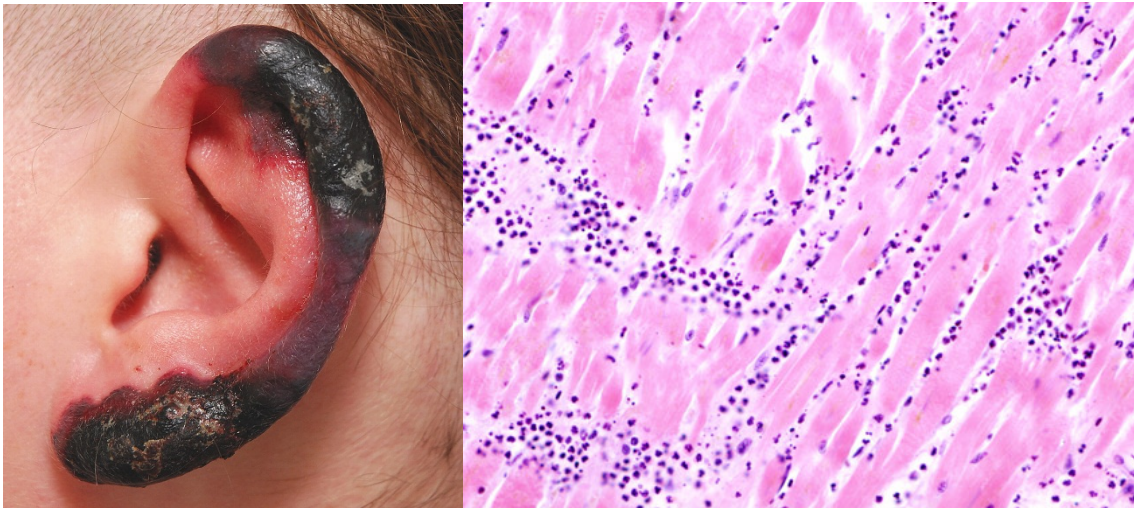


AJK Medical College, Muzaffarabad



Cell Pathology, General & Autonomic Pharmacology Module CPGAP-0202 3rd Year



Duration: 4-Weeks
Starting on:

**DEPARTMENT OF MEDICAL EDUCATION
AJK Medical College, Muzaffarabad**

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Module Development Team

Name	Role
Prof. Sarosh Majid Salaria	Principal/Patron in Chief/Dean
Prof. Muhammad Arif	Module Planner
Prof. Muhammad Munir	Coordinator
Dr. Ziyad Afzal Kayani	DME/Member
Dr. Javed Akhtar Rathore	Member
Dr. Nasir Ahmed Sheikh	Member
Dr. Malik Mehmood	Member
Dr. Uzma Hafeez	Member

Caution

80% mandatory attendance to appear in Modular / Professional / University Examination as per Pakistan Medical and Dental Council (PM&DC) regulation.

Rationale

For the better understanding of various diseases of human beings, it is vital to know the etiology, pathogenesis, morphology and clinical features of lesions. This aspect has got paramount importance for the effective management of health care issues of human being. The basic idea of this module is to familiarize the undergraduate medical students (from the beginning of their training) to the fundamental components of human diseases. The understanding, diagnosis and management of diseases is impossible without the functional cellular and molecular knowledge of human tissue.

Clinical assessment of human diseases is very challenging at times. The basic concepts of pathological aspects of diseases will help in understanding of disease pattern. Similarly basic pharmacological principles will help in formulating treatment plan of human diseases. Medico legal aspects are also important in human diseases.

Module Organization

In second Spiral / 3rd year, we are beginning with basic concepts of pathological aspects of diseases, pharmacological features and medico legal aspects of human lesions. This module integrates horizontally the disciplines of pathology, pharmacology and forensic medicine and vertically integrates with medicine, surgery, Paeds, Eye, ENT, Gynecology and Obstetrics.

This module will run for 4 weeks and comprises 4 Themes and two PBLs (each based on real life clinical problems). Each theme has its explicit outcomes/LOs. In order to enhance the quality of integration of basic science concepts with clinical sciences and to facilitate problem solving skills, this module has been designed and will be delivered by a committed integrated team comprising of pathologist, pharmacologist, forensic experts, Physicians and surgeons.

This module structured in a way that you will attain your objectives by pursuing different clinical themes while following a presented schedule/ time table. Based on these themes will be clinical cases; that you'll discuss in different sessions (SGDs, PBL, SDL, LGIS, Skill Lab) during the program. The time table/ schedule with clinical themes are included in this study guide.

Teaching Strategies

The content of this module will be delivered by a combination of different teaching strategies. These include small group discussions (SGD), large group interactive sessions (LGIS), demonstrations in clinics, lab practical, general club, dissection/ skill videos, clinical skill sessions at skill lab and case discussion in ward rounds and OPD. Entire curriculum will be delivered by clinical case scenarios each covering a theme. Read the cases and the objectives of the theme which you are supposed to encounter next day, understand and explain the case to yourself and read the relevant information. Following learning/teaching strategies will be used in this Module:

Small Group Discussion: Main bulk of the course content will be delivered in small group sessions. Each theme has an associated case. The case will be the centre around which learning will take place. Depending on the case you might be required to deduce

objectives and learning issues or only learning issues. Every group will have a facilitator assigned to it. The facilitator will be there to keep you on track, giving you maximum liberty to discuss and achieve the objectives as a group. Small groups will be followed by a wrap up session to standardize learning. Rest of the information will be in the schedule/ time table. There are no secrets regarding how to make the best of small group sessions. Come prepared for your session and make use of the combined wisdom of the group to achieve your learning objectives. The facilitator is your guide to help achieve them and clarify any misconceptions. Feel free to consult me in person or by e-mail, in case of any problems.

Large Group Interactive Sessions: LGIS will be employed at times to augment small groups. By and large they will be used to pass on general concepts regarding the theme. Large group instruction will be employed at times sparingly. Attend large group sessions with the following focus:

- a. Diagnose learning misconceptions.
- b. Clarify misconceptions.
- c. Standardize learning of all learners.
- d. Scaffolding of important concepts.
- e. Learn concepts not well understood in SGDs.
- f. Measure your learning comprehension.

Hands-on Activities/ Practical: This will be in the form of laboratory practical, spotting on slides, images, radiology and clinical skills. Attend your scheduled lab and take advantage of free time for study. Use your labs to correlate text structures to actual specimens in lab practice.

Videos: Videos demonstration of clinical skills will be given in Skills lab. Video demonstrations on history taking and clinical examination / invasive procedures, will be shown to give you an idea into the disease process, clinical testing and practical aspect of communication with the patients.

Self Directed Learning: A task will be given in SDL regarding the theme to be discussed on the next day. This will help to prepare you a bit before the theme is under discussion. A few SDLs (10%) have been added in between to create an environment for you to search literature as well as to deduce and synthesize information from different sources to meet the learning objectives. It will also help in breaking the monotonous / strenuous schedule and make you life- long learner.

Assessment

A full-fledged summative assessment will be conducted at the end of module. This will give you an idea about the format of the examination that you will go through at the end of the year. Of course, this will be followed by feedback on your performance in the exam. Marks obtained in the module examination will contribute 30% (internal assessment) towards end of year Professional University Examination. **There is no RESIT exam for module written assessment and block IPE** under any circumstances. If you miss them, your internal assessment will be recorded as zero. No excuse of any kind is permissible for absence in module or IPE assessment.

Table of Specification

S.NO	THEME	WEIGHTAGE
1	Energy Crisis	15%
2	Murder She Wrote	15%
3	Invaders	20%
4	Defenders	25%
5	Palpitations	25%

Themes & Learning Outcomes

1. ENERGY CRISIS

At the end of the theme insha Allah, the students will be able to;

- 1) Revisit Cell! Describe Cell's personality, Shape, Fashion and Functions
- 2) Draw the light and electronmicroscopic picture of a typical cell labeling all its components and organelles
- 3) Enumerate the function of each cellular organelle
- 4) Enlighten the role of ischemia causing cell injury
- 5) Draw the cytoplasmic changes in ischemic injury
- 6) Define the roles of ATP depletion, mitochondria, Cytoplasmic calcium, sodium potassium pump, endoplasmic reticulum, lysosomes, phagosomes and nucleus in cell injury
- 7) Define reversible Cell injury giving its types and mechanisms in;
 - a. Hydropic degeneration
 - b. Fatty change
- 8) Define irreversible cell injury. Enumerate its types
 - a. Coagulative necrosis
 - b. Caseation necrosis
 - c. Liquifactive necrosis
 - d. Gangrene, give its types and causes
- 9) Define pathological classification, describe its types and manifestations
- 10) Define aging and its accompanying features
- 11) Explain the role of flawed Nutrition in causing cell injury
- 12) Define the role of Radiation in causing cell injury
- 13) Explain the role of life style and socio ethical factors in causing cell injury
- 14) Describe the role of environment in causing cell injury
- 15) Enumerate Free Radicals and explain their role in causing cell injury
- 16) Explain the mechanism and types of cell death
- 17) Differentiate between necrosis and apoptosis
- 18) Enumerate adaptive responses to physiological and pathological stresses including;
 - a. Atrophy
 - b. Hyperplasia
 - c. Hypertrophy
 - d. Metaplasia
- 19) Enumerate abnormal intracellular accumulations; give examples along with mechanism and manifestations in
 - a. Glycogen storage disease
 - b. Lipid storage diseases

- c. Congenital hyperlipidemia
 - d. Lipofuchsin
- 20) Enumerate causes of damage to the blood cells
- a. Explain the role of cell membrane damage in Erythrocytes
 - b. Explain defects in hemoglobin and enzymes leading to Erythrocytes injury
 - c. Explain how physical trauma causes damage to erythrocytes
 - d. Explain immunological mechanism of cell damage
 - e. Explain genetic abnormalities adversely affecting bactericidal ability of White Blood Cells and macrophages
 - f. Enumerate causes of platelet and megakaryocyte injury
 - g. Explain how parasites damage erythrocytes
- 21) Correlate cell injury with cellular components released in the blood
- a. Define biomarker, Enlist characteristics of good biomarker
 - b. Enumerate different Biomarkers of liver cell injury
 - c. Enumerate different Biomarkers of cardiac cell injury
 - d. Enumerate different Biomarkers of renal cells injury
 - e. State the difference between different biomarkers in terms of;
 - a. Those represent functioning of the cell
 - b. Those represent cell injury

2. MURDER SHE WROTE

At the end of the THEME the student should be able to:

- 1) Define Inquest
- 2) discuss different medico legal system of death investigation in the world
- 3) discuss criminal procedure code regarding postmortem examination in Pakistan
- 4) Define Postmortem examination
- 5) Enlist types of postmortem examinations
- 6) Give objectives of postmortem examination
- 7) discuss prerequisites of postmortem examination
- 8) Enlist different postmortem techniques
- 9) Define postmortem protocol
- 10) Enlist different types of postmortem protocol
- 11) Discuss procedure of postmortem examination
- 12) Visualization of postmortem procedure on multimedia
- 13) Define negative / obscure autopsy
- 14) Enlist causes of negative autopsy
- 15) Define exhumation
- 16) Give objectives of exhumation
- 17) Enlist precautions of exhumation
- 18) Explain the procedure of exhumation and give its limitations
- 19) Correlate cell injury with intracellular accumulations
- 20) Explain cellular responses to stress and noxious stimuli.
- 21) Differentiate between metastatic and dystrophic calcification
- 22) Enumerate and classify various causative agents of infections in humans.
- 23) Differentiate between Prokaryotic & Eukaryotic Cells.
- 24) Classify the medically important bacteria.

3. INVADERS

At the end of the THEME the student should be able to:

- 1) Describe and explain various mechanisms of cell injury by different types of microbial agents including bacteria, mycobacteria, fungi, parasites, viruses and prion

- 2) Classify the medically important bacteria
- 3) Illustrate structure of bacterial cell membrane and enumerate its functions
- 4) Draw and label Gram positive and Gram negative cell wall
- 5) Enlist the cytoplasmic structures of prokaryotes along with their functions
- 6) Define hospital infection control and role of Infection Control Committee in containing infections
- 7) Define SIRS, Sepsis, severe sepsis, and septic shock, discuss pathogenesis of septic shock
- 8) Enlist and explain mechanism of antibiotics resistance
- 9) Perform the Gram Stain. Identify/ draw Gram Positive Cocci and Gram Negative Bacilli.
- 10) Perform correct technique for Hand hygiene
- 11) Define normal flora and enlist microorganisms of normal flora of skin, nasopharynx, gastrointestinal tract and vaginal flora.
- 12) Explain various routes of transmission of microbes. Illustrate the chain of infection. Describe the precautions to prevent the transmission of infections
- 13) Enlist and explain the various diagnostic techniques for infectious agents.
- 14) Define hospital acquired infections. Enlist Common Hospital acquired infections, Name the most Common organisms causing HAIs, and describe the types, steps and techniques of Hand hygiene
- 15) Describe the cell structure of fungi, classify fungal infections, and describe the various laboratory methods for diagnosis of fungal infections
- 16) Describe structure of viruses, classify viruses, and describe the various laboratory methods for diagnosis of viral infections

4. DEFENDERS

At the end of the THEME the student should be able to:

- 1) Define the term Pharmacology.
- 2) Define Different Branches of Pharmacology.
- 3) Define Pharmacopoeias with examples.
- 4) What is drug nomenclature? Illustrate with one example.
- 5) Classify the different sources of drugs with examples.
- 6) Define the active principle and what are their types?
- 7) List the names of active principles
- 8) Define different types of active principles with examples of drugs.
- 9) Give the formulas for calculating the dose of a drug according to age and weight
- 10) Define the different dosage forms
- 11) What are the advantages & disadvantages of the different routes of administration?
- 12) What are different means by which drug transport across cell membrane? Illustrate with examples.
- 13) Define the term Drug Absorption
- 14) How different factors affecting absorption of drugs?
- 15) Define Bioavailability. How it is calculated? What is its clinical importance?
- 16) How different factors affecting Bioavailability?
- 17) Define First- pass metabolism and What is its significance?
- 18) Define the terms Bioequivalence and therapeutic equivalence
- 19) Define drug distribution
- 20) How different factors affecting distribution of a drug?
- 21) Write the distribution of a drug through various body compartments
- 22) Define volume of distribution
- 23) Give the importance of distribution and the volume of distribution
- 24) What are the characteristics/features of a drug that is bound to plasma proteins? What is its clinical significance? Illustrate with examples.
- 25) Define Biotransformation.
- 26) Enlist sites of biotransformation
- 27) What are the aims and outcomes of Biotransformation?

- 28) Define phase I and phase II biotransformation reactions and their types
- 29) Enlist examples of drugs undergoing Phase 1 and phase 2 metabolic reactions
- 30) What are the characteristics of Phase 1 and Phase 2 biotransformation reactions?
- 31) What are microsomal and non- microsomal biotransformation reactions?
- 32) How different factors affecting biotransformation of a Drug?
- 33) Define enzyme induction and its significance with examples?
- 34) Define enzyme inhibition and its significance with examples?
- 35) Define plasma half life
- 36) Enumerate and explain the factors affecting half-life?
- 37) What is the importance of plasma half life
- 38) Define steady state concentration and give its significance
- 39) Define the terms First order kinetics and Zero Order Kinetics with examples
- 40) What are differences between First order kinetics and Zero Order Kinetics?
- 41) Define Maintenance Dose & loading dose, how these are calculated and what is their clinical significance.
- 42) Define the drug excretion and clearance.
- 43) Enlist various routes through which drugs are excreted from the body with examples
- 44) Explain the basic principles by which drugs are excreted from the kidneys?
- 45) How different factors affecting excretion of drugs?
- 46) What are different mechanisms by which drugs produce their action?
- 47) What are different types of mechanisms by which drugs act through receptors, with their examples?
- 48) What are different types of G-Proteins with examples of drugs?
- 49) Write the role of second messengers involved in receptors transduction
- 50) Differentiate between Receptor Down regulation and up regulation with examples. What is their significance?
- 51) Differentiate between the terms potency and efficacy
- 52) What are the characteristics of a Graded Dose Response Curve?
- 53) What information can be obtained from a Graded Dose Response curve?
- 54) What are the characteristics of Quantal Dose Response Curve?
- 55) Define ED_{50} , LD_{50}
- 56) Define Therapeutic Index and Therapeutic Window with examples. What are their clinical significance?
- 57) What are the differences between Graded and Quantal Dose Response curve
- 58) How different factors can affect the actions of different drugs?
- 59) Define with examples the term Agonist, Partial Agonist, Antagonist and Inverse Agonist
- 60) Define Spare Receptors. What is their importance?
- 61) Write different types of Antagonism with examples and clinical significance.
- 62) Define and explain the characteristics of competitive and non-competitive antagonism?
- 63) Enumerate different types of adverse drug reactions? Differentiate with examples.
- 64) Define drug dependence, addiction, abuse.
- 65) Differentiate between psychological dependence and physical dependence with examples
- 66) Define Tolerance. Differentiate different types of Tolerance with their mechanisms and examples.
- 67) Explain difference between Pharmacokinetic and Pharmacodynamic Tolerance
- 68) Define Tachyphylaxis, it's mechanisms and examples.
- 69) Differentiate between Tolerance and Tachyphylaxis
- 70) Define Pharmacogenetics and Pharmacogenomics. What are their importance? Illustrate with examples.
- 71) Define idiosyncrasy with examples.
- 72) Define Drug interactions
- 73) Differentiate between various types with examples?
- 74) Give the different types of Drug Interactions

75) Define Synergism, Summation (addition) and Potentiation with examples.

5. Palpitations

At the end of theme student should be able to:

- 1) Revision of anatomy & physiology of ANS?
- 2) Discuss autonomic receptors and explain their role in functional organization of autonomic activity?
- 3) Explain how different drugs act to modify the autonomic function?
- 4) Discuss the clinical importance of autonomic pharmacology?
- 5) Classify Cholinomimetics and anticholinergic drugs.
- 6) Enlist subtypes & characteristic of cholinergic receptors?
- 7) Enumerate various direct acting cholinomimetics and write their pharmacokinetics properties
- 8) Discuss the mechanism of Action, Pharmacological effects and uses of direct acting cholinergic drugs?
- 9) Enlist the adverse effects, precautions/contraindications and drug interactions of direct acting cholinomimetics.
- 10) Classify indirectly acting cholinomimetics and give their pharmacokinetic properties
- 11) Discuss the mechanism of action, pharmacological effects & therapeutic uses of anti- cholinesterases.
- 12) Discuss the adverse effects, precautions/contraindications & drug interactions of cholinesterase inhibitors?
- 13) Discuss the role of reversible anti-cholinesterases in treatment of Alzheimer's disease
- 14) Enlist the features of overdose toxicity of organophosphates and their management
- 15) Classify Anticholinergic drugs and differentiate between antimuscarinic & anti nicotinic drugs.
- 16) Write down the pharmacokinetic Properties of Natural Belladonna Alkaloids.
- 17) Give mechanism of action, Pharmacological effects & uses of Anti-muscarinics.
- 18) Discuss the adverse effects, precautions /contraindications & drug interactions of antimuscarinic drugs?
- 19) Enlist the features of overdose toxicity of antimuscarinics and how their management
- 20) Discuss the role of Ganglion-blocker drugs in modern pharmacology
- 21) Classify Adrenomimetic drugs.
- 22) Enlist types & subtypes of Adrenoceptors?
- 23) Discuss the pharmacokinetic properties of Adrenomimetic drugs.
- 24) Describe the mechanism of action and pharmacological effects of Adrenomimetics.
- 25) Discuss the therapeutic uses, adverse effects precautions/ contraindications & drugs interactions of Adrenomimetics
- 26) Give classification of Antiadrenergics
- 27) Discuss the pharmacokinetics of alpha – adrenoceptor antagonists
- 28) Write down the mechanism of action, pharmacological effects and clinical uses of alpha- adrenoceptor antagonists.
- 29) Enlist the adverse effect, precautions/contraindications & drug interactions of alpha adrenoceptor antagonists.
- 30) Discuss the logic of using long acting alpha adrenoceptor antagonists in the treatment of pheochromocytoma & benign prostatic Hyperplasia
- 31) Enlist the pharmacokinetic properties of Beta- adrenoceptor antagonists
- 32) Describe the mechanism of action, pharmacological effects & clinical applications of Beta blockers.
- 33) Write down the adverse effects, precautions/contraindications & drug interactions of beta – adrenoceptor antagonists.
- 34) Enlist the drugs used in treatment of Glaucoma along with their mechanism of actions.
- 35) Write down the clinical importance of adrenergic neuron blockers in modern day therapy.

PBL-1

68-year-old man presents with a complaint of light-headedness on standing that is worse after meals and in hot environments. Symptoms started about 4 years ago and have slowly progressed to the point that he is disabled. He has fainted several times, but always recovers consciousness almost as soon as he falls. Review of symptoms reveals slight worsening of constipation, urinary retention out of proportion to prostate size, and decreased sweating. He is otherwise healthy with no history of hypertension, diabetes, or Parkinson's disease. Because of his urinary retention, he was placed on the α 1 antagonist tamsulosin but he could not tolerate it because of worsening of orthostatic hypotension. Physical examination revealed a blood pressure of 167/84 mm Hg supine and 106/55 mm Hg standing. There was an inadequate compensatory increase in heart rate (from 84 to 88 bpm), considering the degree of orthostatic hypotension. Physical examination is otherwise unremarkable with no evidence of peripheral neuropathy or parkinsonian features. Laboratory examinations are negative except for plasma norepinephrine, which is low at 98 pg/mL (normal is 250–400 pg/mL for his age).

PBL-2

A 55 year old man resident of Swat was removing ice from his car screen when he became quite uncomfortable and began excessively sweating. He felt quite dizzy. He was taken to the hospital where his pulse was 120 beats per minute and blood pressure was 90/56. On examination he had tenderness in liver area. His random blood glucose was 275 mg/dl. He had a history of smoking 2 packs per day, naswar and alcohol.

PBL-3

A 26-year-old man is brought by friends to the emergency department of the city hospital because he has been behaving strangely for several days. A known user of methamphetamine, he has not eaten or slept in 48 hours. He threatened to shoot one of his friends because he believes this friend is plotting against him. On admission, the man is extremely agitated, appears to be underweight, and is unable to give a coherent history. He has to be restrained to prevent him from walking out of the emergency department and into traffic on the street. His blood pressure is 160/100 mm Hg, heart rate 100, temperature 39°C, and respirations 30/ min. His arms show evidence of numerous intravenous injections. The remainder of his physical examination is unremarkable. After evaluation, the man is given a sedative, fluids, a diuretic, and ammonium chloride parenterally.

Recommended Books

1. Basic Pathology by Robbins 9th edition
2. Pathologic Basis of Disease 8th edition
3. Jawetz Medical Microbiology 25th edition
4. Warren Levinson Microbiology
5. Pathology illustrated 25th edition
6. Walter & Israel General pathology 7th edition
7. Katzung's Basic and clinical pharmacology 12th edition
8. Rang and Dale Pharmacology, 7th Edition
9. Parikhs text book of medical jurisprudence, forensic medicine and toxicology 6th edition
10. Text book of forensic medicine and toxicology 2nd edition
11. Ansari Public health and community medicine 7th edition
12. Epidemiology and biostatistics by Richard Hebel 7th edition.
13. Tietz Fundamentals of Clinical Chemistry
14. Kaplan Book of Clinical Chemistry

CAUTION!
Eighty percent (80%) attendance is mandatory to appear in Module/Professional/ University Examination as per Pakistan Medical and Dental Council (PMDC) regulations.

AJK Medical College, Muzaffarabad

CPGAP Module – (3rd Year)

Week-1

DATE→			WEEK 1		
TIME↓	MONDAY	TUESDAY	WEDNESDAY	THURSDAY	FRIDAY
8:00am-09:00am	LGIS Introduction to CPGAP Module Prof. Anwar & Module Team	LGIS Red Cell Injury Dr. Malik Mahmood	SGD Pharmacokinetics-II Dr. Arif & Team-4	LGIS Cellular Adaptation & Intracellular Accumulation Dr. Sarosh Majid	SGD Postmortem Examination-I Dr. Naseer & Team
9:00am - 10:00am	CLINICAL ROTATION	LGIS Microbial Cell Injury Dr. Muhammad Munir	Wrap-up Dr. Arif/Dr. Inayat	CLINICAL ROTATION	BREAK
10:00am-10:30am		BREAK			
10:30am-11:30am		SGD Pharmacokinetics-I Dr. Arif & Team-4	SDL Pharmacokinetics		SGD Pharmacokinetics-VI Dr. Arif/Dr. Inayat
11:30am-12:30pm		Wrap-up Dr. Arif/Dr. Inayat	SGD Pharmacokinetics-III Dr. Arif & Team-4		
12:30pm–1:30pm	LGIS Introduction to Forensic Medicine Dr. Humayun/ Dr. Naseer	SGD Reversible Cell Injury Dr. Anwar/ Dr. Wafa	Wrap-up Dr. Arif/Dr. Inayat	LGIS Pharmacokinetics-IV Dr. Arif/Dr. Inayat	SDL
1:30pm–2:00pm	LUNCH BREAK				
2:00pm-3:00pm	LGIS Introduction to Pharmacology Dr. Arif/ Dr. Inayat	PBL-1A Dr. Arif & Team-4	LGIS Irreversible Cell Injury Dr. Sarosh Majid	SGD Pharmacokinetics-V Dr. Arif & Team-4	SDL
3:00pm-4:00pm	LGIS Meet the Healthy Cell Dr. Anwar	LGIS Medicolegal System of Death Investigation Dr. Humayun/ Dr. Naseer	LGIS Bacterial Cell Culture Dr. Muhammad Munir	Wrap-up Dr. Arif/Dr. Inayat	

AJK Medical College, Muzaffarabad

CPGAP Module – (3rd Year)

Week-2

DATE→					
TIME↓	MONDAY	TUESDAY	WEDNESDAY	THURSDAY	FRIDAY
8:00am-09:00am	LGIS Diagnostic Techniques for Infections Agents Dr. Munir	LGIS Normal Flora Dr. Muntaz Khan	LGIS Biomarkers of Cell Injury Dr. Sobia Irum	LGIS Pharmacodynamics-VII Dr.Arif/Dr. Inayat	PBL-1B Dr. Arif & Team-3
9:00am - 10:00am	CLINICAL ROTATION	LGIS Pharmacodynamics-II Dr. Arif/Dr. Inayat	LGIS Pharmacodynamics-V Dr. Arif/Dr. Inayat	CLINICAL ROTATION	
10:00am-10:30am		BREAK			BREAK
10:30am-11:30am		SGD Pharmacodynamics-III Dr.Arif & Team-4	LGIS Exhumation Dr. Humayun/ Dr. Arif		SGD Pharmacodynamics-VIII Dr.Arif& Team-4
11:30am-12:30pm			SGD Pharmacodynamics-VI Dr.Arif & Team-4		
12:30pm–1:30pm	LGIS Postmortem Examination-II Dr. Naseer Sheikh	LGIS Postmortem Examination-III Dr. Naseer Sheikh	WRAP-UP Dr. Arif	LGIS Dr. Humayun/ Dr. Naseer	SDL
1:30pm–2:00pm	LUNCH BREAK				
2:00pm-3:00pm	SGD Pharmacodynamics-I Dr.Arif & Team-4	LGIS Pharmacodynamics-IV Dr.Arif/Dr. Inayat	PBL-2A Dr. Anwar & Team-3	PRACTICAL Gram Stain Dr. Muntaz & Team-3	SDL
3:00pm-4:00pm	WRAP-UP Dr. Arif	SDL	SDL		

AJK Medical College, Muzaffarabad

CPGAP Module – (3rd Year)

Week-3

DATE→					
TIME↓	MONDAY	TUESDAY	WEDNESDAY	THURSDAY	FRIDAY
8:00am-09:00am	LGIS Drug Clearance in Hepatic & Renal Failure Prof. Arif	LGIS Cholinergics-III Prof. Arif / Dr. Inayat	SGD Anticholinergic -II Prof. Arif & Team-4	LGIS Classification of Fungi Prof. Anwar ul Haq	LGIS Management of Myasthenia, Gravis, Alzheimer Disease & Parkinsonism-I Dr. Liaquat Awan
9:00am - 10:00am	CLINICAL ROTATION	LGIS Prevention of Infectious Disease Dr. Ahmed Khan	Wrap-up Prof. Arif / Dr. Inayat		PBL-3A Prof. Arif & Team-4
10:00am-10:30am		BREAK			BREAK
10:30am-11:30am		SGD Cholinergic-IV Prof. Arif & Team-4	LGIS Transmission of Microbes Dr. Mumtaz Khan	CLINICAL ROTATION	SGD Anticholinergic-IV Prof. Arif & Team-4
11:30am-12:30pm		Wrap-up Prof. Arif/ Dr. Inayat	PBL-2B		Wrap-up Prof. Arif / Dr. Inayat
12:30pm–1:30pm	LGIS Cholinergics-I Prof. Arif/ Dr. Inayat	LGIS Antibiotic Resistance Dr. Muhammad Munir	Dr. Anwar & Team-3	LGIS Anticholinergic-III Dr. Arif/ Dr. Inayat	SDL
1:30pm–2:00pm	LUNCH BREAK				
2:00pm-3:00pm	LGIS Bacterial Cell Structure & it's Classification Prof. Munir	LGIS Anticholinergic-I Prof. Arif/ Dr. Inayat	LGIS Role of NMBs in Anesthesia + Management of Organophosphate Poisoning Dr. Ahsan Ul Haque/ Dr. Javed Rathore	PRACTICAL Reversible Cell Injury & Adaptation Dr. Sarosh Majid & Team-3	SDL
3:00pm-4:00pm	LGIS Cholinergics-II Prof. Arif / Dr. Inayat	SDL	SDL		

AJK Medical College, Muzaffarabad

CPGAP Module – Class of 2013-18 (3rd Year)

Week-4

DATE→					
TIME↓	MONDAY	TUESDAY	WEDNESDAY	THURSDAY	FRIDAY
8:00am-09:00am	LGIS Adrenergic-I Dr. Arif/ Dr. Inayat	LGIS Management of Glaucoma and Role of Decongestant in Eye Dr. Bushra	LGIS Management of Overactive Bladder, Stress Incontinence and BPH-I Dr. Mohsin shakeel	LGIS Adrenergic-III Dr. Arif/ Dr. Inayat	SGD Anti-Adrenergic-III Dr. Arif & Team-4 <u>Wrap-up</u> Dr. Arif/ Dr. Inayat
9:00am - 10:00am	CLINICAL ROTATION	LGIS Management of ADHD-I Dr. Ayesha	LGIS Management of Acute & Chronic Heart Failure-I Dr. Rizwan Abid	CLINICAL ROTATION	
10:00am- 10:30am		Role of Decongestants in ENT Dr. Farooq	Asthma-I Dr. Ashfaq Ahmed		
10:30am- 11:30am		BREAK			SGD Anti-Adrenergic-IV Dr. Arif & Team-4 <u>Wrap-up</u> Dr. Arif/ Dr. Inayat
11:30am- 12:30pm		LGIS Classification of Fungi and Transmission of Microbes Dr. Mumtaz Khan	SGD Adrenergic-II Dr. Arif & Team-4 <u>Wrap-up</u> Dr. Arif/ Dr. Inayat		
12:30pm– 1:30pm	LGIS Disinfection & Sterilization Dr. Mumtaz Khan	PBL-3B Dr. Arif & Team-4	LGIS Bacterial Cell Structure & it’s Classification Dr. Mumtaz Khan	LGIS Pheochromocytoma and its Management Dr. Ziyad Afzal Kayani	SDL
1:30pm– 2:00pm	LUNCH BREAK				
2:00pm- 3:00pm	PRACTICAL Necrosis & Apoptosis	PRACTICAL Reversible Cell Injury & Adaptation	LGIS Anti-Adrenergic-II Dr. Arif/ Dr. Inayat	SGD Adrenergic-IV Dr. Arif & Team-4	SDL
3:00pm- 4:00pm	Dr. Anwar & Team- 3	Dr. Sarosh Majid & Team-3	SDL	<u>Wrap-up</u> Dr. Arif/ Dr. Inayat	



For Inquiries & Trouble Shooting please contact:
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