

INFECTION AND INFLAMMATION MODULE 3RD YEAR MBBS

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Khyber Medical University (KMU) Vision:

Khyber Medical University will be the global leader in health sciences academics and research for efficient and compassionate health care.

Khyber Medical University (KMU) Mission:

Khyber Medical University aims to promote professional competence through learning and innovation for providing comprehensive quality health care to the nation.

Institute of Health Professions Education & Research (IHPER) Mission:

To produce leaders, innovators and researchers in health professions education who can apply global knowledge to resolve local issues.

Teaching Hours Allocation

Table 1: Hours Allocation

S. No	Subjects	Hours
1	Pharmacology	35
2	Pathology	46
3	Forensic medicine	12
4	Community medicine	15
5	Family medicine	2
6	Medicine	1
7	Surgery	3
8	Pediatrics	2
9	Gynaecology	2
10	ENT	5
11	EYE	3
12	PRIME	2
13	Research	5
	Total hours	133

Learning Objectives

At the end of this module, the 3rd year students would be able to:

- 1. Describe the process of acute & chronic inflammation with their outcomes
- 2. Relate different aspects of healing and repair
- 3. Differentiate common pathogenic bacteria based on morphology, pathogenesis & lab diagnosis.
- 4. Relate bacterial pathogenic factors to clinical manifestations of commoninfectious diseases.
- 5. Describe the pharmacological details of anti-inflammatory drugs
- 6. Apply/relate the pharmacokinetics & pharmacodynamics of chemotherapeutic agents to their use in infectious diseases
- 7. Construct / Write prescriptions for various inflammatory and infectious diseases
- 8. Describe medico legal aspects of HIV patient.
- 9. Describe mechanism of wound causation.
- 10. Describe medico legal aspects of parameters used for personal identification inreal life situation
- 11. Apply parameters of a person's identification in a simulated environment
- 12. Describe the epidemiology of common infectious diseases.
- 13. Explain the preventive and control measures for infectious diseases.
- 14. Explain the control & preventive measures for nosocomial infections.
- 15. Describe the risks associated with hospital waste and its management.

Theme-1 (Pain	Theme-1 (Pain and Fatigue)				
Subject To	opic	Hours	Learning objectives		
Pharmacology (Note: Co.	Overview to anti- Inflammatory drugs ISAIDs Non-selective cox Inhibitors: Aspirin & other Iommonly Ised NSAIDs)	1	-Classify anti-inflammatory drugs -Describe the role of DMARDs and glucocorticoids as anti-inflammatory agents -Classify NSAIDS -Differentiate between non-selective COXinhibitors and selective COX-2 inhibitors based on mechanism of action. -Name the prototype non-selective COX inhibitor. -Describe the pharmacokinetics of Aspirin -Describe the mechanism of action of aspirin as anti-platelet, analgesic, antipyretic and anti-inflammatory agent. -Give the dose of Aspirin as anti-platelet, analgesic/antipyretic and as anti-inflammatory drug. -Describe clinical uses of NSAIDs.		

		-Describe the pharmacokinetics with emphasis on
		dosage, duration of action and elimination of
		Diclofenac, Ibuprofen, Indomethacin, Mefanamic
		acid and Piroxicam in contrast to Aspirin
		-Relate pharmacokinetics and pharmacodynamics of NSAIDs to their clinical applications
	1	-Describe the mechanism of action of selective
		COX-2 inhibitors.
		-Describe the clinical uses of selective COX-2
Selective COX-2		inhibitors
inhibitors		-Describe the adverse effects of selective COX-2
		inhibitors
		-Describe the merits and demerits of selective
		COX-2 inhibitors and non-selective COXinhibitors.
		-Describe the pharmacokinetics of Paracetamol
		-Describe the mechanism of action of Paracetamol.
Paracetamol		-Describe the clinical uses of Paracetamol.
(Acetaminophen)		-Describe the adverse effects of Paracetamol.
		-Give therapeutic and fatal doses of Paracetamol.
		-Describe the drug treatment of Paracetamolpoisoning

	T _a	D il lice i il ci di
	1	-Describe different cells of inflammation
		-Describe the functions of various cells ofinflammation
Cells of Inflammation		- Enumerate different causes of leukopenia and
		leucocytosis(each neutrophil, lymphocyte, monocyte, eosinophil,
		basophil seperately)
Overview to Acute	1	-Define acute inflammation
Inflammation and vascular		-Describe causes of acute inflammation
phase		-Describe the vascular events of acuteinflammation
	1	-Describe various molecular patterns and appropriate receptors used by
Decognition of microbas		the inflammatorycells to identify microbes
Recognition of flictobes		-Relate the recognition of microbes to the
		initiation of inflammation
Cellular phase of acute	1	-Describe the sequence of events and cellular
inflammation		changes involved in cellular phase of acuteinflammation
	1	-Enumerate plasma derived mediators
Plasma DerivedMediators		-Enlist the functions of each mediator
		-Describe the different cascades involved in thegeneration of mediators
Cell Derived		-Enumerate cell derived mediators
Mediators		-Enlist the functions of each mediator
	Overview to Acute Inflammation and vascular phase Recognition of microbes Cellular phase of acute inflammation Plasma DerivedMediators Cell Derived	Overview to Acute Inflammation and vascular phase 1 Recognition of microbes Cellular phase of acute inflammation 1 Plasma DerivedMediators Cell Derived

Theme (Pain and Fatigue)				
Subjects	Topics	Hours	Los	
Pharmacology	Anti-histamines	1	-Classify anti-histamines -Differentiate between first and second generationanti-histamines -Describe the pharmacologic effects of H1-receptor antagonists. -Describe the clinical uses of H1-receptorantagonists. -Enlist the adverse effects of H1-receptorantagonists. -Describe the drug interactions of H1-receptorantagonists.	
	Serotonin agonistand antagonist	1	 Enlist serotonin agonists Classify serotonin antagonists Describe the mechanism of action of serotonin Describe the organ system effects of serotonin. Describe the clinical uses of serotonin agonists and antagonists Describe the pharmacological basis of ondansetron in chemotherapy induced vomiting 	

			-Enumerate the different morphological patternsof inflammation
	Morphological patterns,	1	-Describe the histological changes in each pattern
	outcomes, defects of	I	- Enlist the outcomes of inflammation
	inflammation		-Enumerate the various defects of inflammation
			-Describe the consequences of the defects of inflammation
			-Define chronic inflammation
	Overview to chronic	1	-Differentiate chronic from acute inflammation
	inflammation		-Describe the causes and morphological features of chronic
			inflammation
	Granulomatous	1	Define granulomatous inflammation
Pathology	inflammation		-Describe the morphological features and
			mediators involved in granulomatousinflammation
	Cells and mediators of	1	-Enlist the cells of chronic inflammation
	chronic inflammation		-Enumerate the mediators of chronicinflammation
			-Describe the function of the mediators
			-Relate the functions of mediators to themorphological changes
			seen in chronic
Sy			inflammation
	Systemic effects o	f 1	-Enumerate the systemic effects of inflammation
	inflammation		-Describe the pathophysiology of the systemic effects of inflammation
L			

	Antidotes	1	Define and classify antidotes
Forensic			Describe the mechanism of action of differentantidotes
Medicine	Steps of management in a case of poisoning	1	Describe general steps of management in a case of poisoning
	Infectious disease epidemiology	1	 Define incubation period Explain the principles of disease eradication and control Define serial intervals Define infectivity period
Community Medicine	Infection control	2	 Define the basic definition related to infectious disease epidemiology Review the role of susceptible host for successful parasitism, modes of transmission and the host defense system List and explain the various classifications of communicable diseases with special reference to the scope and purpose of the International classification of Disease (ICD -10). Enlist the common infectious diseases affecting the population of Pakistan as perNational institute of Health Pakistan. Explain the effect of climate change and seasonal variation on specific diseases globally and in Pakistan. Explain the role of personal hygiene & PPE in infection control.

Define disease careers
Explain the reservoirs of infection
Differentiate between sterilization and disinfection
Explain the types and procedures of disinfection
Discuss Communicable disease control measure (aimed at agent, host,
others, administrative measures and vector control measures

Theme (Trauma and repair)				
Subjects	Topics	Hours	LOs	
Overview to tist and repair Pathology Tissue regeneration	Prostaglandins	1	 Enlist various prostaglandins- Describe the mechanism of action of Prostaglandins. Describe the organ system effects of Prostaglandins. Describe the clinical uses of Prostaglandins. 	
	Overview to tissuehealing and repair		-Differentiate between regeneration and repair -Describe various steps involved in the process of tissue healing and repair	
	Tissue regeneration	1	-Define regeneration -Enlist organs capable of regeneration -Describe the process and mediators involved in regeneration	
	Cell Cycle and itsrole in repair		-Define cell cycle -Describe the initiation, various phases and proteins involved in the cell cycle -Discuss cells capable of entering the cell cycle -Describe proliferative capabilities of various cells	

	Repair by scarring	1	-Describe the various steps involved in process of repair by scarring -Describe the various mediators involved in thesteps of scarring
	Growth factors and receptors ECM	1	-enumerate various growth factors and their receptors -Describe the most common pathways by which growth factors affect tissue repair and regeneration -Classify various components of ECM -Describe the role and importance of ECM in tissue repair
	Factors affecting wound healing/abnormalscarring	1	-Enlist the various factors that influence woundhealing -Describe the mechanism by which these factors affect wound healing -Describe the abnormalities of repair and their consequences
Forensic	Overview to medico-legal aspects of trauma (Wound causation)	1	Describe mechanism of wound causation
Medicine	Toxicity by analgesics	1	Describe the medico legal aspects of toxicity byaspirin and paracetamol

Community Medicine Nosocomial infection & its control	 Describe the prevalence of the nosocomial infections globally and Specifically in Pakistan. Identify the cause of nosocomial infections in Pakistan. Enlist common nosocomial infections. Describe the importance of different modes of transmission for causation of the nosocomial infections. Explain the control & preventive measures for nosocomial infections
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Theme (Feve	er and Infection)		
Subjects	Topics	Hours	
		2	1. Define basic terms like chemotherapy, antibiotic, antimicrobial, MIC,
			MBC, chemoprophylaxis, empirical therapy and post-antibiotic
	Introduction to		effect, bacteriostatic and bactericidal antimicrobials.
	Chemotherapy		2. Explain advantages of drug combinations.
	chemotherapy		3. Describe various mechanisms of bacterial resistance against
			antibiotics.
			4. Differentiate between concentration and time dependent killing
			with examples.
			5. Classify antimicrobials on the basis of mechanism of action (MOA)
	Penicillins	2	Classify beta-lactam antibiotics
			2. Enlist narrow and broad spectrumPenicillins.
			3. Enlist anti-pseudomonal, anti- staphylococcal/ beta lactamase
			resistantPenicillin.
			4. Enlist long- and short-acting Penicillins
			5. Describe anti-bacterial spectrum of Penicillins.
Pharmacology			6. Describe pharmacokinetics in respect of emphasis on route of
			administration and
			excretion of Penicillins
			7. Describe mechanism of action of Penicillins
			8. Describe clinical uses of Penecillins
			9. Describe adverse effects of Penicillins,

Beta lactamaseinhibitors	1	 Enlist beta-lactamase inhibitors Explain the rationale for using beta lactamase inhibitors in combination with 8-lactam antibiotics.
Серникозронніз		 Describe anti-bacterial spectrum of Cephalosporins. Describe pharmacokinetics of Cephalosporins with special emphasis onroute of administration and excretion. Describe clinical uses of Cephalosporins Describe the adverse effects of Cephalosporins. Describe drug interactions of Cephalosporins with Ethanol. Describe the principal bacterial mechanism of resistance to Cephalosporins. Relate pharmacokinetics and pharmacodynamics of Cephalosporin with their clinical applications / uses.
Cephalosporins		 10. Describe contraindications of Penicillins. 11. Describe principal mechanism of bacterial resistance to Penicillins 12. Describe drug interactions of Penicillins 13. Apply formula for interconversion of milligrams and units of Penicillin G. 14. Relate pharmacokinetics and pharmacodynamics of Penicillin with their clinical applications / uses. 1. Classify Cephalosporins

Monobactams & Carbapanem,	1	Describe the antibacterial spectrum of Monobactams and Carbapanem Describe the clinical uses of Monobactams and Carbapanem
Vancomycin	1	 Describe the MOA of Vancomycin. Describe clinical uses of Vancomycin Describe the use of vancomycin in MRSA (Methicillin-resistant Staph aureus). Describe adverse effects of Vancomycin Describe "Red man/Red neck" syndrome.
Fosfomycin Bacitracin & Cycloserine	1	1. Enlist clinical uses of Fosfomycin, Bacitracin & Cycloserine
Protein synthesis inhibitors:	1	Classify bacterial protein synthesis inhibitors
Tetracyclines	1	 Classify Tetracyclines. Describe anti-bacterial spectrum of Tetracyclines. Describe the pharmacokinetics of Tetracycline with special emphasis onabsorption of Tetracyclines. Describe mechanism of action of Tetracyclines. Describe the principal mechanism of resistance to Tetracyclines. Describe clinical uses of Tetracyclines. Describe adverse effects of Tetracyclines Describe Black Bone disease.

Pathology	Bacteria: Pyrogenic Bacteria Bacteria: Rickettsia	1	 Describe the teratogenic effects of Tetracyclines. Describe drug interactions of Tetracyclines. Describe the adverse effect related to theuse of outdated (expired) Tetracycline products. Relate pharmacokinetics and pharmacodynamics of Tetracycline with their clinical applications / uses. Define boil and furuncle Enlist organisms responsible for pyrogenic infections -Describe important properties, pathophysiology, lab diagnosis of GPC & GNC Define Rickettsia -Describe the important properties, pathophysiology, lab diagnosis of diseases caused by Rickettsia
	Spore forming GProds Non Spore forming GP rods	1	-Enumerate spore forming GP rods - Describe the important properties, pathophysiology, clinical features and labdiagnosis of spore forming GP rods Enumerate non spore forming GP rods

		- Describe the important properties, pathophysiology, clinical features and lab diagnosis of non-spore forming GP rods
Chlamydia	1	Describe the important properties, pathophysiology, clinical features and lab diagnosis of chlamydia.
Miscellaneous: Sepsis and Sept	1 ic	-Define sepsis and septic shock -Enlist organisms capable of causing sepsis and inducing septic shock
Shock		-Describe the pathophysiology and clinical features of septic shock
Zoonotic Infections	1	-Enlist organisms causing zoonotic infections -Describe the important properties, pathophysiology, clinical features and lab diagnosis of different zoonotic diseases
General outlines or identification	f 2	Describe methods and parameters of identification
Fetal age determination		Write important physical developmental stages of fetus for age estimation
Age determination skeletal study	by	Write important skeletal points of age estimation
Age estimation dental study	by	Write important dental points for age estimation
Ages of medico leg significance	al	Enlist important ages of legal significance

Hours 1	Los
1	
	Enlist Aminoglycosides.
	Describe anti-bacterial spectrum of Aminoglycosides.
	Describe the pharmacokinetics of Aminoglycosides with special emphasis on
	route of administration, concentration-dependent killing and post-antibiotic
	effect.
	Describe mechanism of action of Aminoglycosides.
	Describe the principal mechanism of resistance to Aminoglycosides.
	Describe clinical uses of
	Aminoglycosides.
	Describe adverse effects of Aminoglycosides.
	Describe the drug interactions of Aminoglycosides.
	Relate pharmacokinetics and pharmacodynamics of Aminoglycosides
	with their clinical applications / uses.

Macrolides & other 2	Enlist Macrolides.
related drugs	Describe anti-microbial spectrum of Macrolides
	Describe pharmacokinetics of Macrolides
	Describe the mechanism of action of Macrolides
	Describe the principal mechanism of resistance to Macrolides
	Describe clinical uses of Macrolides
	Describe adverse effects of Macrolides.
	Describe drug interactions of Macrolides
	Differentiate the salient features of Erythromycin, Clarithromycin and
	Azithromycin in respect of dosing andclinical use.
	Relate pharmacokinetics and pharmacodynamics of Macrolides with their
	clinical applications / uses.
Linezolid 1	Describe mechanism of action ofLinezolid
	Describe clinical uses of Linezolid with special emphasis on methicillin-
	resistant staphylococci and vancomycin-resistant enterococci
Clindamycin	Describe mechanism of action of Clindamycin.
	Enumerate clinical uses of Clindamycin.
	Describe antibiotic-associated (pseudomembranous) colitis.
Streptogramins	Enumerate Streptogramins.
	Describe clinical use of Quinupristin-
	Dalfopristin in VRE (Vancomycin-resistant enterococci).

	1	Describe anti-microbial spectrum of Chloramphenicol
		Describe mechanism of action of Chloramphenicol
Chloramphenicol		Enlist clinical uses of Chloramphenicol
		Describe the reason for obsoleting thesystemic use of Chloramphenicol
		Enlist adverse effects of Chloramphenicol
Quinolones	1	Describe Gray baby syndrome.
		Classify Quinolones.
		• Describe the pharmacokinetics of Fluroquinolones with special emphasis onhalf-
		life of Moxifloxacin
		Enlist respiratory Quinolones.
		Describe anti-microbial spectrum ofFluoroquinolones.
		Describe mechanism of action of Fluoroquinolones.
		 Describe the principal mechanism of resistance to Fluroquinolones,
		Describe clinical uses of Fluroquinolones
		Describe adverse effects of Fluroquinolones
		Describe drug interactions of Fluroquinolones
		Relate pharmacokinetics and pharmacodynamics of Fluoroquinolones
		with their clinical applications / use.

2	Classify Sulfonamides
Sulfonamides and	Describe anti-microbial spectrum of Sulfonamides
Trimethoprim	Describe mechanism of action of Sulfonamides and Trimethoprim
	Describe mechanism of resistance to Sulfonamides
	Describe clinical uses of Sulfonamides and Trimethoprim
	Describe adverse effects of Sulfonamidesand Trimethoprim
	Describe the advantages of combining sulfamethoxazole with trimethoprim (Co-
	Trimoxazole)
	Describe the drug interaction of
	Sulphonamides with Phenytoin.
Parasites: HydatidCyst 1	Describe the life cycle and important properties of Echinococcus
	Relate the pathogenesis to the clinical features and lab work up of Echinococcus
	Identify cysts of Echinococcus in the lab
Leishmania	Describe the life cycle, and important properties of Leishmania
	 Relate the pathogenesis to the clinical features and lab work up of Leishmania

	Toxoplasma	2	Describe the life cycle and important properties of Toxoplasma
Pathology			 Relate the pathogenesis to the clinical features and lab work up of Toxoplasma
	Malaria		 Describe the life cycle and important properties of Malarial parasite Relate the pathogenesis to the clinical features and lab work up of Malaria
	Tenia		 Describe the life cycle, important properties, of Tenia saginata and solium Relate pathogenesis to the clinical features and lab work up of Tenia saginata and solium
	Sex determination	2	Describe parameters of sex determination
Forensic Medicine	Race determination		Describe parameters of race determination
medieme	Examination ofhair		Describe medico legal aspects of hair
	Forensic odontology		Write the application of odontology in forensicmedicine
	Forensic Anthropometry		Describe medico legal aspects of forensicanthropometry

	Epidemiology and control2	Describe the epidemiological determinants, frequency and distribution
	of vector borne diseases	of Malaria
	• Malaria	Compare the prevalence/incidence of malaria in different provinces of
	Dengue and other	Pakistan.
	Viral haemorrhagic	Explain the preventive and controlmeasures of Malaria
Community	fevers and	Describe the scope/function of Malaria control program.
Medicine	Arboviral	Explain the types, risk factors, complications and control measures of
	infections	viral hemorrhagic fevers including Dengue fever
	• Plague	
	 Filariasis 	
	Epidemiology & control 1 of Leishmaniasis	 Describe the epidemiological determinants, frequency and distribution of Leishmaniasis Explain the preventive and controlmeasures of Leishmaniasis
	zoonotic and direct2	Explain the pre and post exposure prophylaxis of Rabies
	contagious diseases	Explain the epidemiology, types of Anthrax and its preventive measures
	 Rabies 	Discuss the history, types and prevention of Plague
	• Anthrax	Explain the etiology, risk factors, clinical features and prevention of
	• Plague	Brucellosis
	 Brucellosis 	Explain the preventive measures of Scabies
	• Tetanus	Discuss the etiology, risk factors, clinical features and prophylaxis of
	 Scabies 	pre and post exposure of Tetanus

Leprosy	Explain the etiology, risk factors, stages and preventive measures of
Trachoma	Leprosy
	Explain the etiology, risk factors, complications and preventive measures of Trachoma
Malaria & Hepatitis1 Family medicine control program teams	Explain the etiology, clinical features, types, investigations and management of Malaria in family practice
	Describe the red-flags in a patient with Malaria for referral to specialty care
	Identify at risk patients of hepatitis and Malaria and offer them screening

Theme (Feve	er and Infection	on)	
Subjects	Topics	Hours	Los
Pharmacology		3	Describe terms like chemoprophylaxis, causal prophylaxis, terminal
	Antimalarials		prophylaxisand radical cure with examples of drugs.
			Classify antimalarial drugs.
			Enlist drugs used for chemoprophylaxis ofmalaria.
			Enlist drugs used for radical cure ofmalaria.
			Describe the pharmacokinetics of Chloroquine with special emphasis
			onvolume of distribution and dosing
			• Describe mechanism of action of Chloroquine, Quinine, Mefloquine,
			Halofantrine, Primaquine, Pyrimethamine and Artemisinins.
			Describe adverse effects of antimalarial drugs
			Describe Cinchonism and Blackwaterfever.
			Enlist the antimalarial drugs relativelysafe in pregnancy.
			Describe the antimalarial drugs contraindicated in G6PD deficiency.
			Relate pharmacokinetics and pharmacodynamics of antimalarial drugs with
			their clinical applications / use.

Antifungal drugs 2	• C • C • T • C	Classify Antifungal drugs. Describe the pharmacokinetics of Amphotericin B and Ketoconazole Describe the advantages of liposomalpreparation of Amphotericin B Describe mechanism of action of Azoles, Amphotericin B, Griseofulvin, Turbinafine, and Nystatin. Describe clinical uses of Azoles, Amphotericin B, Griseofulvin, Turbinafine, and Nystatin. Describe adverse effects of Azoles, Amphotericin B, Griseofulvin, Turbinafine, and Nystatin. Describe drug interactions of Ketoconazole and Amphotericin B
Antivirals 1 Anti-herpes 1	• E • C • C	Classify antiviral drugs Chlist anti- Herpes drugs Describe the pharmacokinetics of Acyclovir Describe mechanism of action of Acyclovir Describe clinical uses of Acyclovir. Describe adverse effects of Acyclovir Describe the role of Ganciclovir in CMV retinitis.
Anti-HIV drugs 3	• (Classify anti-HIV drugs.

			 Describe the role of entry inhibitors, integrase inhibitors, protease inhibitors, NRTIs and NNRTIs in HIV treatment Describe adverse effects of Zidovudine and Indinavir Describe the rationale of HAART therapy.
	Viruses: Corona	1	Describe the structure, important properties, pathogenesis and clinical features along with labwork up of Corona Virus
	Viruses: HIV		Describe the structure, important properties, pathogenesis and clinical features along with labwork up of HIV
	Viruses: Herpesviruses	1	Describe the structure, important properties, pathogenesis and clinical features along with labwork up of Herpesviruses
	Viruses: Tumor Viruses		Describe the structure, important properties, pathogenesis and clinical features along with labwork up of Tumor viruses
Pathology	Viruses: MMR		Describe the structure, important properties, pathogenesis and clinical features along with lab work up of MMR viruses
	Fungi: Aspergillus	1	Describe the structure, important properties, pathogenesis and clinical features along with labwork up of Aspergillus
	Fungi: Candida		Describe the structure, important properties, pathogenesis and clinical features along with lab work up of Candida

	Tenia	Describe the structure, important properties, pathogenesis and clinical features along with labwork up of Tenia
	Medico legal 1 issues related to	Describe legal issues related to HIV patient
	HIV patient Dactylography	Describe medico legal aspects of dactylography
Forensic Medicine	DNA finger printing	 Define DNA finger printing Write its application in forensic practice Write methods of collection of samples and dispatch to laboratory
	Tattoos, scarmarks, Superimposition	 Describe medico legal aspects of tattoo marks, Describe medico legal aspects of scar tissue, Describe medico legal aspects of superimposition

	and facial		Describe medico legal aspects of facial reconstruction
	reconstruction		
	Polygraph		Describe medico legal aspects of polygraph
	Narcoanalysis		Describe medico legal aspects of
			narcoanalysis
Family Medicine	TORCH infections	1	Define TORCH infection
			Describe the steps of investigations for TORCH infections
			Describe the preventive strategies for TORCH infections and their complications

	Epidemiology &	1	Describe the epidemiological determinants, frequency and
	control of airborne		distribution of measles, mumps, chickenpox, rubella,
	diseases		Diphtheria, Pertissus and meningitis
			grand and and and and and and and and and
Community			
Medicine			
			Explain the preventive and control
			measures of measles, mumps & rubella with reference to
			Pakistani context.

Epidemiology & control of Corona virus infection	1	 Describe the epidemiological determinants, frequency and distribution of corona Compare the prevalence/incidence of corona in different parts of the world. Describe the preventive and controlmeasures of corona Describe the role of Pakistani government in corona control program.
Epidemiology and prevention of water borne diseases:	2	 Enumerate common water borne diseases Explain the epidemiology and prevention measures of these diseases describe the current situation of these diseases on Pakistan and worldwide

poisoning	
Amebiasis and	
Giardiasis	
 Brucellosis 	
 Leptospirosis 	
• Worm	
infestations	

Practical Work

Week 1 Practicals			
	Cell of	1.5	Identify Cells of inflammation in themicroscope
Pathology	inflammation		
	Acute Appendicitis	1.5	Identify the histopathological changes
			in acute appendicitis
Forensic	Gastric Lavage	1.5	Demonstrate the steps of gastriclavage
Medicine			
Week 2 Practicals			
	Chronic	1.5	-Identify the morphological changes occurring in chronic
	cholecystitis		cholecystitis
Pathology	Granuloma	1.5	- Identify the various cells and their arrangement in a
			granuloma
Week 3 Practicals			
Pathology	Granulation Tissue	1.5	-Identify the histological features of
			granulation tissue
Week 4 Practicals			
	Catalase test	1.5	-Perform and interpret the result of catalase test by tube and
			slide method
	Coagulase test		-Perform and interpret the result of
			coagulase test by tube method

Pathology	Oxidase test		-Perform and interpret the result of coagulase test
3,	Culture media		-Identify blood agar, Mannitol saltagar, Chocolate media, Cary
			Blair transport media in the lab
			-Identify different types of haemolysison blood agar
Pharmacology			Prescription Writing
	Acute	1.5	Construct a prescription for a patient
	Tonsillitis		with acute tonsillitis.
	Sex determination	1.5	Identify human sex through bones
Forensic	through bones		
Medicine	Hair, Fibre		Identify human hair throughmicroscopy
			Differentiate between hair and fibre
Week 5 Practicals	5		
			Prescription Writing
Pharmacology	Malaria	1.5	Construct a prescription for a patient
			with Malaria
Week 6 Practicals	5		
	Hydatid Cyst	1.5	Identify cysts and ova of
			Echinococcus in the lab
	Leishmania		Identify leishmania in slides of bonemarrow/ skin biopsies
Pathology	Malaria		Identify Malarial parasite trophozoites and gametocytes under microscope

	Taenia		Identify ova of Taenia in the lab
	saginata/solium		
Community medicine	Communicable	3	Identify the models related to the communicable diseases
	diseases models		
			Explain the complication, preventive measures and the
			identification signs of concerned disease

CLINICAL SUBJECTS							
S#	MEDCINE	SURGERY	PAEDS	Obs/Gyn	ENT	EYE	PRIME
1	PUO 1	Surgical infections 1	PUO (better to teach either by Medicine or Paeds if majority content is same/ joint session can be taken) 1	Puerperal pyrexia 1	Acute & chronic Phyrangitis 1	Acute and chronic dacrocystit is	Reactionto illness 1
2		Anesthesia & pain relief	Child with Rash 1	Post- operative wound sepsis 1	Acute & chronic Rhinitis 1	Episcleritis 1	Attributes of professionalismempathy
3		Acute abdomen 1			Acute & chronic Sinusitis 2	Infective conjuncti Vitis 1	Steps of research process 1
4					Acute and chronic tonsillitis		Identifying study question 2
							Literature review 2

Learning Resources

S.No	Subjects	Textbooks		
1.	Community	1.Community Medicine by Parikh		
	Medicine	2. Community Medicine by M Illyas		
		3. Basic Statistics for the Health Sciences by Jan W Kuzma		
2.	Forensic	1. Nasib R. Awan. Principles and practice of Forensic Medicine 1st ed. 2002.		
	Medicine	2. Parikh, C.K. Parikh's Textbook of Medical Jurisprudence, Forensic Medicine and Toxicology. 7th ed.2005.		
		3.Knight B. Simpson's Forensic Medicine. 11th ed.1993.		
		4. Knight and Pekka. Principles of forensic medicine. 3rd ed. 2004		
		5. Krishan VIJ. Text book of forensic medicine and toxicology (principles and practice). 4th ed. 2007		
		6. Dikshit P.C. Text book of forensic medicine and toxicology. 1st ed. 2010		
		7. Polson. Polson's Essential of Forensic Medicine. 4th edition. 2010.		
		8. Rao. Atlas of Forensic Medicine (latest edition).		
		9. Rao.Practical Forensic Medicine 3rd ed ,2007.		
		10. Knight: Jimpson's Forensic Medicine 10th 1991,11th ed.1993		
		11. Taylor's Principles and Practice of Medical Jurisprudence. 15th ed.1999		
3.	Pathology	1. Robbins & Cotran, Pathologic Basis of Disease, 9th edition.		
		2. Rapid Review Pathology, 4th edition by Edward F. Goljan MD		
4.	PHARMACOLOGY	1. Lippincott Illustrated Pharmacology		
		2. Basic and Clinical Pharmacology by Katzung		

Assessment Plan - 3rd Year MBBS

The year-3 will be assessed in 3 blocks

- 1) Block-1 (Foundation 2 and Infection and Inflammation modules) will be assessed in paper-G
- 2) Block-2 (Multisystem, blood and MSK modules) will be assessed in paper-H
- 3) Block-3 (CVS and Respiratory module) will be assessed in paper-I
- 4) Each written paper consists of 120 MCQs and
- 5) Internal assessment will be added to final marks in KMU as shown in below table.
- 6) In OSPE, each station will be allotted 6 marks, and a total of 120 (+10% marks of internal assessment) marks are allocated for each OSPE/OSCE examination.

Year 3 Professional Exam in System-based Curriculum

Theory paper	Modules	Theory marks	Internal assessment theory (10%)	OSPE/OSPE	Internal assessment OSPE/OSPE (10%)	TOTAL MARKS
Paper G	Foundation-II Inf.&Inflamm.	120	14	120	14	268
Paper H	Multisystem Blood MSK-II	120	13	120	14	267
Paper I	CVS-II Respiratory-II	120	13	120	12	265
TOTAL MARKS		360	40	360	40	800

^{*}Research viva of 20 marks will be conducted in paper-L. However, the rest of 15 marks will be decided by the concerned department internally for the contribution of the students in research project/thesis.

Assessment Blueprints

Table 2: Paper G (Foundation II and Infection & Inflammation)

Subjects	Total MCQs
Infection & Inflammation	54
Foundation - II	66
Total	120

Table 3: Paper G OSCEs

Subject	Total OSCE stations
Infection & Inflammation	10
Foundation - II	10
Total	20

A minimum of 20 stations will be used in final exams. Total marks will be 120 (6 marks for each station).