of Gross and microscopic findings	
5	Summarizes the case and analyses the appropriate differential diagnoses	
6	Whether the diagnosis follows logically from history and findings	
7	Investigations required : Completeness of list, relevant order, interpretation of investigations	
8	Management principles and details	
9	Time management	
10	Overall performance – relevant answers to questions, attitude during presentation and confidence	

*Score interpretation – 1-3->Needs improvement; 4-6->Meets expectations; 7-

9->Exceeds expectation; 10->Outstanding.

General Comments:

Highlights in performance (strengths)

Possible suggested areas for improvement (weakness)

Signature:

Annexure 5 - EPA Assessment Form

MAHATMA GANDHI MEDICAL COLLEGE AND RESEARCH INSTITUTE DEPARTMENT OF PATHOLOGY

Entrustable professional activity assessment form

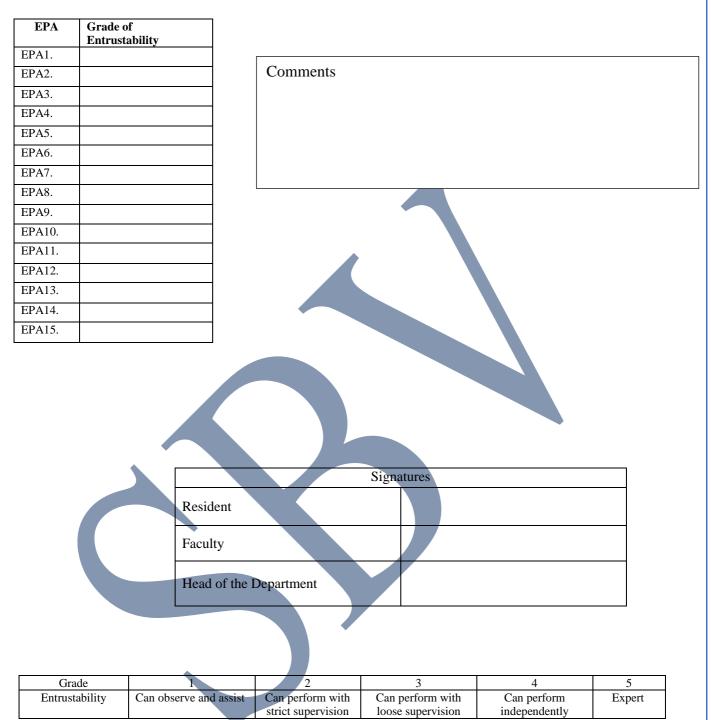
STUDENT NAME: PGY: FACULTY: UIN No: ASSESSMENT No: DATE:

EPA	Entrustable										
No.	Professional Activity										
1.	Perform gross examination of surgical pathology specimens and Interpretation of Microscopic findings in tissue.	PS 1 3	PS2 3	PS3 3	ICS 1 2	ICS2 2	SBP 1 2	PBLI 1 1	P 1 1	MK 1 2	MK 4& 5 2
						\searrow					
2	Perform histotechniques (Tissue processing, embedding, microtomy, Staining and	PS 4 3	ICS 2 2	SBP 1 2	PBLI 1	P1 1	MK 5 2				
	special stainning)										
3.	Perform And Interpretation of FNAC and other Cytological smears (Sputum, Bronchial washings, Serous effusions, etc.	PS5 2	ICS 1 2	ICS 2 2	PC1 2	SBP1 2	PBLI1 1	P 1 1	M K 3 2		
4.	Interpretation of Frozen sections	PS 6 2	ICS 1 2	PC 1 2	SBP 1 2	PBLI 1 1	P 1 1	MK 5 2	2		
5.	Selection, performance and interpretation of appropriate Immunohistochemical markers.	PS 7 2	ICS 1 2	ICS 2 2	SBP 1 2	PBLI 1 1	P1 1	MK 5 2			
6.	Interpretation of Bone Marrow Smears	PS 8 2	ICS 1 2	ICS 2 2	PC 1 2	SBP 1 2	PBLI 1 1	P1 1	M K 7		
									2		

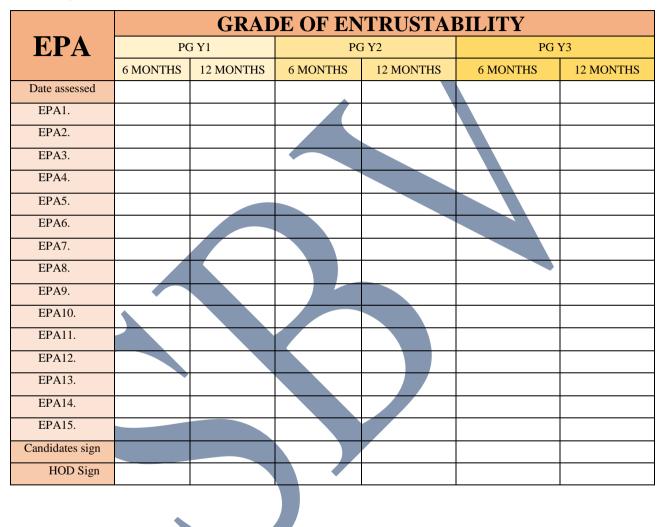
7.	Perform and Interpretation of routine haematological investigations like	PS 9 3	ICS 2 2	SBP 1 2	PBLI 1	P1 1	MK7 2				
	haemoglobin, TLC, DLC, ESR PCV, Blood indices and peripheral smear.										
	Perform and Interpretation of special investigations like Reticulocyte count, Sickling	PS10 2	ICS 2 2	SBP1 2	PBLI1 1	P1 1	MK 7 2				
8.	test, Osmotic Fragility Test, Haemoglobin Electrophoresis, Fetal Haemoglobin, etc.										
9.	Planning investigations for a Clinical case	PS 11 2	ICS 2 2	PC 1 2	SBP 1 2	PBLI 1	P1 1	MK 9 2			
).											
10.	Perform and Interpret Urine Examination, Body fluids and semen analysis.	PS 12 3	ICS 1 2	ICS 2 2	PC 1 2	SBP 1 2	PBLI 1 1	P 1 1	M K9 2		
11	Interpretation of ancillary techniques like Immunoflourescence, Karyotyping, FISH, PCR and	PS 13 2	ICS 2 2	SBP 1 2	PBLI 1 1	P1 1	MK 10 2				
	ElectronMicroscopy.										
12	Demonstration of familiarity within laboratory investigations in Microbiology and Biochemistry	PS 14 2	ICS 2 2	SBP 1 2	PBLI 1 1	P 1 1	MK 9 2				
	Perform and interpret Blood	PS 15	ICS 1	ICS 2	SBP 1	PBLI	P1	MK8			
13	banking tehniques (Bloodgrouping , Rh typing, Cross-matching and Coomb's	2	2	2	2	1 1	1	2			
	test, ELISA for infectious disease like HIV, HbsAg).										
14	Selection of blood donors and Management of adverse donor	PS 16 2	PS17 2	ICS 1 2	ICS 2 2	PC 1 2	SBP 1 2	PBLI 1 1	P1 1	MK8 3	

	reactions. Perform investigation for a case of mismatched blood transfusion.										
	Participation and Presentation	PS 18	PS19	ICS 2	ICS 3	PC 1	SBP 1	PBLI 1	P1	MK1	MK
	in multidisciplinary meetings like tumor boards, CPCs,	2	2	2	2	2	2	1	1	2	5
	Dermato-Pathological										3
15	conferences. Teaching pathology to undergraduates (MBBS), and allied health sciences like BDS, BSc (Nursing), BSc (MLT), BSc (Radiology), etc.										

Key for assigning Grade of entrustability



Annexure 6 – EPA Progress sheet



Annexure 7 – Dissertation evaluation form MAHATMA GANDHI MEDICAL COLLEGE AND RESEARCH INSTITUTE PILLAIYARKUPPAM, PUDUCHERRY – 607 402 Proforma for evaluation of Dissertation

<u>UIN:</u>

Topic of the study :

DISSERTATION COMPONENTS		Grade	
TITLE	•	D	C
Title appropriate and clear	A	В	C
INTRODUCTION		D	G
Purpose of the Study	A	B	C
Hypothesis/Research Question	A	B	C
Aims & Objectives	A	В	С
REVIEW OF LITERATURE			~
Appropriate	A	В	C
Complete and current	A	В	C
METHODS			I
Study subjects, controls, Inclusion and Exclusion criteria	A	В	C
Materials/Apparatus/Cases	А	В	С
Methodology used	А	В	С
Procedure for data collection	А	В	C
Appropriate statistical methods employed	A	В	С
Handling of ethical issues	A	В	С
RESULTS			
Logical organization of data	A	В	С
Appropriate use of charts, tables, Graphs, figures, etc.	А	В	С
Statistical/Clinical interpretation	А	В	С
DISCUSSION		<u>.</u>	
Appropriate to data	A	В	C
Discussion and implication of results	A	В	С
Comparison with other studies	A	В	С
Satisfactory explanation of deviations if any	A	В	С
Limitations of the study	A	В	С
Recommendation for future studies	A	В	С
CONCLUSION		<u> </u>	
Relevance, are they in line with aims	A	В	C
SUMMARY	1	<u> </u>	
Clear and Concise	A	В	C
REFERENCES	1		
	•	π	С
Vancouver Format and appropriately cited in text.	А	В	U

Key for grading – A – Exceeds expectation, B – Meets expectation, C – Needs Improvement

Overall Impression

(Please Check the appropriate box)

Accepted as submitted

Accepted pending modification as suggested below

Not Accepted for reasons specified below

Remarks:

