

MGM INSTITUTE OF HEALTH SCIENCES

(Deemed to be University u/s 3 of UGC Act, 1956) Grade 'A' Accredited by NAAC Sector-01, Kamothe, Navi Mumbai -410 209 Tel 022-27432471, 022-27432994, Fax 022 -27431094

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Amended History

- 1. Approved as per BOM 04/2007, Item No. 4, Dated 14/12/2007.
- 2. Amended upto BOM 23/2012, resolution No. 4, dated 30/03/2012.
- 3. Amended upto BOM 40/2015, Resolution No. 3.2(c), Resolution No. 3.2(d), Resolution No. 3.2(f), Dated 13/03/2015.
- 4. Amended upto BOM-43/2015, Resolution No. 3.2 (a), Dated 06/11/2015.
- 5. Amended upto BOM 45/2016, Resolution No. 3.2(b), Dated 28/04/2016.
- Amended upto BOM 51/2017, Resolution No. 1.3.7.1, Resolution No. 1.3.7.5, Resolution No. 1.3.8.1, Resolution No. 1.3.8.8, Resolution No. 1.3.8.11, Resolution No. 1.3.8.13, dated 28/08/2017.
- 7. Amended upto BOM 52/2018, Resolution No. 3.5.9; Dated 13/01/2018.
- 8. Amended upto BOM 53/2018, Resolution No. 4.2.1, Resolution No. 4.3.5 Dated 19/05/2018.
- 9. Amended upto BOM 55/2018, Resolution No. 4.5.2.1, Resolution No. 4.5.2.2, Resolution No. 4.5.2.3, Resolution No. 4.13, Dated 27/11/2018.
- 10. Amended upto BOM 57/2019, Resolution No. 3.1.4.2, dated 26/04/2019.

GENERAL CONSIDERATIONS AND TEACHING APPROACH

- (1) Graduate medical curriculum is oriented towards training students to undertake the responsibilities of a physician of first contact who is capable of looking after the preventive, promotive, curative & rehabilitative aspect of medicine.
- (2) With wide range of career opportunities available today, a graduate has a wide choice of career opportunities. The training, though broad based and flexible should aim to provide an educational experience of the essentials required for health care in our country.

"Training should be able to meet internationally acceptable standards."

- (3) To undertake the responsibilities of service situations which is a changing condition and of various types, it is essential to provide adequate placement training tailored to the needs of such services as to enable the graduates to become effective instruments of implementation of those requirements. To avail of opportunities and be able to conduct professional requirements, the graduate shall endeavour to have acquired basic training in different aspects of medical care.
- (4) The importance of the community aspects of health care and of rural health care services is to be recognized. This aspect of education & training of graduates should be adequately recognized in the prescribed curriculum. Its importance has been systematically upgraded over the past years and adequate exposure to such experiences should be available throughout all the three phases of education & training. This has to be further emphasized and intensified by providing exposure to field practice areas and training during the internship period. The aim of the period of rural training during internship is to enable the fresh graduates to function efficiently under such settings.
- (5) The educational experience should emphasize health and community orientation instead of only disease and hospital orientation or being concentrated on curative aspects. As such all the basic concepts of modern scientific medical education are to be adequately dealt with.
- (6) There must be enough experiences to be provided for self learning. The methods and techniques that would ensure this must become a part of teaching learning process.
- (7) The medical graduate of modern scientific medicine shall endeavour to become capable of functioning independently in both urban and rural environment. He/she shall endeavour to give emphasis on fundamental aspects of the subjects taught and on common problems of health and disease avoiding unnecessary details of specialization.
- (8) The importance of social factors in relation to the problem of health and diseases should receive proper emphasis throughout the course and to achieve this purpose, the

educational process should also be community based than only hospital based. The importance of population control and family welfare planning should be emphasized throughout the period of training with the importance of health and development duly emphasized.

- (9) Adequate emphasis is to be placed on cultivating logical and scientific habits of thought, clarity of expression and independence of judgment, ability to collect and analyze information and to correlate them.
- (10) The educational process should be placed in a historic background as an evolving process and not merely as an acquisition of a large number of disjointed facts without a proper perspective. The history of Medicine with reference to the evolution of medical knowledge both in this country and the rest of the world should form a part of this process.
- (11) Lectures alone are generally not adequate as a method of training and are a poor means of transferring/acquiring information and even less effective at skill development and in generating the appropriate attitudes. Every effort should be made to encourage the use of active methods related to demonstration and on firsthand experience. Students will be encouraged to learn in small groups, through peer interactions so as to gain maximal experience through contacts with patients and the communities in which they live. While the curriculum objectives often refer to areas of knowledge or science, they are best taught in a setting of clinical relevance and hands on experience for students who assimilate and make this knowledge a part of their own working skills.
- (12) The graduate medical education in clinical subjects should be based primarily on outpatient teaching, emergency departments and within the community including peripheral health care institutions. The out-patient departments should be suitably planned to provide training to graduates in small groups.
- (13) Clinics should be organized in small groups of preferably not more than 10 students so that a teacher can give personal attention to each student with a view to improve his skill and competence in handling of the patients.
- (14) Proper records of the work should be maintained which will form the basis for the students' internal assessment and should be available to the inspectors at the time of inspection of the college by the Medical Council of India.
- (15) Maximal efforts have to be made to encourage integrated teaching between traditional subject areas using a problem based learning approach starting with clinical or community cases and exploring the relevance of various preclinical disciplines in both understanding and resolution of the problem. Every attempt be made to de-emphasize compartmentalization of disciplines so as to achieve both horizontal and vertical integration in different phases.

- (16) Every attempt is to be made to encourage students to participate in group discussions and seminars to enable them to develop personality, character, expression and other faculties which are necessary for a medical graduate to function either in solo practice or as a team leader when he begins his independent career. A discussion group should not have more than 20 students.
- (17) Faculty member should avail of modern educational technology while teaching the students and to attain this objective, Medical Education Units/ Departments be established in all medical colleges for faculty development and providing learning resource material to teachers.
- (18) To derive maximum advantage out of this revised curriculum, the vacation period to students in one calendar year should not exceed one month, during the 4 ½ years Bachelor of Medicine and Bachelor of Surgery (MBBS) Course.
- (19) In order to implement the revised curriculum in Toto, State Govts. and Institution Bodies must ensure that adequate financial and technical inputs are provided.
- (20) HISTORY OF MEDICINE –The students will be given an outline on "History of Medicine". This will be taught in an integrated manner by subject specialists and will be coordinated by the Medical Education Unit of the College.
- (21) All medical institutions should have curriculum committee which would plan curricula and instructional method which will be regularly updated.
- (22) Integration of ICT in learning process will be implemented.

OBJECTIVE OF MEDICAL GRADUATE TRAINING PROGRAMME:

- (1) **NATIONAL GOALS** : At the end of undergraduate program, the medical student should be able to :
- (a) Recognize `health for all' as a national goal and health right of all citizens and by undergoing training for medical profession fulfill his/her social obligations towards realization of this goal.
- (b) Learn every aspect of National policies on health and devote himself / herself to its practical implementation.
- (c) Achieve competence in practice of holistic medicine, encompassing promotive, preventive, curative and rehabilitative aspects of common diseases.
- (d) Develop scientific temper, acquire educational experience for proficiency in profession and promote healthy living.
- (e) Become exemplary citizen by observation of medical ethics and fulfilling social and professional obligations, so as to respond to national aspirations.
- (2) **INSTITUTIONAL GOALS:** (1) In consonance with the goals each medical institution should evolve institutional goals to define the manpower (or professionals) they intend to produce. The undergraduate students coming out of a medical institute should:
 - (a) Be competent in diagnosis and management of common health problems of the individual and the community, commensurate with his/her position as a member of the health team at the primary, secondary or tertiary levels, using his/her clinical skills based on history, physical examination and relevant investigations.
 - (b) Be competent to practice preventive, promotive, curative and rehabilitative medicine in respect to the commonly encountered health problems.
 - (c) Appreciate rationale for different therapeutic modalities; be familiar with the administration of the "essential drugs" and their common side effects.
 - (d) Be able to appreciate the socio-psychological, cultural, economic and environmental factors affecting health and develop humane attitude towards the patients in discharging one's professional responsibilities.
 - (e) Possess the attitude for continued self learning and to seek further expertise or to pursue research in any chosen area of medicine, action research and documentation skills.
 - (f) be familiar with the basic factors which are essential for the implementation of the National Health Programmes including practical aspects of the following:-
 - (i) Family Welfare and Material and Child Health(MCH)
 - (ii) Sanitation and water supply

- (iii) Prevention and control of communicable and non-communicable diseases (iv)
- Immunization (v)
- Health Education (vi)
- IPHS standard of health at various level of service delivery, medical waste disposal. (vii)
- Organizational institutional arrangements.
- Acquire basic management skills in the area of human resources, materials (g) and resource management related to health care delivery, General and hospital management, principal inventory skills and counseling (h)
- Be able to identify community health problems and learn to work to resolve these by designing, instituting corrective steps and evaluating outcome of such measures.
- Be able to work as a leading partner in health care teams and acquire (i) proficiency in communication skills. (j)
- Be competent to work in a variety of health care settings. (k)
 - Have personal characteristics and attitudes required for professional life such as personal integrity, sense of responsibility and dependability and ability to relate to or show concern for other individuals.

All efforts must be made to equip the medical graduate to acquire the skills as detailed as under:

A comprehensive list of skills recommended as desirable for Bachelor of Medicine and Bachelor of Surgery (MBBS) Graduate:

- 1. Clinical Evaluation:
 - To be able to take a proper and detailed history.
 - To perform a complete and thorough physical examination and elicit clinical signs. (a)
 - To be able to properly use the stethoscope, Blood Pressure, Apparatus Auroscope, Thermometer, Nasal Speculum, Tongue Depressor, Weighing Scales, Vaginal (b) (c)
 - To be able to perform internal examination-Per Rectum (PR), Per Vaginum (PV) etc.
 - To arrive at a proper provisional clinical diagnosis. (d) (e)
 - Bed side Diagnostic Tests: II.
 - To do and interpret Haemoglobin (HB), Total Count (TC), Erythrocytic Sedimentation Rate (ESR), Blood smear for parasites, Urine examination /albumin (a) /sugar /ketones /microscopic:
 - Stool exam for ova and cysts;
 - Gram, staining and Siehl-Nielsen staining for AFB; (b)
 - (c) To do skin smear for lepra bacilli
 - To do and examine a wet film vaginal smear for Trichomonas (d)
 - To do a skin scraping and Potassium Hydroxide (KOH) stain for fungus infections; (e)
 - (f)
 - To perform and read Montoux Test. (g)
 - Ability to Carry Out Procedures: III.
 - To conduct CPR (Cardiopulmonary resuscitation) and First aid in newborns, children (a)
 - To give Subcutaneous (SC) /Intramuscular (IM) /Intravenous (IV) injections and start (b)

- Intravenous (IV) infusions. To pass a Nasogastric tube and give gastric leavage.
- To administer oxygen-by masic/catheter (c)
- (d) To administer enema
- To pass a ruinary catheter-male and female (e)
- (f) To insert flatus tube
- To do pleural tap, Ascitic tap & lumbar puncture (g)
- Insert intercostal tube to relieve tension pneumothorax (h)
- (i) To control external Haemorrhage.
- (j)
- Anaesthetic Procedure IV
 - Administer local anaesthesia and nerve block (a)

Be able to secure airway potency, administer Oxygen by Ambu bag. (b) V

Surgical Procedures

- To apply splints, bandages and Plaster of Paris (POP) slabs; (a)
- To do incision and drainage of abscesses; (b)
- To perform the management and suturing of superficial wounds; (c)
- To carry on minor surgical procedures, e.g. excision of small cysts and nodules, (d)
 - circumcision, reduction of paraphimosis, debridement of wounds etc
- (e) To perform vasectomy;
- (f) To manage anal fissures and give injection for piles.
- VI Mechanical Procedures
 - To perform thorough antenatal examination and identify high risk pregnancies. (a)
 - (b) To conduct a normal delivery;
 - To apply low forceps and perform and suture episiotomies; (c)
 - (d) To insert and remove IUD's and to perform tubectomy

VII **Paediatrics**

- To assess new borns and recognize abnormalities and I.U. retardation (a)
- (b) To perform Immunization;
- (c) To teach infant feeding to mothers;
- To monitor growth by the use of 'road to health chart' and to recognize development (d) retardation;
- To assess dehydration and prepare and administer Oral Rehydration Therapy (ORT) (e)
- (f) To recognize ARI clinically;

ENT Procedures: VIII

- (a) To be able to remove foreign bodies;
- To perform nasal packing for epistaxis; (b)
- To perform trachesotomy (c)

IX **Ophthalmic Procedures:**

- (a) To invert eye-lids;
- To give Subconjunctival injection; (b)
- (c) To perform appellation of eye-lashes;
- (d) To measure the refractive error and advise correctional glasses;
- To perform nasolacrimal duct syringing for potency (e)

X. **Dental Procedures:**

To perform dental extraction

Community Healthy: XI

- To be able to supervise and motivate, community and para-professionals for corporate (a) efforts for the health care;
- To be able to carry on managerial responsibilities, e.g. Management of stores, (b) indenting and stock keeping and accounting
- Planning and management of health camps; (c)
- Implementation of national health programmes; (d)
- To effect proper sanitation measures in the community, e.g. disposal of infected (e) garbage, chlorination of drinking water;
- To identify and institute and institute control measures for epidemics including its (f) proper data collecting and reporting.

Forensic Medicine Including Toxicology XII

- To be able to carry on proper medico legal examination and documentation of injury (a) and age reports.
- To be able to conduct examination for sexual offences and intoxication; (b)
- To be able to preserve relevant ancillary material for medico legal examination; (c)
- To be able to identify important post-mortem findings in common un-natural deaths. (d)

Management of Emergency XIII

- To manage acute anaphylactic shock; (a)
- To manage peripheral vascular failure and shock; (b)
- To manage acute pulmonary oedema and LVF; (c)
- Emergency management of drowning, poisoning and seizures (d)
- Emergency management of bronchial asthma and status asthmaticus; (e)
- Emergency management of hyperpyrexia; (f)
- Emergency management of comatose patients regarding airways, positioning (g) prevention of aspiration and injuries
- Assess and administer emergency management of burns (h)

Syllabus for MICROBIOLOGY

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1	Broad Curriculum As Per MCI Guidelines for Microbiology	35-36
2	Syllabus of Microbiology	37-70

BROAD CURRICULUM AS PER MCI GUIDELINES (MICROBIOLOGY)

i) GOAL

The broad goal of the teaching of undergraduate students Microbiology is to provide an understanding of the natural history of infectious disease in order to deal with the etiology, pathologenesis, laboratory diagnosis, treatment and control of infections in community.

ii) OBJECTIVES

a. KNOWLEDGE

At the end of the course, the student should be able to:

- 1) State the infective micro-organisms of the human body and describe the host parasite relationship.
- List pathogenic micro-organisms (bacteria, viruses, parasites, fungi) and describe the pathogenesis of the diseases produced by them.
- State or indicate the modes of transmission of pathogenic and opportunistic organisms and their sources, including insect vectors responsible for transmission of infection.
- 4) Describe the mechanisms of immunity to infections.
- Acquire knowledge on suitable antimicrobial agents for treatment of infections and scope of immunotherapy and different vaccines available for prevention of communicable diseases.
- 6) Apply methods of disinfection and sterilization to control and prevent hospital and community acquired infections.
 7) Recommend laborations.
- 7) Recommend laboratory investigations regarding bacteriological examination of food, water, milk and air.

b. SKILLS

At the end of the course, the student should be able to:

- 1. Plan and interpret laboratory investigations for the diagnosis of infectious diseases and to correlate the clinical manifestations with the etiological agent.
- 2. Identify the common infectious agents with the help of laboratory procedures and use antimicrobial sensitivity tests to select suitable antimicrobial agents.

- 3. Perform commonly employed bed-side tests for detection of infectious agents such as blood film for malaria, filaria, gram staining and AFB staining and stool sample for
- 4. Use the correct method of collection, storage and transport of clinical material for
- microbiological investigations.

INTEGRATION (b)

The student should understand infectious diseases of national importance in relation to the clinical, therapeutic and preventive aspects.

IInd MBBS Syllabus for Microbiology, MGMIHS

GENERAL MICROBIOLOGY [n=17]

Sr. No.	Торіс	МК	DK	NK	No. of Hrs.
1	History & Microscopy				1hr
	Definitions of Medical Microbiology, Pathogen Commensal, Symbiotic infection,	√			
	Contribution of Louis Pasture				
	Robert Koch	√			
	Lister				
	Names of scientists who discovered common bacteria				
	Importance of Microbiology	\checkmark			
2	Morphology of bacteria I	\checkmark			1hr
	Difference between Prokaryotes & Eukaryotes	\checkmark			
	 Microscopy – Basic principles and applications of all microscopes 	\checkmark			
	Classification of staining techniques	\checkmark			
	Gram's stain and ZN stain in detail(with examples)	\checkmark	· 5,.		
	Negative staining, Impregnation method	\checkmark			
	Albert's stain	\checkmark			
3	Morphology II	\checkmark			1hr
	Morphology of Bacteria	$\overline{}$			
	Bacterial cell anatomy in detail	$\overline{\mathbf{v}}$			
	Bacterial Spore				·····

4	Physiology of bacteria				
	Bacterial cell division, Generation time, Bacterial growth curve	\checkmark	*****		
	Bacterial growth requirements				1.hr
	Bacterial Metabolism		\checkmark		
5	Sterilization	$\overline{\mathbf{A}}$			2hrs
&	 Definitions of Sterilization, disinfection, asepsis, antiseptics 				
6	 Need of Sterilization / Disinfection in various fields – Medical, Food & Pharma Industry 	\checkmark			
	 Physical methods of Sterilization Sunlight, Heat (dry & moist heat), Filtration, Radiation in details 	V			
	Working and efficacy testing of autoclave and hot air oven				
	Plasma sterilization central sterile supply department concept only			\checkmark	
7	Disinfection	****			1hr
	Characteristics of ideal chemical disinfectant	\checkmark			
	Factors influencing potency of a disinfectant	\checkmark			
	 Disinfectants like Aldehydes , Alcohols, phenols, Halogens, Oxidising agents, Salts, Surface Active Agent, Gases, Dyes (Concentration, Mode of action and uses only) 	V	** <u>*</u>		
	Testing of Disinfectants		\checkmark		
8	Culture Media				1hr
	Types of Media and their uses	\checkmark		1	
	Composition of Media	<u> </u>			-

9	Culture Methods				1hr
	Types of aerobic culture methods and their uses	√			
	Types of anaerobic culture methods and their uses	$\overline{}$			
	 Mc Intosh Filde's Jar – Functioning and uses 		******		
10	Identification of bacteria	$\overline{\mathbf{v}}$			1hr
	Morphology of bacteria (Gram stain), Motility				
	Biochemical tests (Principle and examples)				
	Morphology of Bacterial Colony		√		
	Biochemical tests (Procedure)				
	Typing Method		V		
	Pathogenicity tests				
11	 Antimicrobial therapy Mode of action of antimicrobial agents Antibiotic Sensitivity tests : Kirby Bauer disc diffusion Importance of making an antibiotic Policy Strict Adherences to antibiotic Policy 	√			1hr
	Antibiotic Sensitivity tests: Stokes Disc diffusion E-test		\checkmark		
	 Dilution test MIC , MBC procedure other than Kirby- Baur AST testing 		^н ъ.	V	
12	Bacterial Genetics				
	Basic structure of Bacterial DNA				
&	Definitions of Gene, Codon, Nonsense codons	√			2hr
13	Extrachromosomal elements – Plasmids, Episomes, Transposons				

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	Difference between Phenotypic & Genotypic variation (\checkmark			
	Mutation) Genetic Variation (Mutation)	√			
	 Gene transfer (Transformation, Transduction, Lysogenic 				
	 conversion, Conjugation) 	√			
	 Differences between Mutational & Transferable drug resistance 	\checkmark			
	 mechanism of drug resistance 	\checkmark			
	 Synthesis of Polypeptide (Transcription and translation) 			\checkmark	
	Genetic Engineering		√		
	DNA Probes				
	• PCR			_√	
	Gene therapy			\checkmark	
.4	Bio- medical waste disposal				1hr
	Definition of Biomedical waste	$\overline{\mathbf{A}}$			
	Classification & disposal as per categories	\checkmark			
	Importance of segregation	\checkmark			
15	Universal Safety Precautions & health care associated infections		· N.		1hr
	Universal safety precautions	$\overline{\mathbf{A}}$			
	Hand Hygiene	\checkmark			
	Definition of Hospital acquired infection.	\checkmark			
	 Sources, types, prevention and control of health care associated infections 	1			

16	Normal microbial flora of human body		1hr
	 Introduction – Various sites, types & role 	√	
17	Infection /		1hr
	Host parasite relationship		
	 Definitions of Saprophytes, Parasitic, Commensals, Pathogen, Opportunistic Pathogens, Pathogenicity, Virulence 	V	
	Types of infection, Routes of transmission	\checkmark	
	Sources of Infection	 √	
		V	
	Difference between Exotoxins & Endotoxins	\checkmark	
	Types of Infectious diseases Localized, Generalized, Endemic, Epidemic, Pandemic	√	
	Factors Predisposing to Microbial Pathogenicity		

SYSTEMIC BACTERIOLOGY [n=29]

Sr.	Торіс	МК	DK	NK	No. of
no.	Must Know				Hrs
	Subtopics				
1	Staphylococci	\checkmark			1hr
	A: Classification	\checkmark			
	B: Morphology	√			
	C: Culture Characteristics	\checkmark			
	D: Biochemical reactions				
	E: Antigens	√			
	F: Pathogenesis & diseases caused in detail	\checkmark			
	G: Laboratory diagnosis	\checkmark			
	H: Prevention & control	√			
	I: Special identification tests	√			
	MRSA				
2	Streptococci				1hr
	A: Classification	√			
	B: Morphology	√			
	C: Culture Characteristics	√			
	D: Biochemical reactions		$\overline{\mathbf{v}}$	_	
	E: Antigens	\checkmark			
	F: Pathogenesis & diseases caused in detail	√			
	G: Laboratory diagnosis	√			
	H: Prevention & control	\checkmark			
	I: Special identification tests				

				1hr
Other streptococci and Pneumococci				
A: Classification				
B: Morphology	\checkmark			
C: Culture Characteristics	\checkmark			
D: Biochemical reactions				
E: Antigens	\checkmark			×
F: Pathogenesis & diseases caused in detail	\checkmark			
G: Laboratory diagnosis	\checkmark			
H: Prevention & control	\checkmark			
I: Special identification tests	\checkmark			
Group - B streptococci			\checkmark	1
Neisseria				1hr
A: Classification	\checkmark			
B: Morphology	\checkmark			
C: Culture Characteristics	\checkmark			
D: Biochemical reactions		V		
E: Antigens	\checkmark			
F: Pathogenesis & diseases caused in detail	\checkmark			2
G: Laboratory diagnosis	\checkmark	7		
H: Prevention & control	\checkmark			
I: Special identification tests	\checkmark			

				1hr
C. diptheriae				
A: Classification	\checkmark			
B: Morphology				
C: Culture Characteristics				
D: Biochemical reactions		1		
E: Antigens	\checkmark			
F: Pathogenesis & diseases caused in detail	7			
G: Laboratory diagnosis	√			
H: Prevention & control	√			
I: Special identification tests	7			
Diptheroides				
M. tuberculosis				1hr
A: Classification	1			
B: Morphology	√			
C: Culture Characteristics	\checkmark			
D: Biochemical reactions				
E: Antigens	\checkmark			
F: Pathogenesis & diseases caused in detail	\checkmark			
G: Laboratory diagnosis	√			
H: Prevention & control	\checkmark		_	
I: Special identification tests	7			
MDR, XDR			\downarrow	

7	Atypical mycobacteria				1hr
	Name of the Species				
	Names of the diseases caused				
	 Brief outline of lab diagnosis 				
	Special tests for identification		V		
8	M. leprae				1hr
	A: Classification				
	B: Morphology	$\overline{\mathbf{A}}$			
	C: Culture Characteristics	\checkmark			
	D: Biochemical reactions				
	E: Antigens	\checkmark			
	F: Pathogenesis & diseases caused in detail	\checkmark			
	G: Laboratory diagnosis	\checkmark			
	H: Prevention & control				
	I: Special identification tests	\checkmark			
9	 Bacillus Name of the Species Names of the diseases caused 	V			1hr
	Brief outline of lab diagnosis		15		
	Special tests for identification		\checkmark		
10	 Method of anaerobiasis & Nonsporing anaerobes Method of anaerobiasis Nonsporing anaerobes { Name of the Species, Names of the diseases caused Brief outline of lab diagnosis } 	√		,	1hr
	Special tests for identification		√		

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.1	Clostridium – I			1hr	
_	A: Classification				
	B: Morphology				
	C: Culture Characteristics		-		
	D: Biochemical reactions		$\overline{\mathbf{V}}$		
	E: Antigens				
	F: Pathogenesis & diseases caused in detail				
	G: Laboratory diagnosis	\checkmark			
	H: Prevention & control	\checkmark			
	I: Special identification tests	\checkmark			
2	Clostridium – II				
-	A: Classification				
	B: Morphology		- <u>·</u>	1hr	
	C: Culture Characteristics				
	D: Biochemical reactions				
-	E: Antigens		N		
-	F: Pathogenesis & diseases caused in detail		1		
	G: Laboratory diagnosis				
ŀ	H: Prevention & control				
	I: Special identification tests				
	Cl.botulinum				

<u> </u>				
13	Enterobacteriacae – I			1hr
	(E. coli.)			
	A: Classification			·····
	B: Morphology	$\overline{}$		
	C: Culture Characteristics			
	D: Biochemical reactions			
	E: Antigens	$\overline{}$		
	F: Pathogenesis & diseases caused in detail			
	G: Laboratory diagnosis			
	H: Prevention & control			
	I: Special identification tests			
	Enterobacteriacae – II Proteus & Klebsiella			
	A: Classification	$\overline{}$		
	B: Morphology	√		
	C: Culture Characteristics			
	D: Biochemical reactions		$\overline{\mathbf{v}}$	
	E: Antigens		· · · · · · · · · · · · · · · · · · ·	
	F: Pathogenesis & diseases caused in detail			
	G: Laboratory diagnosis	$\overline{}$		
	H: Prevention & control			
	I: Special identification tests	$\sqrt{1}$		

 5					14-
	Enterobacteriacae – III				1hr
	Salmonella				
	A: Classification				
	B: Morphology				
	C: Culture Characteristics	1			
	D: Biochemical reactions				
	E: Antigens	1			
	F: Pathogenesis & diseases caused in detail	√			
	G: Laboratory diagnosis	\checkmark			-
	H: Prevention & control	√			
	I: Special identification tests	1			
	Antigenic variation		\checkmark		
~~~	Shigella				1hr
	A: Classification				
	B: Morphology				
	C: Culture Characteristics	1			
ľ	D: Biochemical reactions				
	E: Antigens	√			
	F: Pathogenesis & diseases caused in detail	√			
	G: Laboratory diagnosis	√			
ſ	H: Prevention & control	$\checkmark$		1	
	I: Special identification tests				

17	Vibrio			1hr
	A: Classification	√		
	B: Morphology	$\overline{\mathbf{v}}$		
	C: Culture Characteristics	√		
	D: Biochemical reactions		1	
	E: Antigens	√		
	F: Pathogenesis & diseases caused in detail	$\checkmark$		
	G: Laboratory diagnosis	$\checkmark$		
	H: Prevention & control	$\checkmark$		
	I: Special identification tests	7		
	Halophilic vibrios			
18	Campylobacter & Helicobacter	1		1hr
	Name of the Species			
	Names of the diseases caused			
	Special tests for identification			
	Brief outline of lab diagnosis			
19	Pseudomonas			1hr
	A: Classification	$\checkmark$		
	B: Morphology	$\checkmark$		
	C: Culture Characteristics	1		
	D: Biochemical reactions		$\checkmark$	
	E: Antigens	$\checkmark$		
	F: Pathogenesis & diseases caused in detail	$\checkmark$		
	G: Laboratory diagnosis	$\overline{\mathbf{v}}$		
	H: Prevention & control	√		· · · · · · · · · · · · · · · · · · ·

	I: Special identification tests	$\checkmark$			
	Burkholderia species				
20					
20	Other GNB I	9			1hr
	(Yersinia, Pasteurella, Francisella, Bordetella)				2.11
	Name of the Species				
	Names of the diseases caused				
	Special tests for identification				
	Brief outline of lab diagnosis		-		
	Pastaurella & Francisella - infections caused				-
21	Other GNB II			V	1hr
	(Haemophilus, Brucella)				1111
	Name of the Species	$\checkmark$			_
	Names of the diseases caused	v			
	Brief outline of lab diagnosis				
	Special tests for identification		V		
22	Miscellaneous Bacteria		v		
	(Newer bacteria's)				1hr
	Name of the Species				
	Names of the diseases caused	$\checkmark$			
	Special tests for identification				
	Brief outline of lab diagnosis				
3	Spirochaete - I				
	(Treponema species )				1hr
	A: Classification				
		V			
	B: Morphology	$\checkmark$	100		
	C: Culture Characteristics		14 ¹¹		
		$\checkmark$			
	E: Antigens	V			
	F: Pathogenesis & dispasso poursed in the stat				
-	F: Pathogenesis & diseases caused in detail	$\checkmark$	4		
(	G: Laboratory diagnosis	$\checkmark$			
	H: Prevention & control	V			
	. revention & control				
1	: Special identification tests				
			$\checkmark$		

Contraction of the second s	i i			1hr
Spirochaete –II (Borrelia, Leptospira)				
(borrena, reprospira)				
A: Classification	<b>√</b>			
B: Morphology	√			
C: Culture Characteristics	√			
E: Antigens	√			
F: Pathogenesis & diseases caused in detail	√			
G: Laboratory diagnosis	$\checkmark$			
H: Prevention & control	$\checkmark$			
I: Special identification tests	$\checkmark$		-	
Actinomycete and Nocardia				1hr
A: Classification	$\checkmark$			
B: Morphology	$\checkmark$			
C: Culture Characteristics	$\checkmark$			
D: Biochemical reactions		<b>√</b>		
E: Antigens	7			
F: Pathogenesis & diseases caused in detail	$\overline{}$			
G: Laboratory diagnosis	√	_		
H: Prevention & control	$\checkmark$			
I: Special identification tests	√			
				ļ
••	A: Classification         B: Morphology         C: Culture Characteristics         E: Antigens         F: Pathogenesis & diseases caused in detail         G: Laboratory diagnosis         H: Prevention & control         I: Special identification tests         Actinomycete and Nocardia         A: Classification         B: Morphology         C: Culture Characteristics         D: Biochemical reactions         E: Antigens	A: Classification       √         B: Morphology       √         C: Culture Characteristics       √         E: Antigens       √         F: Pathogenesis & diseases caused in detail       √         G: Laboratory diagnosis       √         H: Prevention & control       √         I: Special identification tests       √         Actinomycete and Nocardia       √         R: Morphology       √         C: Culture Characteristics       √         B: Morphology       √         C: Culture Characteristics       √         P: Biochemical reactions       ✓         E: Antigens       √         F: Pathogenesis & diseases caused in detail       √         H: Prevention & control       √	A: Classification       √         B: Morphology       √         C: Culture Characteristics       √         E: Antigens       √         F: Pathogenesis & diseases caused in detail       √         G: Laboratory diagnosis       √         H: Prevention & control       √         I: Special identification tests       √         Actinomycete and Nocardia       √         B: Morphology       √         C: Culture Characteristics       √         D: Biochemical reactions       √         E: Antigens       √         F: Pathogenesis & diseases caused in detail       √         A: Classification       √         B: Morphology       √         C: Culture Characteristics       √         D: Biochemical reactions       √         E: Antigens       √         F: Pathogenesis & diseases caused in detail       √         G: Laboratory diagnosis       √         H: Prevention & control       √	A: Classification       Image: Constraint of the second seco

26	Rickettsia			1hr
	Name of the Species			
	Names of the diseases caused			
	Special tests for identification			
	<ul> <li>Brief outline of lab diagnosis</li> </ul>			
27	Chlamydia			1hr
	Name of the Species			 
	<ul> <li>Names of the diseases caused</li> </ul>			
	Brief outline of lab diagnosis			
28	Mycoplasma			1hr
	Name of the Species	$\overline{}$		
	Names of the diseases caused			
	Brief outline of lab diagnosis			
	Special tests for identification		$\checkmark$	
29	Bacteriology of water, Air & Milk	$\checkmark$		 1hr
	Bacteriological Examination of Air			 
	Acceptable limit of Air pollution			
	Bacterial flora in water			
	Water Borne Pathogens		,	
	Bacteriological Examination of water		$\checkmark$	
	Milk Borne diseases			
	Bacteriological Examination of milk			
	Procedures for bacteriological examination of milk & water			

# MYCOLOGY [n=4]

Sr.	Торіс	МК	DK	NK	No. of Hrs
No.					
1	Introduction to mycology	$\checkmark$			1hr
	<ul> <li>Introduction to Mycology</li> </ul>				
	Difference between fungus & Bacteria				
	<ul> <li>Classification of fungi with examples</li> </ul>				
	<ul> <li>Reproduction &amp; Sporulation</li> </ul>				
	<ul> <li>Lab diagnosis of mycosis</li> </ul>				
	Classification of Fungal diseases				
2	Agents of superficial mycosis (Dermatophytes)				1hr
	Superficial Mycosis				
	a. Enumerate agents				
	b. Predisposing factors				
	c. Lab diagnosis (Outline)				
	<ul> <li>Colony characteristics of dermatophytes</li> </ul>		V		
3	Subcutaneous mycosis & Candida	$\checkmark$			1hr
	Subcutaneous Mycosis & Candida in detail				
	a. Enumerate agents				
	b. Predisposing factors				
	c. Lab diagnosis (Outline)				
4	Systemic mycosis & opportunistic fungal infections & P. Carinii				1hr
	Systemic & Opportunistic Mycosis				
	a. Enumerate agents				
	b. Predisposing factors				
	c. Candida, Cryptococcus in detail				
	d. Mucor, Aspergillus		-		
	Histoplasma		$\checkmark$		
	P. Carinii				
	Mycetism				

# VIROLOGY [n=16]

Sr.n o.	Торіс	МК	DK	NK	No.of hrs
1	General virology – I				 1hr
	Morphology of Viruses				
	Replication of viruses				
	<ul> <li>Chemical properties of viruses</li> <li>Susceptibility to physical and chemical agents</li> <li>Viral Haemagglutinin</li> </ul>		√		
2	General virology – II				1hr
	<ul> <li>Cultivation of Viruses ,Viral assays</li> <li>Outline of diagnosis of viral diseases</li> </ul>	$\checkmark$			
;	Virus-host interactions				1hr
	<ul> <li>Inclusion Bodies</li> <li>Routes of transmission of viral infections</li> <li>Interferons</li> <li>Immunity in viral diseases</li> </ul>	√			
	Host responses to virus infections				······
	Viral vaccines and antiviral agents				1hr
	<ul> <li>Commonly used viral vaccines <ul> <li>Types and Schedule</li> <li>List of antiviral agents.</li> </ul> </li> </ul>	~			
	Mode of preparation				
	Chemoprophylaxis				
-	Chemotherapy of viral diseases			<b>v</b>	

1	1			1	
5	Pox viruses				1hrs
	Bacteriophage				
	<ul> <li>Morphology</li> </ul>				
	<ul> <li>Names of poxviruses and diseases caused</li> </ul>				
	<ul> <li>Bacteriophage[Basic structure and Significance]</li> </ul>				
	Cultivation		$\checkmark$		
6	Herpes simplex & Varicella zoster CMV, EBV				1hr
	Morphology	$\checkmark$			
	Classification				
	<ul> <li>HSV (Infections caused and Lab diagnosis)</li> </ul>				
	<ul> <li>Varicella – Zoster (Infections caused and Lab diagnosis)</li> </ul>				
	EBV (Infections caused and Lab diagnosis)		$\checkmark$		
	CMV (Infections caused and Lab diagnosis				
7	Other DNA viruses				1hr
	(Papova, Adeno, )				
	Basic morphology, diseases caused				
	Outline of lab diagnosis		$\checkmark$	31.	
8	Orthomyxoviruses				1hr
	Differences between Orthomyxo & paramyxo viruses	$\checkmark$	1		
	Influenza Virus				
	a. Morphology				
	b. Antigenic classification and structure				
	c. Antigenic shift and Antigenic drift				
	d. Pathogenesis and lab diagnosis				
	Influenza Virus				
	Antigenic classification				
	Prophylaxis				L.
	Bird flu, Swine flu				
	avense gar pasar * 18-1 titt"'titt pitter			1576	

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9	Paramyxoviruses				1h
	Morphology				
	Measles virus and Mump Virus				
	Parainfluenza virus			_	
	• RSV				
10	Picornaviruses				1hr
	Classification	$\checkmark$			
	<ul> <li>Polio virus in detail</li> </ul>				
	<ul> <li>Differences between killed and live vaccines</li> </ul>				
	<ul> <li>Eradication and Prophylaxis of Polio virus</li> </ul>				
	Coxsackie viruses		$\checkmark$		
	Rhino virus			$\checkmark$	
11				-	1hr
	Hepatitis viruses				
	<ul> <li>HAV (Pathogenesis and Lab diagnosis)</li> </ul>	$\overline{\mathbf{v}}$			
	HBV (Morphology, Mode of transmission, Clinical features, Lab	V			
	diagnosis)				
	<ul> <li>HCV (Morphology, Mode of transmission, Clinical features, Lab diagnosis)</li> </ul>				
	<ul> <li>HDV &amp; HEV (Pathogenesis &amp; Lab diagnosis)</li> </ul>				
12	Arboviruses		shir		1hr
	Classification , Names of Arboviruses and diseases caused			-	
	Dengue Virus in detail	V			
	- 18687 9				
	JE yellow fever KFD				
			1	1	1

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Y.				
	Morphology			
	<ul> <li>Pathogenesis and Lab diagnosis</li> </ul>			
	<ul> <li>Prophylaxis</li> </ul>			
4	Retro Viruses – HIV			  1hr
	<ul> <li>Morphology</li> </ul>	$\overline{\mathbf{v}}$		 
	Resistance			
	<ul> <li>Modes of Transmission</li> </ul>			
	Pathogenesis			
	<ul> <li>Opportunistic infections and malignancies</li> </ul>			
	Lab diagnosis in detail			
	Prevention			
	• PEP			
	Viral genes & antigens			
	• ART		$\checkmark$	
	Strategies for HIV testing			
.5	Miscellaneous viruses			 1hr
	<ul> <li>Viruses (Only names) causing gastroenteritis</li> </ul>	$\overline{}$		 
	<ul> <li>Viruses causing viral hemorrhagic fevers (only names)</li> </ul>	<b>v</b>		
	<ul> <li>Slow virus diseases (Only names)</li> </ul>			
.6	Oncogenic viruses	1		1hr
	Papilloma Virus	$\overline{\mathbf{v}}$		 
	Only names of oncogenic viruses and malignancies caused			

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## IMMUNOLOGY [n=12]

Sr. No.	Торіс	МК	DK	NK	No. of Hrs
1	Immunity			-	1hr
	<ul> <li>Innate Immunity – Types, Factors influencing innate immunity, Mechanisms</li> <li>Acquire Immunity -         <ul> <li>a. Active Immunity</li> <li>b. Passive Immunity</li> </ul> </li> <li>Combined immunization</li> <li>Adoptive immunity</li> <li>Local Immunity</li> <li>Herd Immunity</li> </ul>	<b>√</b>	√		
2	Antigen				1hr
	<ul> <li>Types of Antigens</li> <li>Factors determining antigenicity</li> <li>Super antigens</li> </ul>	$\checkmark$			
	Antibody		V		1hr
	<ul> <li>Properties of antibodies</li> <li>Structure of Immunoglobulin classes</li> <li>IgG, IgM, IgA, IgD, IgE <ul> <li>a) Basic structure, function</li> <li>&amp; distribution</li> </ul> </li> </ul>	√	2. 2.		
	<ul> <li>IgG, IgM, IgA, IgD, IgE Mol. Wt., Sedimentation Coefficient</li> </ul>		$\checkmark$		
	Abnormal Immunoglobulins				
4	Complement			1hr	
--------	--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------	--------------	-------------	---------	
	Components of complement				
	Classical Pathway				
	Alternative Pathway	$\checkmark$			
	Biological effects of complement	•			
	Deficiencies of complement				
	Regulation of complement activation				
	<ul> <li>Biosynthesis of Complement</li> </ul>				
	Quantitation of complement				
5	Ag-Ab reactions I			 	
& 6	<ul> <li>Types of Ag – Ab reactions, precipitation, Agglutination, CFT, Neutralization, Opsonisation, Immunoflourescence, ELISA Immunochromatography (Principle, Types &amp; uses only)</li> </ul>	$\checkmark$		2hrs	
7	Structure & function of Immune system			 1hr	
	<ul> <li>Central Lymphoid Organs</li> <li>Peripheral Lymphoid Organs</li> <li>Cells of Lymphoreticular System</li> <li>HLA</li> <li>Differences between T &amp; B cells</li> </ul>	√			
	Lymphocytic recirculation			 	
8	Immune response			1hr	
	<ul> <li>Humoral Immune Response Primary and secondary responses Production of Antibodies Factors influencing antibody production</li> <li>Cell mediated Immune Response Cytokines &amp; Lymphokines – Types &amp; functions only</li> </ul>	√	** <u>.</u>		
	Cell mediated Immune Response     Detection of CMI		√		
	Immunological tolerance     Monoclonal antibodies				

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9	Hypersensitivity				1\   r
	Definition & Classification				
	Type 1 Reaction				
	Differences between Immediate & delayed hypersensitivity	$\checkmark$			
	• Type 2, 3 & 4 Reactions				
10	Autoimmunity				1hr
	Definition & Mechanisms     Classification with examples	√			
	Classification with examples				1hr
11	Immunodeficiency diseases				
	<ul> <li>Definition, classification and examples of diseases</li> </ul>				
	Laboratory test for detection.			$\checkmark$	
12	Transplantation & Tumor immunity				1hr
	Types of Transplants				
	Allograft reaction				
	Histocompatibility Antigens	$\checkmark$			
	Histocompatibility Testing				
	Graft - Versus - Host reaction				
	Tumor antigens				
	Immunosurveillance		$\checkmark$		
	Immune response to malignancy			$\bigvee$	
	Immunotherapy of cancer				

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## PARASITOLOGY [n=10]

Sr. no.	Topic	МК	DK	NK	No. of
					Hrs.
1	Introduction to parasitology				1hr
	Classification of Parasites				
	Type of Parasites				
	Hosts (Definitive & Intermediate)				
	<ul> <li>Host – Parasitic relationship</li> </ul>				
	Sources of Infection				
	Lab diagnosis in general				
2	E.Histolytica				1hr
	Morphology				
	Life cycle				
	Pathogenesis & Complications				
	<ul> <li>Lab diagnosis &amp; treatment</li> </ul>				
	Non Pathogenic amoebae				
	Free living amoebae				
3	Giardia, Trichomonas				1hr
	<ul> <li>Giardia lamblia (Morphology, life Cycle, Pathogenesis, Lab diagnosis &amp; tractment)</li> </ul>				
	treatment)				
	<ul> <li>Trichomonas vaginalis (Morphology Pathogenesis, Lab diagnosis &amp; treatment</li> </ul>				
4					1hr
	Malaria		s a _{n c}		
	Life Cycle, Morphology, Pathogenecity & Lab diagnosis,				
	prevention				
5	Haemoflagellates				1hr
	<ul> <li>Leishmania (Classification, diseases caused)</li> <li>L. donovani in details</li> </ul>				
	Morphology, Lifecycle, Pathogenesis, Lab diagnosis				

	Trypanosomes		<u> </u>		<u> </u>
~			$\checkmark$		
5	Miscellaneous protozoa				
					1
	<ul> <li>Toxoplasma (Morphology life Cycle, Pathogenesis, Lab diagnosis)</li> <li>Cryptosporidium, Isospora</li> </ul>	$\overline{\mathbf{v}}$			
	Cryptosporidium, Isospora	v			
	B.coli				
	Cestodes				
					1h
	<ul> <li>Taenia &amp; Echinococcus (Morphology life Cycle, Pathogenecity, diagnosis)</li> </ul>				
	diagnosis)	, Lab   √			
	Brief mention about other cestodes				
	Trematodes				
	Schistosomes				1hr
	Names & diseases caused				
	<ul> <li>Morphology, Life Cycle, Pathogenicity &amp; Lab diagnosis</li> <li>Easciele benetic</li> </ul>				
	<ul> <li>Fasciola hepatica</li> </ul>				
	Parognimus westermani				
	Nematodes (Intestinal) I				
ŕ	• A duodonalo A hard i i i				1hr
	A. duodenale, A. lumbricoides,     E. vermicularis, T. tricker, //				
	E. vermicularis, T. trichura (in details)				
ſ	S. stercoralis				
					+
	Tissue Nematodes II				
	& Stool concentration techniques				1hr
1	W. bancrofti ( in details)		_		
	T. Spiralis				
•	- incumensis ( in decails)				
•	stool concentration techniques				
•	Name of parasites in stool				
•	Names of parasites affecting CNS				
•	Names of parasites affecting liver				
•	Names of parasites entering through skin				
	ble stained eggs& Eggs which float in saturated as the last second			}	
W	hich do not	1		1	

# APPLIED MICROBIOLOGY (To be taken in the form of UG seminars/Tutorials)

(n=8)

Topic	Торіс	No. of Hrs.
No		
	Only Causative agents & Brief Outline of Lab diagnosis in	
1	Gastrointestinal infections	1hr
2	• URTI	1hr
3	• LRTI	1hr
4	• UTI	1hr
5	CNS Infections	1hr ·
5	• Wound & Pyogenic infections	1hr
	PUO & infections	1hr
•	STDs	1hr

# Theory: (n=96)

Section	No of lectures
General Microbiology	17
Systemic bacteriology	29
Mycology	04
Virology	16
Immunology	12
Parasitology	10
eminars/Tutorials on Applied Microbiology	08
Total	96

Practicals: Including Extra coaching,	Revisions & classroom account	6.000	
No Function	and the classific of assessment	(CRA)	(n=132)

No	Experiments	No.of Hrs
1.	Microscopy	4hrs
2.	Morphology of bacteria	4hrs
3.	Sterilisation and Disinfection	4/1/5 4/hrs
4.	Principles in diagnostic Microbiology 1	4hrs
5.	Principles in diagnostic Microbiology 2	4///s
6.	Immunology and Serologigal methods	
7.	Staphylococci	4hrs
8.	Streptococci and Pneumococci	4hrs
9.	Neisseria	4hrs
10.	Corynebacteria	4hrs
11.	Bacillus	4hrs
12.		4hrs
13.	M.tuberculosis and Atypical Mycobacteria M.leprae	4hrs
14.		4hrs
	E.coli,Klebsiella and Proteus	4hrs
15.	Salmonella	4hrs
16.	Shigella and Vibrio	4hrs
17.	Pseudomonas and Hospital infections	4hrs
18.	Yersinia and Brucella	4hrs
19.	Haemophillus and Bordetella	4hrs
20.	Clostridia	4hrs
21.	Non-sporing anaerobes	4hrs
2.	Spirochaetes	4hrs 4hrs
3.	Actinomycetes and Nocardia	4hrs 4hrs
4.	Virology	
5.	Intestinal protozoa	4hrs
6.	Blood and tissue protozoa	4hrs
		4hrs

	TOTAL	132Hrs
33.	Mycology	4hrs
32.	Medical Entamology	4hrs
31.	Tissue nematodes	4hrs
30.	Intestinal nematodes	4hrs
29.	Trematodes	4hrs
28.	Cestodes	4hrs
27.	Blood and tissue flagellates	4hrs

# Total Teaching Hours: 250 hours (As per MCI)

Lectures + Seminars/Tutorials	96Hrs
Practicals Including Extra coaching & Revisions	132Hrs
Assessments	22Hrs
Total	250Hrs

#### **Books Recommended :**

Sr. No.	Name of the Book	Author
1	Textbook of Microbiology	R. Ananthanarayan C K Jayaram Panikar
2	A Textbook of Microbiology	P. Chakraborty
3	Textbook of Medical Microbiology	Rajesh Bhatia & Itchpujani
4	Textbook of Medical Microbiology	Prof C.P. Baveja
5	Textbook of Medical Parasitology	C K Jayaram Panikar
6	Medical Parasitology	C.P.Baveja V.Baveja
7	Textbook of Medical Parasitology	S C Parija
8	Textbook of Parasitology	Damle and Karyakarte
8	A Textbook of Parasitology	Dr.K.D. Chatterjee.
9	Practical Microbiology	Dr. Anuradha De

### **Reference Books :**

Sr. No.	Name of the Book	Author
1	Mackie McCartney practical Medical Microbiology	Colle JG, Fraser AG
2	Principles of Bacteriology, Virology & Immunology Vol. 1, 2, 3, 4, 5	Topley Wilsons
3	Medical Mycology (Emmons)	Kwon – Chung
4	Review of Medical Microbiology (Lange)	Jawetz
5	Immunology	Weir DM
6	Medical Microbiology	David Greenwood, Richard Stack, John Pentherer
7	Medical Virology	Timbury MC
8	Mackie McCartney Medical Microbiology Vol.1	Duguid JP
9.	Textbook of Microbiology	Monica Cheesebrough

#### Evaluation

a. Methods

Theory, Practical & Viva

		Particulars	Total Marks
No.	Theory (Total out of 95 Marks)	Theory ( 2 Papers – 40 Marks each)	80 Marks
1		Oral (Viva)	15 Marks
2	Practical (Total out of 25	Practical	25 Marks
2	Marks) Internal Assessment (Total out	Internal Assessment (Theory – 15 Practical – 15)	30 Marks
5	of 30 Marks)	TOTAL	150 Marks

**Passing:** A candidate has to obtain minimum of 47 Marks out of 95 in Theory, 13 marks out of 25 in Practical, 11 marks out of 30 in Internal Assessment and 75marks out of 150 Total to be declared as passed.

## Nature of Question Paper :- Theory (Total 80 Marks)

Paper - 1	General Microbiology , Systemic Bacteriology & Related Applied Microbiology	40 Marks
	Immunology, Virology, Parasitology , Mycology &	40 Marks
Paper – II	Related Applied Microbiology	

Section	Question Description	<b>Division of Marks</b>	Total Marks
A	MCQs (16)	16 x 0.5 Marks	08 Marks
B	Brief Answer Questions (4/5)	4 x 4 Marks	16 Marks
c	Long Answer Questions (2/3)	2 x 8 Marks	16 Marks
TOTAL	(2/3)		40 Marks

## Practical Examination Marks distribution : -

No.	Particulars	Marks	
1	Grams Staining	5 Marks	
2	ZN Staining	5 Marks	
3	Stool Examination	5 Marks	
4 Spots (10)		10 Marks	
	TOTAL	25 Marks	

#### Viva (Two Tables)

	Total	15 Marks
В	Parasitology, Virology , Mycology, Immunology	7 Marks
A	General Microbiology, Systemic Bacteriology and Applied Microbiology	8 Marks

## Distribution of MCQs:

PAPER 1: 16 MCQs , Marks 0.5 each= 8Marks

General Microbiology	06 MCQs	
Systemic Bacteriology	10MCQs	
Total	16 MCQs	

PAPER 2: 16 MCQs , Marks 0.5 each= 8Marks

Parasitology	05MCQs		
Mycology	03MCQs		
Virology	04MCQs		
Immunology	04MCQs		
Total	16 MCQs		

#### DIRECTION:-

For paper setting out of total marks ,70%, 20% and 10 % marks must be from must know , desirable to know, and nice to know portion, respectively

However all LAQs and MCQs are to be from must know area.

Internal Assessment shall be computed on the basis of three term ending examinations (Two Terminals & One Preliminary examination).

Examination	No. of Papers	Pattern	Duration of each paper	Total Marks
l st Terminal	1 (40 Marks)	MCQs = 16 (8 Marks) BAQs = 4/5 (16 Marks) LAQs = 2/3 (16 Marks)	2 Hours	40 Marks
ll nd Terminal	1 (40 Marks)	MCQs = 16 (8 Marks) BAQs = 4/5 (16 Marks) LAQs = 2/3 (16 Marks)	2 Hours	40 Marks
Prelim	2 (40 Marks each)	MCQs = 16 (8 Marks) BAQs = 4/5 (16 Marks) LAQs = 2/3 (16 Marks)	2 Hours each paper	80 Marks

D:\Microbiolgy dept\d\Revised Syllabus for MBBS\Lanscape Final revised MBBS Syllabus 15.07.2015.docx

HYProved in Bom 23/2012, Duted 30/03/2012.

Resolution 210. - 4

Resolved to approve the following recommendations [Sr. Nos. 1 to 39] of the Academic Council [AC-12/2012] dated 24.03.2012

5. In Microbiology theory syllabus for MBBS-II, the topic of Mycology be shifted from re-TV Paper – II to Paper – I. [Annexure - IV].

Proposed M.D Micro Exam Scheme for 2 days

#### DAY 1

- 1. Short case
- 2. Long case Plating from broth and work up with discussion and further tests for identification from solid media.
- 3. Serology A ) WIDAL test or VDRL test
- 4. Mycology
  - Slant LPCB with slide culture e
    - Yeast identification 65
- 5. HIV (ELISA) Procedure, result & discussion
- 6. Pedagogy

DAY 2

1. Short case final identification.

- 2. Long case final identification.
- 3. Serology WIDAL test Reading
- 4. Fungal final identification.
- 5. Parasitology
  - Stool examination R & M

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 Stool Cryptosporidium – Modified ZN staining / Malarial parasite – Leishman staining

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#### 6. 10 slides

7. Grand viva

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ISTILIS ANNEXURG-4 (Microbiology)

MGM INSTITUTE OF HEALTH SCIENCES (Deemed University u/s 3 of UGC Act, 1956) Grade 'A' Accredited by NAAC Sector-1, Kamothe, Navi Mumbai - 410209 Tel. No. 022-27432471, 022-27432994, Fax No. 022 - 27431094 E-mail : registrar@mgmuhs.com ; Website : www.mgmuhs.com

# SECOND YEAR MBBS

1

FE

Annexure - 4

# PARA-CLINICAL

SYLLABUS FOR THE SUBJECT OF SECOND YEAR MBBS COURSE AT CONSTITUENT COLLEGES OF MGM INSTITUTE OF HEALTH SCIENCES, NAVI MUMBAI / AURANGABAD

(Approved in Born 40/2015, 2014ed 13.03.2015, Resortation No. - 3.2 (F))

MGM Institute Of Health Sciences	
INWARD NO. 485G	
CATE: 1.5/7/15	

## EXAMINATION PATTERN FOR PATHOLOGY, MICROBIOLOGY & PHARMACOLOGY GENERAL SECTION

#### A. PASSING:-

- i. A candidate must obtain 50% in aggregate with a minimum of 50% in Theory including oral and minimum of 50% in practical and 35% in internal assessment combined theory and practical.
- Prelims examination on the basis of University pattern (Theory, Practical and viva): Minimum 3-4 weeks gap between Prelims and University examination.
- iii. The total time will be 2 hours each for theory papers of 40 marks.
- iv. Practical (total time 3 hours). The details of Practical examination exercises will be notified by Head of the department / Head of Institution.
- v. Prelim pattern will be as per the University exam with 2 papers in theory each of 2 hours duration.

## B. CALCULATION OF INTERNAL ASSESSMENT MARKS:

- Calculation of Theory and Practical Internal Assessment marks for Pathology, Microbiology & Pharmacology shall be as per following rule
- 1. Distribution of 15 marks in theory shall be as follows:
  - 1.1 5 marks for attendance as per the following guidelines:
    - Below 75% -0

Upto75% -2.5

Above 75% proportionately higher marks at pro-rate basis (multiplication factor is 0.1)

- 1.2 10 marks for academic performance in theory in 2 term and prelim exam-(average of all the 3 internal examination shall be taken)
- 1.3 Marks in decimal computed in 1.1, 1.2 & 1.3 should be converted into whole number at the end.

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- 2. Distribution of 15 marks in practical shall be as follow:
  - 2.1 5 marks for attendance as per the following guidelines:

Below 75%-0

Upto 75% -2.5

Above 75% proportionately higher marks at pro -rate basis (multiplication factor is 0.1)

- 2.2 10 marks for academic performance in Practicals in 2 term and prelim exam-(average of all the internal examination shall be taken).
- 2.3 Marks in decimal computed in 1.1, 1.2 & 1.3 should be converted into whole number at the end.

Minimum marks required by a candidate to be declared as pass will be as follows:

Subject	Theory and		Practical		Internal assessment		Total	
	Max	Dral Min Passing	Max	Min Passing	Max	Min Passing	Max	Min Passing
	95	47	25	13	30	11	150	75
Pathology		47	25	13	30	1.1	150	75
Microbiology	95		25	13	30	11	. 150	75
Pharmacology		47		15	20	7	100	50
FMT	50	25	30	15	20			

#### MICROBIOLOGY

#### 1. THEORY

The computation of internal assessment marks shall be as per rule No 2 and 3 mentioned in this rule and regulation

#### University Examination

- Pattern of Theory Examination including Distribution of Marks, Questions and Time. 2.
  - a. Distribution of Marks

		Total marks
Sr.No	10 la cash)	80
1	Theory ( 2 papers - 40 marks each)	15
2	Oral (Viva)	25
3	Practical	30
4	Internal assessment (Theory -15, Practicals -15)	150
	TOTAL	150

- b. Total duration 4 hrs (each paper of 2 hrs or 120 minutes)
- c. Each paper will have 3 sections.
- d. Pattern and marking for each paper of 40 marks as provided in the table
- e. One compulsory question of 7 marks on applied Microbiology in each paper

Sections	Two Theory Papers	Total No. of Questions	Mark (s) per Question	Total Marks
A)	Multiple Choice Questions	16	1/2	00
B)	(MCQs) Brief Answer Questions (BAQs)	4 out of 5	4	16
/	Long Answer Question (LAQ)	2 out of 3	8	16
C)	Long Answer Question (DAQ) Total			40

**Topic Distribution** 3.

A) MICROBIOLOGY PAPER I:- General Microbiology, Systematic bacteriology including Rickettsia, Chlamydia and Mycoplasma, Related applied microbiology.

B) MICROBIOLOGY PAPER II:- Parasitology, Virology, Mycology, Related applied Microbiology, and Immunology.

-1. University examination Nature of practicals and duration

#### a. Practical examination in MICROBIOLOGY will be of 25 Marks and oral (viva) of 15 Marks of THREE hours duration.

Q.1: Gram staining	5 Marks
Q.2: Zeihl-Neelson's staining	5 Marks
Q.3: Stool examination for Ova/cyst	5 Marks
Q.4: Spots identification (Ten Spots)	10 Marks
	Total-25 Marks

b. Viva (Two tables) 15 Marks

VIVA 1	General Microbiology, Systemic Bacteriology and Applied microbiology	8 Marks
VIVA 2	Parasitology, Virology, Mycology, Immunology.	7 Marks

(*Spots-Bacteriology slide, Culture media, Biochemical, Sterilization and Disinfection, Mycology, Virology, Parasitology, Serological test, Vaccine, Experimental Animal/Vector)

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Final Revised MBBS Syllabus for Microbiology, MGMIHS

(Proposed in BOS, March 2015)

GENERAL MICROBIOLOGY [n=17]

		non/ of oldering	NO. OT NIS.
Topic	Must Know	Desirable to Milow	
History & Microscopy	<ul> <li>Definitions of Medical Microbiology, Pathogen Commensal,</li> </ul>	-	
	Symbiotic.	a	
	<ul> <li>Contribution of</li> </ul>		
	Louis Pasture		JUT
	Robert Koch		
1	Lister		
	<ul> <li>Names of scientists who discovered common bacteria</li> </ul>		
	<ul> <li>Importance of Microbiology</li> </ul>		
Morphology of	<ul> <li>Difference between Prokaryotes &amp; Eukaryotes</li> </ul>		
bacteria l	<ul> <li>Microscopy – Basic principle and applications of all microscopes</li> </ul>		
	<ul> <li>Classification of staining techniques</li> </ul>		
	<ul> <li>Gram's stain and ZN stain in detail( with examples)</li> </ul>		
	<ul> <li>Negative staining, Impregnation method</li> </ul>		2hr
	<ul> <li>Albert's stain</li> </ul>		
	3		1
Morphology II	<ul> <li>Morphology of Bacteria</li> </ul>		
	<ul> <li>Bacterial cell anatomy in detail</li> </ul>		
	Bacterial Spore		
nt	<ul> <li>Bacterial cell division Generation time, Bacterial growth curve</li> </ul>	Bacterial Metabolism	1hr
Physiology of pacteria			
	-		

		đ	
Sterilization	<ul> <li>Definitions of Sterilization, disinfection, asepsis, antiseptics</li> <li>Need of Sterilization / Disinfection in various fields – Medical, Food &amp; Pharma Industry</li> <li>Physical methods of Sterilization</li> <li>Physical methods of Sterilization</li> </ul>		2hrs
Disinfection	<ul> <li>Sunlight, heat (ury without the set of ideal chemical disinfectant</li> <li>Characteristics of ideal chemical disinfectant</li> <li>Factors influencing potency of a disinfectant</li> <li>Disinfectants like Aldehydes , Alcohols, Halogens, Oxidising agents, O</li></ul>	Testing of Disinfectants	1hr
	only) Trynes of Media and their uses	Composition of Media	1hr
Culture Methods	<ul> <li>Types of aerobic culture methods and their uses</li> <li>Types of an aerobic culture methods and their uses</li> </ul>		1hr
ldentification of bacteria	<ul> <li>Mc Intosh Filde's Jar – Functioning and uses Morphology of bacteria (Gram stain), Motility Biochemical tests (Principle and examples)</li> </ul>	Morphology of Bacterial Colony Biochemical tests (Procedure)	1hr
		Typing Method Pathogenicity tests	
Antimicrobial therapy	Mode of action of antimicrobial agents Antibiotic Sensitivity tests : Kirby Bauer disc diffusion Importance of making an antibiotic Policy Strict Adherences to antibiotic Policy	Antibiotic Sensitivity tests: Stokes Disc diffusion E-test Dilution test	Thr
Bacterial Genetics	Basic structure of Bacterial DNA	<ul> <li>Synthesis of Polypeptide (Transcription and</li> </ul>	2hrs

	Definitions of Gene, Codon, Nonsense codons	translation) Genetic Engineering	
	Extrachromosomal elements – Plasmids, Episomes, Transposons	DNA Probes	
	<ul> <li>Difference between Phenotypic &amp; Genotypic variation (Mutation)</li> </ul>	<ul> <li>PCK</li> <li>Gene therapy</li> </ul>	
	<ul> <li>Genetic Variation (Mutation)</li> <li>Gene transfer (Transformation, Transduction, Lysogenic conversion,</li> </ul>		
-	<ul> <li>Conjugation)</li> <li>Differences between Mutational &amp; Transferable drug resistance</li> </ul>		
	<ul> <li>mechanism of drug resistance</li> </ul>		14
Bio- medical waste	<ul> <li>Definition of Biomedical waste</li> <li>Classification &amp; disposal as per categories</li> </ul>		Tur
disposal	<ul> <li>Importance of segregation</li> </ul>		
	<ul> <li>Universal safety precautions</li> </ul>		
	<ul> <li>Hand Hygiene</li> <li>Hand Hygiene</li> </ul>		1hr
Universal Safety	<ul> <li>Definition of Hospital acquired intection.</li> </ul>		
Precautions & health	<ul> <li>Sources, types, prevention and control of fiealth care associated</li> </ul>		
care associated	infections	5	
infections			
Normal microbial flora	Introduction – Various sites, types & role		1hr
of human body			
		Factors Predisposing to	
Infection	<ul> <li>Definitions of Saprophytes, Parasitic, Commensals, Paulogen, Concernington Pathogens, Pathogenicity, Virulence</li> </ul>	Microbial Pathogenicity	
Host parasite	<ul> <li>Types of infection, Routes of transmission</li> </ul>		1hr
relationship	<ul> <li>Sources of Infection</li> </ul>		
	<ul> <li>Difference between Exotoxins &amp; Endotoxins</li> </ul>		
	<ul> <li>Types of Infectious diseases Localised, Generalised, Endening,</li> </ul>		
	Epidemic, Panaetitic		

		Desirable to Know No. of Hrs	No. of Hrs
Topic	Must Know		
•	Subtopics	D	Thr
Staphylococci	A,B,C,,E,F,G,H,I	D	Thr
Streptococci	A,B,C,,E,F,G,H,I	0	Thr
Other streptococci and	A,B,C,,E,F,G,H,I	2	
Pneumococci	4		

SYSTEMIC BACTERIOLOGY [n=29]

Formaț of study -

A :Classification

B; Morphology

C:Culture Characteristics

D: Biochemical reactions

E:Antigens

F:Pathogenesis & diseases caused in detail

G:Laboratory diagnosis

H:Prevention & control

I: Special identification tests

		0	1hr
Neisseria	A,B,C,,E,F,G,H,I		1hr
C. diptheriae	A,B,C,,E,F,G,H,I	0	1hr
M. tuberculosis	A,B,C,,E,F,G,H,I		1hr
Atypical mycobacteria	Name of the Species Names of the diseases caused Special tests for identification		
M. leprae	Brief outline of lab diagnosis A,B,C,,E,F,G,H,I	Δ	1hr
Bacillus	Name of the Species		1hr
	Names of the diseases caused Special tests for identification		
	Brief outline of lab diagnosis		1hr
Method of anaerobiasis &	Method of anaerobiasis Nonsporing anaerobes { Name of the Species, Names of the diseases caused		
	Special tests for identification, Brief outline of lab ulagroups J A,B,C,,E,F,G,H,I	G	Ţhr
Closefeidium – II	A,B,C,,E,F,G,H,I	Ω	1hr
	A B.C.E.F.G.H,I		1hr
( E.coli. )		D	Thr
Enterobacteriacae – Il proteus & Klebsiella	A,B,C,,E,F,G,H,I		
	5	_	

X.	 1hr	1hr	1hr		1hr	1hr		Thr		Thr		1hr	1hr	
					٥									
	A,B,C,,E,F,G,H, ¹	A,B,C,,E,F,G,H,I	A,B,C,,E,F,G,H,I	Name of the Species Names of the diseases caused Special tests for identification	Brief outline of lab diagnosis	А,В,С,Е,Е,Е,Б,Н,І	Name of the Species Names of the diseases caused	breed outline of lab diagnosis	Name of the Species Names of the diseases caused Special tests for identification	Brief outline of lab diagnosis	Name of the Species Names of the diseases caused	Special tests for lucitudes and Brief outline of lab diagnosis	A,B,C,,E,F,G,H,I	A,B,C,,E,F,G,H,I
	Enterobacteriacae – III A,B Salmonella		Vibrio A,B	Campylobacter & Na Helicobacter		Pseudomonas A,I		Francisella, Bordetella) Br	Other GNB II ( Haemophilus, Brucella)		Miscellaneous Bacteria N (Newer bacterias)		Spirochaete - I	Spirochaete -II

# MYCOLOGY [n=4]

		d. Mucor, Aspergillus	
		c. Candida, Cryptococcus in detail	
		b. Predisposing factors	infections & P. Carinii
	p Carinii	a. Enumerate agents	opportunistic fungal
TUL	Histoplasma,	Systemic & Opportunistic Mycosis	Systemic mycosis &
	-	c. Lab diagnosis (Outline)	
		b. Predisposing factors	
		a. Enumerate agents	Candida
Tur		<ul> <li>Subcutaneous Mycosis &amp; Candida in detail</li> </ul>	Subcutaneous mycosis &
		c. Lab diagnosis (Outline)	
	dermatophytes	b. Predisposing factors	
	characteristics of	a. Enumerate agents	mycosis (Dermatophytes)
TUL		<ul> <li>Superficial Mycosis</li> </ul>	Agents of superficial
		Classification of Fungal diseases	
		<ul> <li>Lab diagnosis of mycosis</li> </ul>	
	3	<ul> <li>Reproduction &amp; Sporulation</li> </ul>	
		<ul> <li>Classification of fungi with examples</li> </ul>	
		<ul> <li>Difference between fungus &amp; Bacteria</li> </ul>	
TU	3	<ul> <li>Introduction to Mycology</li> </ul>	Introduction to mycology
		Subtopics	
	Know		
No. of Hrs	Desirable to	Must Know	Topic

							Ş	& Milk	Barteriology of water, Air			Mycoplasma			Chlamydia				Rickettsia		Actinuinycere		
milk	Examination of	Bacteriological	diseases	Milk Borne	water	Examination of	Acceptable littlic of Americal Bacteriological		Bacteriology of Air Water Borne	Bilei ouring - water water	Special courtine of lab diagnosis Bacterial flora in	 Name of the spore	Bile curine	cherial tests for identification	Names of the diseases caused	Name of the Species	Brief outline of lab diagnosis	Special tests for identification 1hr	Names of the diseases caused	Name of the Species		A,B,C,,E,F,G,H,I 1hr	1hr
																						-	

12	11	10	60	80	07	
Arboviruses	Hepatitis viruses	Picornaviruses	Paramyxoviruses	Orthomyxoviruses	Other DNA viruses (Papova, Adeno, )	
<ul> <li>Classification, Names of Arboviruses and diseases</li> <li>caused</li> <li>Dengue Virus in detail</li> </ul>	<ul> <li>HAV (Pathogenesis and Lab diagnosis)</li> <li>HBV (Morphology, Mode of transmission, Clinical features, Lab diagnosis)</li> <li>HCV (Morphology, Mode of transmission, Clinical features, Lab diagnosis)</li> <li>HDV &amp; HEV (Pathogenesis &amp; Lab diagnosis)</li> </ul>	<ul> <li>Classification</li> <li>Polio virus in detail</li> <li>Differences between killed and live vaccines</li> <li>Eradication and Prophylaxis of Polio virus</li> </ul>	<ul> <li>Morphology</li> <li>Measles virus and Mump Virus</li> </ul>	<ul> <li>Differences between Orthomyxo and paramyxo virus</li> <li>Influenza Virus <ul> <li>Morphology</li> <li>Antigenic classification and structure</li> <li>Antigenic shift and Antigenic drift</li> <li>Pathogenesis and lab diagnosis</li> </ul> </li> </ul>	Basic morphology, diseases caused	diagnosis)
JE yellow tever new		<ul> <li>Coxsackie viruses</li> <li>Rhino virus</li> </ul>	Parainfluenza virus     RSV	Influenza Virus Antigenic classification and structure Prophylaxis Bird flu		
1			1hr	1 hr	1hr	

VIROLOGY [n=16]

	<ul> <li>CMV (Infections caused and Lab diagnosis)</li> </ul>	<ul> <li>Morphowey</li> <li>Classification</li> <li>HSV (Infections caused and Lab diagnosis)</li> <li>Varicella – Zoster (Infections caused and Lab</li> </ul>	Herpes simplex & Varicella zoster CMV, EBV	06
111	<ul> <li>EBV (Infections caused and</li> </ul>			
Thr	-	<ul> <li>Names of poxviruses and diseases causes</li> <li>Bacteriophage[Basic structure and Significance]</li> </ul>	Bacteriophage	C
IUT	Cultivation	Morphology	Pox viruses	D _n
	diseases			
	Chemotherapy of viral	<ul> <li>a. Types and Schedule</li> <li>list of antiviral agents.</li> </ul>	antiviral agents	
	Chemoprophylaxis &	Commonly used viral vaccines	Viral vaccines and	04
TUL	Mode of preparation	<ul> <li>Immunity in viral diseases</li> </ul>		
		<ul> <li>Interferons</li> </ul>	111/61 404010	
	infections	<ul> <li>Inclusion bodies</li> <li>Routes of transmission of viral infections</li> </ul>	Virus-host	03
7111	<ul> <li>Host responses to virus</li> </ul>	Outline of diagnosis of vital discussion		1
Jhr.		Cultivation of Viruses ,Viral assays	General virology - Il	02
1hr	• VII di nacilia66.	Replication of viruses		
2	and chemical agents			
	<ul> <li>Susceptibility to physical</li> </ul>	<ul> <li>Morphology of Viruses</li> </ul>	General Virology -	01
				NO
1hr	<ul> <li>Chemical properties of</li> </ul>	Subtopics	Tobic	DIC
		Must Know		-
No.of nrs	Desirable to Know			

13	Rhabdoviruses	<ul> <li>Iviorphology</li> </ul>		Thr
		<ul> <li>Pathogenesis and Lab diagnosis</li> </ul>		
		<ul> <li>Prophylaxis</li> </ul>		
14	Retro Viruses – HIV	<ul> <li>Morphology</li> </ul>	<ul> <li>Viral genes &amp; antigens</li> </ul>	1hr
		<ul> <li>Resistance</li> </ul>	• ART	
		<ul> <li>Modes of Transmission</li> </ul>		
		<ul> <li>Pathogenesis</li> </ul>		
		<ul> <li>Opportunistic infections and malignancies</li> </ul>	5	
		<ul> <li>Lab diagnosis in detail</li> </ul>		
		<ul> <li>Prevention</li> </ul>		
		• PEP		5
15	Miscellaneous	<ul> <li>Viruses (Only names) causing gastroenteritis</li> </ul>		1hr
	viruses	<ul> <li>Viruses causing viral hemorrhagic fevers (only</li> </ul>		
		names)		
		<ul> <li>Slow virus diseases (Only names)</li> </ul>		
16	Oncogenic viruses	Papilloma Virus		1hr
		Only names of oncogenic viruses and malignancies		
		caused		

		t	- 5 U vo
		Desirable to Know	611 10 .ON
	Must Know		
Topic			
	Subtopics		1hr
	T Loc Contollo	1	
	<ul> <li>Innate Immunity - Iypes, raccord</li> </ul>		
Immunity	influencing innate immunity,		
	Mechanisms		
	<ul> <li>Acquire Immunity -</li> </ul>		
	a. Active Immunity		
	b. Passive Immunity		
	<ul> <li>Combined immunization</li> </ul>		
	<ul> <li>Adoptive immunity</li> </ul>		
	- Incal Immunity		
	uord Immunity	cer antigens	Thr
	The second		
	• Iypes of contracting antigenicity		
Antigen	· Factors allecons anecons	Tinc IoM. IgA, IgD,	1117
	<ul> <li>properties of antibodies</li> </ul>	10L 10L 10L 1	
Antibody	<ul> <li>Structure of Immunoglobulin</li> </ul>	Mol. Wt., Sed.	
	classes		
	a leG. IgM, IgA, IgD, IgE	Coefficient,	
		. Abnormal	
	a) Basic structure, function	Immunoglobulins	
	& distribution	" Regulation of	Inr
	<ul> <li>Components of complement</li> </ul>	complement	
Complement	<ul> <li>Classical Pathway</li> </ul>	activation	
	<ul> <li>Alternative Pathway</li> </ul>		
	12		

[11=n] YOOLOGY [n=11]

	" Biological offects of complement	<ul> <li>Biosynthesis of</li> </ul>	
		Complement	
		<ul> <li>Quantitation of</li> </ul>	
		complement	
Δα-Δh reactions I	<ul> <li>Types of Ag – Ab reactions,</li> </ul>		2hrs
	precipitation, Agglutination, CFT,		15
	Neutralization, Opsonisation,		
	Immunoflourescence, ELISA		
	Immunochromatography	23	
	(Principle, Types & uses only)		
Structure & function of Immune system	<ul> <li>Central Lymphoid Organs</li> </ul>	<ul> <li>Lymphocytic</li> </ul>	Ihr
	<ul> <li>Peripheral Lymphoid Organs</li> </ul>	recirculation	
	<ul> <li>Cells of Lymphoreticular System</li> </ul>		
	<ul> <li>HLA</li> </ul>		
	<ul> <li>Differences between T &amp; B cells</li> </ul>		
	Human Response	<ul> <li>Cell mediated</li> </ul>	Thr
Immune response		Immine Recoonse	
	a) Primary and secondary		
	responses	a) Detection of	
2	b) Production of Antibodies	CMI	
	c) Monoclonal antibodies	<ul> <li>Immunological</li> </ul>	
	d) Factors influencing antibody	tolerance	
	production		
	<ul> <li>Cell mediated Immune Response</li> </ul>		
	a) Cytokines & Lymphokines –		
	Types & functions only	-	1.
Hvnersensitivity	<ul> <li>Definition &amp; Classification</li> </ul>	3	Ihr
	<ul> <li>Type 1 Reaction</li> </ul>		
	<ul> <li>Differences between Immediate &amp;</li> </ul>		
	delayed hypersensitivity		
	<ul> <li>Type 2, 3 &amp; 4 Reactions</li> </ul>		

artion & Tumor immunity	<ul> <li>Primary Immunodeficiency Syndromes &amp; Secondary Immunodeficiency (Only classification)</li> <li>(Only classification)</li> <li>(Only classification)</li> <li>Immunosurveillance</li> <li>Immunosurveillance</li> <li>Immunosurveillance</li> <li>Immunosurveillance</li> <li>Immunosurveillance</li> <li>Immunosurveillance</li> </ul>		Contraction & Mechanisms	Thr	
rimmunodeficiency hmmunodeficiency syndromes & Secondary hmmunodeficiency (Only classification) e Immunosurveillance histocompatibility Antigens Histocompatibility Testing e Graft - Versus – Host reaction e Tumor antigens histocompatibility Testing e Tumor antigens	<ul> <li>Primary Immunodeficiency Syndromes &amp; Secondary Immunodeficiency (Only classification)</li> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Timmunodeficiency (Only classification)</li> <li>Allograft reaction</li> <li>Immunorveillance</li> <li>Immunorveillance</li> <li>Immunorveillance</li> <li>Immunorveillance</li> <li>Immunorveillance</li> <li>Immunorveillance</li> <li>Types of Transplants</li> <li>Tumorartifity Testing</li> <li>Tumor antigens</li> <li>Tumor antigens</li> </ul>	nunity	<ul> <li>Definition &amp; monoconstruction</li> <li>Classification with examples</li> </ul>		
r immunity Types of Transplants & Secondary Immunodeficiency Syndromes & Secondary Immunodeficiency (Only classification) (Only classification) & Allograft reaction & Immunosurveillance & Immunosurv	rimmunity Types of Transplants & Secondary Immunodeficiency Syndromes & Secondary Immunodeficiency (Only classification) (Only classification) & Allograft reaction & Immunosurveillance & Immunosurve			<ul> <li>Primary</li> </ul>	
<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>	<ul> <li>Types of Transplants</li> <li>Types of Types of Ty</li></ul>	odeficiency diseases		Immunodeficiency sundromes &	
<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>	<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>			e Secondary	
<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus – Host reaction</li> <li>Tumor antigens</li> </ul>	<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>			Immunodeficiency	
<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>	<ul> <li>Types of Transplants</li> <li>Types of Transplants</li> <li>Allograft reaction</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>				
<ul> <li>Iypes of itemplation</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>	<ul> <li>Iypes of itensprend</li> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>				
<ul> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>	<ul> <li>Allograft reaction</li> <li>Histocompatibility Antigens</li> <li>Histocompatibility Testing</li> <li>Graft - Versus - Host reaction</li> <li>Tumor antigens</li> </ul>	splantation & Tumor immunity		<ul> <li>Immune response to</li> </ul>	
y Testing Aost reaction	y Testing Aost reaction		<ul> <li>Allograft reaction</li> <li>Allograft reaction</li> </ul>	malignancy	
iy resume Host reaction	y teaction Host reaction		<ul> <li>Histocompatibility Anticedia</li> </ul>	<ul> <li>Immunotherapy of</li> </ul>	
lost reaction	lost reaction		<ul> <li>Histocompatibility lesuits</li> </ul>	cancer	
• Tumor antigens	<ul> <li>Tumor antigens</li> </ul>		<ul> <li>Graft - Versus - Host reaction</li> </ul>		
			<ul> <li>Tumor antigens</li> </ul>	•	

PARASITOLOGY [n=10]

No. of Hrs. 1hr 1hr 1hr Thr Thr & Trypanosoma Cruzi Trypanosoma brucci Desirable to Know in details Morpho, Lifecycle, Pathogenesis, Lab diagnosis Leishmania (Classification, diseases caused)
 L. donovani in details Life Cycle, Morphology, Pathogenecity & Lab Pathogenesis, Lab diagnosis & treatment) Pathogenesis, Lab diagnosis & treatment Giardia lamblia (Morphology, life Cycle, Trichomonas vaginalis (Morphology , Hosts (Definitive & Intermediate) Pathogenesis & Complications . Host - Parasitic relationship Lab diagnosis & treatment Non Pathogenic amoebae Classification of Parasites Lab diagnosis in general Free living amoebae diagnosis, prevention Sources of Infection Type of Parasites Morphology Life cycle Must Know Subtopics Introduction to parasitology Giardia, Trichomonas Haemoflagellates E.Histolytica Malaria Topic Topic 02 S 4 No m 01

Miscellaneous protozoa     Toxoplasma (Morphology life Cycle, Pathogenesis, Lab diagnosis)       Cestodes     Taenia & Echinococcus (Morphology life Cycle, Pathogenesity, Lab diagnosis)       Trematodes     Taenia & Echinococcus (Morphology life Cycle, Pathogenesity, Lab diagnosis)       Nematodes     Trematodes       Nematodes     - Schistosomes a. Names & diseases caused a. Names & diseases caused       Nematodes ( Intestinal)     - A. duodenale, A. lumbricoides, E. vermicularis, T. trichura (in details)       IO     Tissue Nematodes I       Trissue Nematodes I     W. bancrofti ( in details)       10     Tissue Nematodes I       8< stool concentration     • D. medinensis ( in details)       11     Tissue Nematodes II       8< stool concentration     • Names of parasites affecting liver ( Names of parasites effecting liver)				Cryptosporidium,	Thr
Image: Contract of the second seco	9	Miscellaneous protozoa	Toxoplasma (Morphology life Cycle,	Isospora	
Taenia & Echinococcus (Morphology life Cycle, Pathogenesity, Lab diagnosis)     Brifer mention about.       Trematodes     Taenia & Echinococcus (Morphology life Cycle, Pathogenesity, Lab diagnosis)     Morphology, Life       Trematodes     Antono about.     1       Trematodes     Schistosomes     Schistosomes       Normatodes     Schistosomes     Schistosomes       Names & diseases caused     Parogenicity       Nematodes     A. duodenale, A. lumbricoides, westermani       Nematodes (Intestinal)     A. duodenale, A. lumbricoides, westermani       Nematodes I     Tr.spue Nematodes I       Tissue Nematodes I     Tr.spiralis       S. stercoralis     S. stercoralis       A. duodenale, A. lumbricoides, techniques     S. stercoralis       B. Nematodes I     M. bancrofti (in details)       T. Spiralis     S. stercoralis       S. stercoralis     S. stercoralis       R. Stool concentration     Numes of parasites in stool       R. Stool concentration     Names of parasites file of files       R. Stool concentration     Names of parasites in stool       R. Stool concentration     Names of parasites file of folore in saturated safe solution which float in			Tatilogenesis, ter and	B.coli	
Cestodes         Taenia & connoccuo (non-proces)         othercestodes           Pathogenesity, Lab diagnosis)         Morphology, Life         1           Trematodes         Schistosomes         R.Lab diagnosis         1           Schistosomes         a. Names & diseases caused         Parognimus         1           Morphology, Life         A. duodenale, A. lumbricoides,         S. stercoralis         1           Mematodes ( Intestinal)         A. duodenale, A. lumbricoides,         S. stercoralis         1           Morphology ( Intestinal)         C. vermicularis, T. trichura (in details)         S. stercoralis         1           Morpholos ( Intestinal)         C. vermicularis, T. trichura (in details)         S. stercoralis         1           Morpholos ( Intestinal)         C. vermicularis, T. trichura (in details)         S. stercoralis         1           Intestina         W. bancroft ( in details)         S. stercoralis         1         1           Intestina         W. bancroft ( in details)         S. stercoralis         1         1           Intestina         W. bancroft ( in details)         S. stercoralis         1         1           Intestodes II         P. D. medinensis ( in details)         S. stercoralis         1         1           Intestolutiones         N				Brief mention about.	1hr
Trematodes     Morphology, Life     1       Trematodes     Trematodes     Morphology, Life     1       Trematodes     • Schistosomes     Rasciola hepatica       • Schistosomes     • Schistosomes     Parognimus       • Names & diseases caused     Parognimus       • Nematodes (Intestinal)     • A. duodenale, A. lumbricoides,     S. stercoralis       • Nematodes (Intestinal)     • A. duodenale, A. lumbricoides,     S. stercoralis       • Nematodes I     W. bancrofti (in details)     S. stercoralis       • Stool concentration     • Numention details)     S. stercoralis       • Stool concentration     • Names of parasites affecting liver     Numentiones       • Rechniques     • Names of parasites affecting liver     • Names of parasites affecting liver       • Names of parasites affecting liver     • Names of parasites affecting liver       • Names of parasites affecting liver     • Names of parasites affecting liver       • Names of parasites affecting liver     • Names of parasites affecting liver       • Names of parasites affecting liver     • Names of parasites affecting liver		Cestodes	Taenia & Echinococcus (workproved a pathogenesity, Lab diagnosis)	othercestodes	
TrematodesTrematodesCycle, Pathogenicity (a. Lab diagnosis a. Names & diseases caused a. Names & diseases caused a. Names & diseases caused b. Names & diseases caused b. A. duodenale, A. lumbricoides, westermaniFasciola hepatica parognimus westermani0Nematodes (Intestinal) Tissue Nematodes I Tissue Nematodes I• A. duodenale, A. lumbricoides, westermaniS. stercoralis s. stercoralis10Tissue Nematodes I Tissue Nematodes IW. bancrofti ( in details)S. stercoralis s. stercoralisS. stercoralis s. stercoralis11Tissue Nematodes II & S. tool concentration techniques techniques• D. medinensis ( in details) s. steol concentration techniques which float in e. Names of parasites affecting fiver heren is the olution & names of parasites affecting fiver saturated salt solution & those which float in saturated salt solution & those which float in				Morphology, Life	1hr
<ul> <li>Schistosomes</li> <li>Schistosomes</li> <li>Schistosomes</li> <li>Schistosomes</li> <li>Schistosomes</li> <li>Names &amp; diseases caused</li> <li>Parognimus</li> <li>A. duodenale, A. lumbricoides, westermani</li> <li>M. bancrofti (in details)</li> <li>Tissue Nematodes I</li> <li>T. Spiralis</li> <li>Tissue Nematodes II</li> <li>S. stercoralis</li> <li>S. ste</li></ul>	00	Trematodes		Cycle, Pathogenicity	
a. Schistosomes       Fasciola hepatica         a. Names & diseases caused       Parognimus         b. Namatodes ( Intestinal)       A. duodenale, A. lumbricoides, westermani         b. Mematodes ( Intestinal)       A. duodenale, A. lumbricoides, mestermani         construction       A. duodenale, A. lumbricoides, mestermani         b. Mematodes ( Intestinal)       F. vermicularis, T. trichura (in details)         construction       W. bancrofti ( in details)         construction       W. bancrofti ( in details)         construction       M. bancrofti ( in details)         construction       W. bancrofti ( in details)         construction       N. bancrofti ( in details)         construction       N. bancrofti ( in details)         constructi       N. bancrofti ( in details)				& Lab diagnosis	
a. Names & diseases caused       Parognimus         b       A. duodenale, A. lumbricoides,       parognimus         Nematodes (Intestinal)       • A. duodenale, A. lumbricoides,       S. stercoralis         0       Tissue Nematodes I       W. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes I       W. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes I       W. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes I       W. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes I       W. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       W. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       N. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       N. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       N. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       N. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       N. bancrofti (in details)       S. stercoralis         1       Tissue Nematodes II       • D. medinensis (in details)       S. stercoralis				Fasciola hepatica	
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Nematodes ( Intestinal)       A. duodenale, A. lumbricoides,       S. stercordins         Nematodes ( Intestinal)       E. vermicularis, T. trichura (in details)       S. stercordins         Tissue Nematodes I       W. bancrofti ( in details)       S. stercordins         Tissue Nematodes I       W. bancrofti ( in details)       S. stercordins         Tissue Nematodes I       W. bancrofti ( in details)       S. stercordins         Tissue Nematodes II       W. bancrofti ( in details)       S. stercordins         Tissue Nematodes II       • D. medinensis ( in details)       S. stercordins         Tissue Nematodes II       • D. medinensis ( in details)       S. storol concentration techniques         R Stool concentration       • Names of parasites affecting CNS       Names of parasites affecting liver         • Names of parasites affecting liver       • Names of parasites entering through skin       Names of parasites entering through skin         • Bile stained eggs& Eggs which float in       • Bile stained eggs& regs which float in       Saturated salt solution & those which do not					1hr
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Tissue Nematodes II <ul> <li>D. medinensis ( in details)</li> <li>Stool concentration techniques</li> <li>stool concentration</li> <li>Name of parasites in stool</li> <li>Names of parasites affecting CNS</li> <li>Names of parasites entering through skin</li> <li>Bile stained eggs&amp; Eggs which float in</li> <li>saturated salt solution &amp; those which do not</li> </ul>	10	Tissue Nematodes I	W. bancrofti ( in details)		
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Tissue Nematodes II & Stool concentration techniques			- details)		Thr
3 3 3 3 3	11	Tissue Nematodes II	<ul> <li>D. medinensis (m) documents</li> <li>stool concentration techniques</li> </ul>		
9983		& Stool concentration	<ul> <li>Name of parasites in stool</li> </ul>		
		techniques			
saturated sati societion					
			saturated sail solution &		

APPLIED IVIICROBIOLOGY (To be taken in the form of UG seminars/Tutorials)

(n=8)

		No. of Hrs.
Topic	Topic	
No	-	
	Only Causative agents & Brief Outline of Lab diagnosis in	
		1hr
Н	<ul> <li>Gastrointestinal infections</li> </ul>	
		1hr .
2	• URTI	
		1hr
m	• LRTI	
		1hr
4	• UTI	
		1hr
S	CNS Infections	
		Ihr
9	<ul> <li>Wound &amp; Pyogenic infections</li> </ul>	
		Thr
2	<ul> <li>PUO &amp; infections</li> </ul>	
		1hr
∞	• STDs	

Theory: (n=96)

No of lectures	17	29	. 04	16	11	11	08	96	
Section	General Microbiology	Systemic bacteriology	Mycology	Virology	Immunology	Parasitology	Seminars/Tutorials on Applied Microbiology	Total	
100	170								
-----	-----								
1	1								
1 .	=								
10	dis								
	TIC								
-	a								
2	7								

11	Eventiments	No.of Hrs
ON	Experimence	
H	Microscopy	4hrs 5
2.	Morphology of bacteria	4hrs
m.	Sterilisation and Disinfection	4hrs
4.	Principles in diagnostic Microbiology 1	4hrs
5.	Principles in diagnostic Microbiology 2	4hrs
6.	Immunology and Serologigal methods	4hrs
7.	Staphylococci	4hrs
∞.	Streptococci and Pneumococci	4hrs
9.	Neisseria	4hrs
10.	Corynebacteria	4hrs
11.	Bacillus	4hrs
12.	M.tuberculosis and Atypical Mycobacteria	4hrs
13.	M.leprae	4hrs
14.	E.coli,Klebsiella and Proteus	4hrs
15.	Salmonella	4hrs
16.	Shigella and Vibrio	4hrs
17.	Pseudomonas and Hospital infections	4hrs
18.	Yersinia and Brucella	4hrs
19.	Haemophillus and Bordetella	4hrs
20.	Clostridia	4hrs

4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	4hrs	132Hrs	
	Non-sporing anaerobes	Spirochaetes	Actinomycetes and Nocardia	Virology	Intestinal protozoa	Blood and tissue protozoa	Blood and tissue flagellates	Cestodes	Trematodes	Intestinal nematodes	Tissue nematodes	Medical Entamology	Mycology	TOTAL
	21.	22.	23.	24.	25.	26.	27.	28.	29.	30.	31.	32.	33.	

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(As per MCI)
1SF
2
s: 250 hours (
2501
urs:
PHO
Teaching Hours:
ach
Te
57
Tot

96Hrs	132Hrs	22Hrs		250Hrs	20
	Lectures + Seminars/Tutoriais	Practicals	Assessments		Total

3.

Books Recommended :

Sr.	Name of the Book	Author
No.		R Ananthanaravan
Ч	Textbook of Microbiology	C K Jayaram Panikar
c	A Textbook of Microbiology	P. Chakraborty
4 0	Touthook of Medical Microbiology	Rajesh Bhatia & Itchpujani
n		Prof C.P. Baveia
4	Textbook of Medical Milci University	
L.	Textbook of Medical Parasitology	C K Jayaram Panikar
,		C P. Baveia
9	Medical Parasitology	V.Baveja
7	Textbook of Medical Parasitology	S C Parija
00	Textbook of Parasitology	Damle and Karyakarte
~	A Textbook of Parasitology	Dr.K.D. Chatterjee.
σ	Practical Microbiology	Dr. Anuradha De

Refe	Reference Books :	
Sr. No.	Name of the Book	Author
7	Mackie McCartney practical Medical	Colle JG, Fraser AG
-1	Microbiology	
2	Principles of Bacteriology, Virology &	Topley Wilsons
0	Medical Mycology (Emmons)	Kwon – Chung
n .	Review of Medical Microbiology	Jawetz
4	(lange)	
L		Weir DM
ñ		David Greenwood, Richard Stack, John
Q	Medical Microbiology	pentherer
r	Modical Mirology	Timbury MC
-		
0	Mackie McCartney Medical	Duguid JP
0	Microbiology Vol.1	······
<u>ہ</u>	Textbook of Microbiology	Monica Uneeseniousi

e

Books:

J

Evaluation

a. Methods

Theory, Practical & Viva

No.		Particulars	Total Marks
	Theory (Total out of 95 Marks) Theory (2 Papers – 40 Marks each)	Theory ( 2 Papers – 40 Marks each)	80 Marks
		Oral (Viva)	15 Marks
7	Practical (Total out of 25 Marks)	Practical	25 Marks
m	Internal Assessment (Total out Internal Assessment (Theory – of 30 Marks) 15 Practical – 15)	Internal Assessment (Theory – 15 Practical – 15)	30 Marks
		TOTAL	TOTAL 150 Marks
	and the second se		

Passing: A candidate has to obtain minimum of 47 Marks out of 95 in Theory, 13 marks out of 25 in Practical, 11 marks out of 30 in Internal Assessment and 75marks out of 150 Total to be declared as passed.

Nature of Question Paper :- Theory (Total 80 Marks)

Paper - I	General Microbiology , Systemic Bacteriology & Related Applied Microbiology	40 Marks
Paper – II	Immunology, Virology, Parasitology , Mycology & 40 Marks Related Applied Microbiology	40 Marks

	Currention Description	Division of Marks Total Marks	Total Marks
Section	Cression construction		oo narie
	MCOS (16)	16 x 0.5 Marks	CN INIGINO
A			
	Brief Answer Questions	4 x 4 Marks	CV INIAI OT
D.	(4/5)		
	Long Answer Questions	2 x 8 Marks	16 Marks
U	(2/3)		
			40 Marks

Practical Examination Marks distribution : -

		Marks
NO.	No. Particulars	
	2	5 Miarks
1	Grams Staining	5 Marks
2	ZN Staining	
		CV IPIAI C
m	Stool Examination	10 Marks
	C==== (10)	
4		25 Marks
	TOTAL	

Viva (Two Tables)

	*				
 8 Marks		7 Marks		15 Marks	
General Microbiology, Systemic	Bacteriology and Applieu Microsco	Parasitology, Virology , Wycology,	Immunology	Tota	10141
<	I		â		

# Distribution of MCQs:

PAPER 1: 16 MCQs , Marks 0.5 each= 8Marks

Microbiology 06 MCQs	c Bacteriology 10MCQs	16 MCQs	
General Microbiology	Systemic Bacteriology	Total	

4.

PAPER 2: 16 MCQs , Marks 0.5 each= 8Marks

Parasitology	05MCQs
Mycology	03MCQs
Virology	04MCQs
mmunology	04MCQs
Total	16 MCQs

Internal Assessment shall be computed on the basis of three term ending examinations (Two Terminals & One Preliminary

evamination).

	No. of Papers	Pattern	Duration of each paper	Total Marks
Examination				
		MCQs = 16 (8 Marks)		
1 st Terminal	1 (40 Marks)	BAQs = 4/5 (16 Marks)	2 Hours	40 Marks
		LAQs = 2/3 (16 Marks)		
		-		
		MCQs = 16 (8 Marks)		
n nd Terminal	1 (40 Marks)	BAQs = 4/5 (16 Marks)	2 Hours	40 Marks
		LAQs = 2/3 (16 Marks)		
		MCQs = 16 (8 Marks)		
Prelim 2	2 (40 Marks each)	BAQs = 4/5 (16 Marks)	2 Hours each paper	80 Marks
-		LAQs = 2/3 (16 Marks)		

南

Do. A. Softwark, M.D. Prof. & HOD Microbiology MOM Medical College & Hospital Kamolhe, Navi Mumbai–410209.

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Dr. Anchrist, Bhesania Houiwala MBBS M.D. Reen. No. 2000/05/1733 (BDAGSSO: Jept. of Microbiology, MGM. Medical College & Hospital, Navi Mumbal, Kamothe



Approved in BOM 40/215, dated 13/03/2015, Resolution No. 3.2 (c)

Approved in BOM 40/215, dated 13/03/2015, Resolution No. 3.2 (c)



Resolution No. 3.2 (J)

Resolution No. 3.2(d): Resolved to accept the following distribution of MCQ for UG examination so as to cover the syllabus properly :

.

MCQ Paper I 16 MCQ -

General Microbiology 06 MCQ Marks 0.5 each = 08 Systemic Bacteriology 10 MCQ

Paper II 16 MCQ

Parasitology - 5 MCQ Mycology - 3 MCQ Virology - 4 MCQ Immunology - 4 MCQ Approved in Bom 43/2015, Dated 06/11/2015-Resolution No. - 3.2 (a)

HHE

ANNEXURG -IV

# Photo 1: Life Cycle of E.histolytica



[116]

# Photo 2: Life Cycle of Giardia lamblia



#### Photo 3: Life Cycle of Plasmodium



[118]

# Photo 4: Life Cycle of Leishmania donovani



#### Photo 5: Life Cycle of Taenia



[120]





[121]



[122]



# Photo 8: Life Cycle of Ascaris lumbricoides

[123]

Photo 9: Life Cycle of Ancylostoma duodenale



[724]

## Photo 10: Life Cycle of Trichuris trichura



[577]



[126]



# Photo 12: Life Cycle of Strongyloides stercoralis

[127]





[128]



# Photo 14: Life Cycle of Dracunculus medinensis

[129]





#### CONTENTS

	IIIrd Semest			
lo l	Experiments	Page	Date	Sign
1	Microscopy	1		
2	Morphology of bacteria	4		
3	Sterilisation and disinfection	7		
4	Principles of Diagnostic Microbiology-I	11		
5	Principles of Diagnostic Microbiology-II	16		
6	Immunological and Serological methods	21		
	IVth Semes			
No	Experiments	Page	Date	Sign
1	Staphylococci	27		
2	Streptococci	30		
3	Neisseria	33		
4	Corynebacteria	35		
5	Bacillus	38		
6	M.tuberculosis and Atypical Mycobacteria	40		
7	M.leprae	44		
8	E.coli, Klebsiella, Proteus	46		
9	Salmonella	49		
10	Shigella and Vibrio	53		
11	Pseudomonas and Hospital Infections	58		
12	Yersinia and Brucella	61		
13	Hemophilus and Bordetella	64		
14	Clostridia	66		
15	Non sporing anaerobes	70		
16	Spirochaetes	71		
17		76		
18		78		
10	Vth Seme	ester		
No	Experiments	Page	Date	Sign
1	Intestinal protozoa	82		
2		88		
3		92		
4		95		
5		98		
6		100		
7		106		
8		109		
5	A second s	112		

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1	Life cycle of E.histolytica	116
2	Life cycle of G.lamblia	117
3	Life cycle of Plasmodium	118
4	Life cycle of L.donovani	119
5	Life cycle of Taenia	120
6	Life cycle of E.granulosus	121
7	Life cycle of Schistosomes	122
8	Life cycle of A. humbricoides	123
9	Life cycle of A.duodenale	124
10	Life cycle of T.trichura	125
11	Life cycle of E.vermicularis	126
12	Life cycle of S.stercoralis	127
13	Life cycle of W.bancrofti	128
14	Life cycle of D.medinensis	129

Page 15 and 16 which is showing coloured pictures related to Microbiology are to be printed as backside of cover page and frontside of last page respectively ( in colour).

#### Ref: MGM/Micro/2016/O-

Date: 15.01.2016

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To, The Registrar MGM Institute of Health Sciences KAmothe, Navi Mumbai.

Subject : Regarding correction in the II-MBBS syllabus

Respected Sir,

Herewith forwarding the text of some changes to be done in making system in University Examination, as suggested by Dr. Shroff, Dr. Deshmukh & Dr. Bhalchandra for needful.

Thanking you,

Yours Sincerely,

AA

Dr. A.D. Urhekar Professor & Head Department of Microbiology **Dr. A.D. Urhekar, M.D.** Professor & Head Department of Microbiology MGM Medical College & Hospital Kamothe, Navi Mumbai-410209.

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COE. for inflomentation & immediate estat-



MGM Institute Of	Health Sciences
INWARD NO	3)0
DATE: 15/	116

To,

The Registrar

MGMIHS

Subject: Changes to be made in Microbiology Syllabus,

Respected Sir,

There are some changes which need to be made on page No 32 of Microbiology syllabus sent to you.

The following things need to be changed:

Sr No:	Existing	Changes
1.	A candidate has to obtain minimum of 47 marks out of 95 in theory, 13 marks out of 25 in practical, 11marks out of 30 in internal assessment and 75 marks out of 150 total to be declared as passed	A candidate has to obtain minimum of 47.5 marks out of 95 in theory, 12.5 marks out of 25 in practical, 10.5 marks out of 30 in internal assessment and 75 marks out of 150 total to be declared as passed

The hard copy of the page has been attached herewith for your reference.

BOS Dr A.D. Urhekar

Dr Deshmukh Dr M. Bhalchandra

Shall

HOD Microbiology

HOD Microbiology MGMMC, Aurangabad Chairperson BOS, Paraclinical Dr A.Shroff Chairperson Faculty of Medicine



# 111 10000 101 BOM 45/2016, Dated 28/04/2016 Resolution No. 3.2(b)

Resolution No. 3.2(b): Resolved to accept revised method to calculate internal assessment marks for IInd MBBS Exam effective from batch entering into 2nd MBBS from August 2016 onwards.

For Theory: ſ

	Microbiology	Pharmacology	Pathology	FMT
Day to day assessment as per MCI norma	10	10	10	07
Total marks	05	05	05	03
or Practical:	15	15	15	10

Microbiology IIIrd, IVth, Sem. & Prelim Exam. Pharmacology Pathology FMT 10 Day to day assessment as per MCI norms 10 10 07 05 05 05 Total marks 03 15 15 15 10

Prinded On. 14.12.2016

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From:	anahita BHESANIA [anahitapb@hotmail.com]
Sent:	Wednesday, December 14, 2016 1:12 PM
^j To:	REGISTRAR
Cc:	mgmihsaurangabad@gmail.com; anahita BHESANIA
Subject:	Model question papers and Integrated teaching topics to b included in Microbiology syllabus
Attachments:	Model question paper 1 for Microbiology Syllabus.docx; Model Question Paper 2 for Microbiology Syllabus.docx; Integrated teaching topics.docx

As asked by Dr Goel, Herewith sending

- Model question papers 1 and 2 and
- Integrated teaching topics

to be included in Microbiology syllabus. Shall be sending the hard copies of the same tomorrow.

Regards Dr Anahita V Bhesania Hodiwala Professor, Department of Microbiology. MGMMC, NM

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MOM Institute Of Health	Sciences 79 4121C
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## MGM INSTITUTE OF HEALTH SCIENCES, NAVI MUMBAI

#### **MBBS** university examination

# Paper 1: General Microbiology, Systemic Bacteriology and Applied Microbiology

#### Subject: Microbiology

Duration : 2 hrs

#### Instructions:

- 1. Attempt all questions
- 2. Mark the most appropriate answer in Sec –A (MCQS) by shading the respective circle option.
- 3. Maximum marks are indicated in the right
- 4. Illustrate the answers with suitable diagrams wherever necessary.
- 5. Please surrender your SWICHED OFF cell phones at entry point into examination hall.
- 6. Mobile phones, pagers, blue tooth or any other such communication devices are not allowed in the examination premises and in adjacent area.

#### Multiple Choice Questions:

(Darken the correct choice answer on the response sheet)

- 1. Pure growth is a growth of a single organism belonging to the same
  - a. Order
  - b. Family
  - c. Species
  - d. Genus
- 2. Spores are visible in
  - a. Grams staining
  - b. Negative staining
  - c. Modified ZN staining
  - d. Alberts staining
- 3. All are capsulated organisms except
  - a. B.anthracis
  - b. Y.pestis
  - c. H.influenzae
  - d. P.aeruginosa
- 4. Percentage of agar-agar in Loeffler's serum slope is
  - a. 0.2%
  - b. 1%
  - c. 2%

# Marks: 16x0.5=8mks

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Marks:40

Month/Year: _____

d. Nil

- 5. Dark field microscopy is useful to identify
  - a. Mycoplsma
  - b. Rickettisia
  - c. Spirochaetes
  - d. Brucella
- 6. Who is called "Father of antiseptic surgery"
  - a. Robert Koch
  - b. Louis Pasteur
  - c. Joseph Lister
  - d. Antony Von Leuwenhok
- 7. Red fluorescence when exposed to UV light is characteristic of

Ý

- a. B.uniformis
- b. B.stercoris
- c. B.fragilis
- d. B.melaninogenicus
- 8. The single most frequent etiologic agent of ascending UTI is
  - a. K.pneumoniae
  - b. E.coli
  - c. E.cloacae
  - d. S.marcescens
- 9. Pathogenesis of which disease does not involve an exotoxin
  - a. Typhoid fever
  - b. Botulism
  - c. Scarlet fever
  - d. Toxic shock syndrome
- 10. Pseudomonas are classified on basis of
  - a. Phage typing
  - b. Pyocin typing
  - c. Serology
  - d. Neutralisation
- 11. Number of serotypes of H.influenzae are
  - a. 4
  - b. 6
  - c. 10
  - d. 13
- 12. Xenodiagnosis is used for
  - a. Chlamydia
  - b. Rickettsiae
  - c. Brucella
  - d. Yersinia
- 13. Chicken cholera is caused by

- a. Pasteurella
- b. Yersinia
- c. Francisella
- d. Vibrio
- 14. Tularaemia is also called as
  - a. Malta fever
  - b. Haemorrhagic fever
  - c. Rabbit fever
  - d. Rift valley fever
- 15. Food poisoning due to ice cream is most probably due to
  - a. S.aureus
  - b. S.typhimurium
  - c. C.botulinum
  - d. Cl.perfringens
- 16. Which of the below organism does not have vertical transmission?

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- a. Syphilis
- b. Tuberculosis
- c. Measles
- d. Toxoplasmosis
### **SECTION B**

### Brief answer questions: (Answer any 4)

Marks: 4x4=16mks

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- 1. Enumerate all anaerobic culture methods. Add a note on Mc Intosh Fildes jar.
- 2. Plasmids
- 3. Give the principle and uses of dark ground microscope
- 4. Vapour phase disinfectants
- 5. Newer techniques for diagnosis of pulmonary tuberculosis.

SECTION C

### Long answer questions: (Answer any 2) Marks: 2x8=16 mks

- 1. Define Hospital Acquired infections (HAI), Sources of HAI, Write a note on Infection control policy.
- 2. Classify Streptococci ,Discuss pathogenesis of Streptococcus pyogenes ,Discuss lab diagnosis in a case of sore throat .
- 3. Classify pathogenic Clostridia , Add a note on lab diagnosis of gas gangrene ,Discuss morphology and toxins produced by Cl. Welchii.

## MGM INSTITUTE OF HEALTH SCIENCES, NAVI MUMBAI

### **MBBS** university examination

## Paper 2: Immunology, Virology, Parasitology, mycology

Subject: Microbiology

Duration : 2 hrs

### Instructions:

- 1. Attempt all questions
- 2. Mark the most appropriate answer in Sec –A (MCQS) by shading the respective circle option.
- 3. Maximum marks are indicated in the right
- 4. Illustrate the answers with suitable diagrams wherever necessary.
- 5. Please surrender your SWICHED OFF cell phones at entry point into examination hall.
- 6. Mobile phones, pagers, blue tooth or any other such communication devices are not allowed in the examination premises and in adjacent area.

### Multiple Choice Questions:

Marks: 16x0.5=8mks

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(Darken the correct choice answer on the response sheet)

- 1. Fab fragment of an immunoglobulin is made of
  - a. H chains
  - b. L chains
  - c. K chains
  - d. H and L chains
- 2. The predominant class of immunoglobulin in the blood of a newborn is
  - a. IgA
  - b. IgG
  - c. IgM
  - d. IgD
- 3. HLA is usually detected on
  - a. Neutrophils
  - b. Monocytes
  - c. Lymphocytes
  - d. Macrophages
- 4. Antigen presenting cells (APC) in the body include
  - a. Macrophage
  - b. B cells
  - c. T cells

Marks:40

Month/Year:

- d. NK cells
- 5. Romana's sign is positive in

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- a. Filariasis
- b. Chaga's disease
- c. Malaria
- d. Dracunculosis
- 6. Trichuris trichura is also known as
  - a. Pinworm
  - b. Seatworm
  - c. Whipworm
  - d. Roundworm
- 7. Stool examination for ova is not diagnostic in
  - a. Strongyloides
  - b. Trichuris
  - c. Ascaris
  - d. Ancyclostoma
- 8. Pruritis ani in children is caused by
  - a. Necator americanus
  - b. Ascaris lumbricoides
  - c. Trichuris trichura
  - d. Enterobius vermicularis
- 9. How many types of herpes viruses have been recognised?
  - a. 3
  - b. 6
  - c. 12
  - d. 40
- 10. Virus having affinity to lymphoid tissue is
  - a. Pox virus
  - b. Herpes virus
  - c. Cytomegalo virus
  - d. Epstein barr virus
- 11. Hepatitis A is an enteric virus of serotype
  - a. 68
  - b. 69
  - c. 70
  - d. 72
- 12. Number of segments in ssRNA genome in Orthomyxovirus is
  - a. 2
  - b. 8
  - c. 11

- d. 20
- 13. Candida infection is more commonly associated with

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- a. Diabetes
- b. Immunosuppression
- c. Both the above
- d. None of the above

## 14. Example of zoophilic dermatophytes include

- a. T.rubrum
- b. T.violecium
- c. M.audonii
- d. M.canis
- 15. The largest worm is
  - a. Ascaris
  - b. Hymenolepsis
  - c. Echinococcus
  - d. Taenia
- 16. Pathogenic free living amoebae include
  - a. Nagleria
  - b. B.coli
  - c. H.nana
  - d. Giardia

### SECTION B

### Short answer questions: (Attempt any 4)

### 4x4marks=16marks

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- 1. Differences between bacillary and amoebic dysentry
- 2. Lab diagnosis of Candida infection
- 3. Lab diagnosis of Human immunodeficiency virus
- 4. Innate immunity
- 5. Rabies vaccine

### Long answer questions: (Attempt any 2)

### 2x8marks= 16marks

- 1. Discuss pathogenicity, complications and lab diagnosis of E.histolytica in detail.
- 2. Define Agglutination. Discuss different types of agglutination reactions with their applications.
- 3. Name the viruses causing respiratory tract infections. Discuss Pathogenesis and Lab diagnosis of Influenza virus in detail.

**Resolution No. 1.3.7.1 of BOM-51/2017:** Resolved to continue the current Internal Assessment pattern for MBBS (i.e. 5 marks for Day-to-day assessment) for Pre and Para Clinical subjects (Anatomy, Physiology, Biochemistry, Microbiology, Pharmacology, Pathology and FMT). For rest of the subjects, Internal Assessment is to be calculated from terminal/Post end exam marks and Prelims examination, with immediate effect.

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**Resolution No. 1.3.8.13 of BOM-51/2017:** Resolved to approve the topics for vertical and horizontal integrated teaching in IInd MBBS Curriculum from batch entering in IInd MBBS in 2017-18 onwards.

		2. Microbiology	
Horizont	al Teaching:		
Sr no.	Topic	-Hrs	Departments
	Malaria	2hrs	Microbiology, Pathology,
2,	Tuberculosis	2 Hrs	Pharmacology Microbiology, Pathology
1			Pharmacology

## Vertical Teaching:

Sr no.	Topic	Hrs	Departments
1.	Typhoid and typhoid Ulcers	2 Hrs	Microbiology, Pathology,
2.	Meningitis	2 Hrs	Medicine Anatomy, Microbiology, Medicine
3.	Dermatophytes	2 Hrs	Microbiology and Dermatology

# **Syllabus of MBBS in Microbiology**

# **Topics for Integrated Teaching.**

# Horizontal Teaching:

Sr no.	Торіс	Hrs
1.	Malaria	2hrs
2.	Autoimmune Diseases	2 hrs
3.	Tuberculosis	2 Hrs

# Vertical Teaching:

Sr no.	Торіс	Hrs
1.	Typhoid and typhoid Ulcers	2 Hrs
2.	HIV	2 Hrs
3.	Meningitis	2hrs
4.	Fungal Infection	2 Hrs
5.	Rheumatic heart disease	2hrs

**Resolution No. 1.3.8.11 of BOM-51/2017:** Resolved to approve the topics to be included under Bioethics in UG. [Annexure-IX]

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**Bioethics** Topics for UG/PG

# Microbiology

# For Under-graduates (MBBS):

- 1. Universal principles
- 2. Sterilization techniques
- 3. Drug resistance minimization

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### Resolution No. 1.3.8.8 of BOM-51/2017: Resolved to:

(i) Introduce problem case discussion (problem based learning) in all paraclinical subjects on topics identified from batch entering in IInd MBBS in 2017-18 onwards. Annexure-VI

Problem based learning inputs for undergraduates (MINES)

### 1 Migrabiology

Propose a caredular a name devotiant a factore trablem based tearing in the factor. I discussion on ease biology or on their contraction in printical distribution of the completion of the particular topic in theory.

A minimum of 2 Problem based learning classes shall be scheduled in 4thSem and 5thSem MBBS each Covering the Following topics :

1) Discussion on clinical case history of enteric fever, Gonorrhoea&Leptospira.

- 2) Discussion on clinical history along with slide presentation :-
  - Malaria- Peripheral smear
  - Rhinosporidiasis
  - Molluscumcontagiosum / Negri bodies

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**Resolution No. 1.3.8.1 of BOM-51/2017:** Resolved that in absence of positive findings in stool mounts, students may be asked to draw diagrams/identify the possible findings of Ova / Cyst / Trophozoites Microbiology practical examination to be effective immediately.

### Resolution No. 3.5.9 of BOM-52/2018:

a) BOM reiterated the earlier BOM resolution as mentioned below:

Resolution No. 1.3.7.5 of BOM-51/2017: It was resolved that

- i) In all the subjects of all courses, MCQ weightage (Section A) shall be a maximum of 20% of the total marks in each paper.
- ii) BOS will have to accordingly workout the changes in Section B & C weightage and put up in forthcoming BOS meeting.
- iii) Further University Examination section must validate the MCQ Question Bank by Faculties before giving it to question paper-setter.

### b) To be effective from:

(i) Ist MBBS - Batch appearing in University August/September 2018 examination onwards.

(ii) <u>IInd MBBS</u> - Batch appearing in University January 2019 examination onwards.
 (iii) <u>IIIrd MBBS (Part I)</u> and IIIrd MBBS (Part II) - Batch appearing in University.

<u>IIIrd MBBS (Part I)</u> and IIIrd MBBS (Part II) - Batch appearing in University January 2019 examination onwards.

**Resolution No. 4.2.1 of BOM-53/2018:** Resolved that the printed format of the Medico-legal examination proforma (sexual violence) may be provided to 2nd MBBS students during practical's in formative and summative assessments **[Annexure-X]**, to be applicable from batch entering into 2nd MBBS 2017-18 onwards.

# Anneslure 30 For item NO. 4

CONFIDENTIAL

Annexure - X

31

### Medico-legal Examination Report of Sexual Violence

	Name of the Hospital OPD No Inpatient No.	<b>) .</b>	,		
2.	Name	••••		• • • • • • •	
З.	Address				
4.	Age (as reported) Date of Birth (if known)				
5.	Sex (M/F/Others)				
6.	Date and Time of arrival in the hospilal				
7.	Date and Time of commencement of examination				
8.	Brought by (Name & signature	s)			
9.	MLC NoPolice Station.				
10.	Whether conscious, priented in time and place and person Manual			• • • • • • • • •	
	Any physical/intellectual/psychosocial disability				
(Int	erpreters or special educators will be needed where the survivor ha	s spec	iatre	eds	such
12. I	nearing/speech disability, language barriers, intellectual or psychos Informed Consent/refusal D/o or S/oD/o elby give my consent for: medical examination for treatment this medico legal examination	ociald	lisəbi	lity.)	 E i
12. I her a)	nearing/speech disability, language barriers, intellectual or psychos informed Consent/refusal D/o or S/o eby give my consent for: medical examination for treatment	ocial d	lisabi	lity.) No	
12. her a) b) c) l sl: exp	nearing/speech disability, language barriers, intellectual or psychos Informed Consent/refusal 	ves Yes Yes Yes Yes		lity.) No No No	
12. her a) b) c) l sl: exp	nearing/speech disability, language barriers, intellectual or psychos Informed Consent/refusal 	ves Yes Yes Yes Yes	this	lity.) No No No	

If special educator/Interpreter/support person has helped, then his/her name and signature.....

Name & signature of survivor or parent/Guardian/person in whom the child reposes trust in case of child (<12 yrs) With date, time & place Name & signature/thumb impression of Witness .......... With Date, time and place 13. Marks of identification (Any scar/mole) (1)..... (2)..... Left Thumb impression 14. Relevant Medical/Surgical history : Onset of menarche (in case of girls) Yes No Age of onset..... Menstrual history - Cycle length and duration ...... Last menstrual period..... Menstruation at the time of incident - Yes/ No, Menstruation at the time of examination - Yes/ No Was the survivor pregnant at time of incident - Yes/No. If yes duration of pregnancy ....... weeks Contraception use: Yes/No...... If yes - method used: ..... Vaccination status - Tetanus (vaccinated/not vaccinated). Hepatitis B (vaccinated/not) (vaccinated) 63

3

(I) Date of incident/s being reported (ii) Time	e of incident/s	(iii) Location/s
(iv)Estimated duration : 1-7 days 1 week to 2-6 months	2 months	• • • • • • • • • • • • • • • • • • •
(v) Number of Assailant(s) and	·····	
name/s (vi) Sex of assallant(s) (s)	Approx or – relationship w	. Age of assallant ith the
(vir) Description of Micident in the words of the national Narrator of the incident: survivor/informant (speci	mator: fy name and relation	on to survivor)
If this space is insufficient use extra page		
5 B. Type of physical violence used if any (De	scribe);	
5 B. Type of physical violence used if any (De	scribe): Burned with	۱ 
5 B. Type of physical violence used if any (De Hit with (Hand, fist, blunt object, sharp object)	••• •••••••	
5 B. Type of physical violence used if any (De Hit with (Hand, fist, blunt object, sharp object) Biting	Burned with	
5 B. Type of physical violence used if any (De Hit with (Hand, fist, blunt object, sharp object) Biting Pinching	Burned with Kicking	۰
^	Burned with Kicking Pulling Hair	
5 B. Type of physical violence used if any (De Hit with (Hand, fist, blunt object, sharp object) Biting Pinching	Burned with Kicking Pulling Hair Banging head	
5 B. Type of physical violence used if any (De Hit with (Hand, fist, blunt object, sharp object) Biting Pinching Violent shaking	Burned with Kicking Pulling Hair Banging head	

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15 C.

١.	Emotional abuse or violence if any (insulting, cursing, belittling, terrorizing)
ü.	Use of restraints if any
₩i.	Used or threatened the use of weapon(s) or objects if any
ív.	Verbal threats (for example, threats of killing or hurting survivor or any other person in whom the survivor is interested; use of photographs for blackmalling, etc.) If any:
v. vi.	Luring (sweets, chocolates, money, job) if any: Any other:

### 15 D.

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i. Any H/O drug/alcohol intoxication:

ii. Whether sleeping or unconscious at the time of the Incident: .....

15 E. If survivor has left any marks of injury on assailant/s, enter details:

15 F. Details regarding sexual violence:

Was penetration by penis, fingers or object or other body parts (Write Y=Yes, N=No, DNK=Don't know) Mention and describe body part/s and/or object/s used for penetration.

	Penetralion			Т Ел	Emission of Semen		
Orifice of Victim	By Penis	By body part of self or assailant or third party (finger, tongue or any other)	By Object	Yes	NO	Don't know	
Genitalia (Vagina and/or urethra)							
Anus	······································						
Mouth							
Oral sex pe	erformed by a	assallant on survivor				·····	
			Y		N	DNK	
Forced Masturbation of self by survivor		Y		N	DNK		

Maaturbatur	· · · · · · · · · · · · · · · · · · ·	) N	] DNK
Masturbation of Assailant by Survivor, Forced Manipulation of genitals of assailant by survivor	Υ Υ	N	DNK
Exhibitionism (perpetrator displaying genitals)	Y	N	DNK
Did ejaculation occur outside body orifice (vagina/anus/mouth/urethra)?	Y	N	DNK -

Y	N	If Yes, describe
Ŷ	N	lf Yeş, describe
Y	N	DNK
Y	N	DNK
Y	N	DNK
and counts and form of the Bournes's services	·····	
		1999 - Angel State and State
	Y Y Y Y Y	Y N Y N Y N

* Explain what condom and lubricant is to the survivor

and a construction of the second s

Post Incident has the survivor	Yes/No/Do Not know	Remarks
Changed clothes		
Changed undergarments		-
Cleaned/washed clothes		
Cleaned/washed undergarments		
Bathed		:
Douched		
Passed Urine		· ·
Passed stoots	1	
Rinsing of mouth/Brushing/ Vomiting (Circle any or all as appropriate)		

H/o vaginal/anal/oral bleeding/discharge since the incident of sexual violence.....

H/o painful urination/ painful defecation/ fissures/ abdominal pain/pain in genitals or any other part since the incident of sexual violence

16.	General Physical Examination-
i,	Is this the first examination.
il.	Pulse
iii.	TempResp. Rate
Ī٧.	Pupils
V,	Any observation in terms of general physical wellbeing of the survivor

# 17. Examination for injuries on the body if any

(

The pattern of injuries sustained during an incident of sexual violence may show considerable variation. This may range from complete absence of injuries (more frequently) to grievous injuries (very rare).

(Look for bruises, physical torture injuries, nail abrasions, teeth bite marks, cuts, lacerations, fracture, tenderness, any other injury, boils, lesions, discharge specially on the scalp, face, neck, shoulders, breast, wrists, forearms, medial aspect of upper arms, thighs and buttocks) Note the Injury type, site, size, shape, colour, swelling signs of healing simple/grievous, dimensions.)

Scalp examination for areas of tenderness (if hair pulled out/ dragged by hair)	
Facial bone injury: orbital blackening, tenderness	
Petechial haemorrage in eyes and other places	
Lips and Buccal Mucosa / Gums	•
Behind the ears	
Ear drum	
Neck, Shoulders and Breast	
Upper limb	
Inner aspect of upper arms	·
Inner aspect of thighs	· · · · · · · · · · · · · · · · · · ·
Lower limbButtocks	
Other, please specify	
m Nath Press of spectra (), & Main of a long of the fact of the spectra state of the spectra	1 Į





18. Local examination of genital parts/other orifices':

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THE REAL PROPERTY.

A. External Genitalia: Record findings and state NA where not applicable.

Body parts to be examined	Findings	
Urethral meatus & vestibute		
Labia majora		
Labia minora	***************************************	
Fourchette & Introitus		
Hymen Perineum		
External Urethral Meatus	**************************************	
Penis	······································	······································
Scrolum	Malannagerging and a second and an an an and a second and a	
Tesles	-	
Clitoropenis		
Labioscrotum		
Any Other		
P/S findings if performed P/V findings if performed Record reasons if P/V of P/S ex C. Anus and Rectum (encircle Bleeding/tear/discharge/oe	amination performed	
D. Oral Cavity - (encircle the re Bleeding/ discharge/ tear/oe	blevant) dema/tenderness	
9. Systemic examination:		
Central Nervous System: Cardio Vascular System:		



- 20. Sample collection/investigations for hospital laboratory/ Clinical laboratory
- 1) Blood for HIV, VDRL, HbsAg
- 2) Urine test for Pregnancy/
- 3) Ultrasound for pregnancy/internal Injury
- 4) X-ray for Injury
- 21. Samples Collection for Central/State Forensic Science Laboratory
- 1) Debris collection paper
- Clothing evidence where available (to be packed in separate paper bags after air drying)

List and Details of clothing worn by the survivor at time of incident of sexual violence

## 3) Body evidence samples as appropriate (duly labeled and packed separately)

Collected/Not Collected	Reason for not collecting
	· · · · · · · · · · · · · · · · · · ·
······································	
	1
*** ······	
	Collected/Not Collected

4) Genital and Anal evidence (Each sample to be packed, sealed, and labeled separately-to be placed in a bag)

* Swab sticks for collecting samples should be moistened with distilled water provided.

	Collected/Not Collected	Reason for not collecting
Matted pubic hair		
Pubic hair combing (mention if shaved)		
Cutting of pubic hair (mention if shaved)		at <u>en non en en</u>
Two Vulval swabs (for semen examination and DNA testing)		
Two Vaginal swabs (for semen examination and DNA testing)		
Two Anal swabs (for semen examination and DNA testing)		
Vaginal smear (alr-dried) for semen examination		······································
Vaginal washing	······································	
Urethral swab		
Swab from glans of penis/clitoropenis		

*Samples to be preserved as directed till handed over to police along with duly attested sample seal.

22. Provisional medical opinion

Samples collected (for FSL), awaiting reports

Samples collected (for hospital laboratory)

Clinical lindings

Additional observations (if any)

23. Treatment prescribed:

Treatment	Yes	NO	Type and comments
ST) prevention treatment			
Emergency contraception			
Wound treatment			
Tetanus prophylaxis			
Hepatitis B vaccination		·····	
Post exposure prophylaxis for HIV	-l		
Counselling		! !	
Olher			

24. Date and time of completion of examination ...... This report contains ....... number of sheets and ...... number of envelopes.

> Signature of Examining Doctor Name of Examining Doctor Seal

Place:

## 25. Final Opinion (After receiving Lab reports)

Place:

Signature of Examining Doctor

1'

Name of Examining Doctor

Seal

### COPY OF THE ENTIRE MEDICAL REPORT MUST BE GIVEN TO THE SURVIVOR/ VICTIM FREE OF COST IMMEDIATELY

**Resolution No. 4.3.5 of BOM-53/2018:** Resolved to add reference book entitled "ESSENTIAL IN RESPIRATORY MEDICINE" by Dr. S.H. Talib in the UG/PG curriculum in medicine and allied subjects

**Resolution No. 4.5.2.1 of BOM-55/2018:** Resolved to introduce training in 'Research Methodology' for 3rd Semester MBBS students entering in 3rd Semester from September 2018 onwards. It was further resolved that responsibility of this training will be with Pharmacology department.

**Resolution No. 4.5.2.2 of BOM-55/2018:** Resolved to include the topic on 'Emerging and Reemerging infections' in MBBS Microbiology syllabus with immediate effect. **Resolution No. 4.5.2.3 of BOM-55/2018:** Resolved to provide the printed standard format of the Medico-legal examination (Age,Alcoholic,Weapon,Injury,Death,Potency,Sickness,Fitness) to 2nd MBBS students during practical examination in formative and summative assessments. **[Annexure-34-A,B,C,D,E,F,G,H]** 

Recd. on 18/11/2018 **Examination for Determination/Estimation of Age** Annexure - 34-A To. The Reference : Your Letter No. _____ Dated_____ Name : Age stated : ; Sex : ; Occupation : _____ Marital status : Address : Brought by Police Constable : ______ No. : _____; P.S. _____ Identified by : . Date and Time of Examination : Place of Examination : Consent : Signature of Examinee (If minor below 12 yrs. consent of Parents/Guardian) Examined in presence of :_____ (If female) (Signature of female attendant) Identification marks : 1._____ 2._____ Education : Birth Date : **Physical Examination :** 2. Weight : _____kg 1. Height : ____cm 3. Chest girth at the level of nipple : cm 4. Abdominal girth at the level of navel : cm 5. General build and appearance : 6. Hairs : Pubic : _____, Axillary : _____, Facial : _____, Scalp : _____

7. Development of breasts :	
8. Development of genitals :	
9. Onset of Puberty :	
Voice :	Adam's apple :
Date of menarche : 10. Dental Status :	Adam's apple : Regularity of menses :
10. Dental Status :	
	Upper Jaw (Maxillary Teeth)
	Lower Jaw (Mandibular Teeth)
	isower saw (manufoular Teeth)
11. Advised X-ray :	
a.	
Ь.	
С.	
X-ray' plate No.: a	bc.
Dated :	bc

# Provisional Age Certificate

On clinical examination of the individual, age is about ______ years. However, the final opinion regarding the age should be collected from this office after submission of the Radiological report and the birth certificate.

Signature

(Dr.

)

Designation & Seal

Place : _____
Date : _____

## Age Certificate

То			
The			
Reference : Age estimation of		, Dated _	
Sir,			
I, Dr		, after going thro	ough the findings
of			· · · · · · · · · · · · · · · · · · ·
Physical examination report No.			
X' ray plate No.		, Dated	
Radiological Examination report No		, Dated	Į
and the Date of Birth Certificate No		, Datec	£
produced before me,			
I am of the opinion that the indivi	dual's age is a	bout	years
		Signature	
	(Dr.	-	)
		Designation & Sea	1

Place : _____
Date : _____

· .

### Examination / Certification of Alcoholic

	A Mod	el Scheme of Examination	Annequire -34-B
То,			(Anneaure Sta
The Investigating Officer I	P.S.		
Reference : Your letter No.		Dated :	
I am forwarding herewith	the result of	my examination of	
Name :		Son / daughter / wife / widow	of
Age :	Sex : M/F	Weight :	
Address :			
Consent for examination :			
n a		Signature / Thumb impression	of Examinee
Identification Marks : 1. 2.			
Brought by P.C. Name :		No. P.S.	
Date and time of examinat	ion :		
Place of examination :			
History :			
a. Alleged case -			
<ul> <li>b. Related to alcohol -</li> <li>c. Illness -</li> </ul>			
General behaviour :			
Clothing :			
Attitude :			
Memory :		Mental alertness :	
Pulse :		Respiration :	
Temperature :		Blood pressure :	
Skin :			
Smell of alcohol, if any :			

Lips :	Tongue :
Eye :	Pupils :
Conjunctiva :	
Muscle co-ordination :	
Gait :	Speech :
Handwriting	
Reflexes :	
Systemic examination :	
Respiratory System :	
Cardio-vascular System :	
Gastro-intestinal Tract :	
Laboratory investigations : <b>a. Blood</b> (5 to 10 ml venous blood) <b>Preservat</b> <b>b. Urine</b> (10 to 20 ml - 2 samples) <b>Preservati</b> <b>c. Expired air :</b>	
Diagnosis :	
Opinion : I am of the opinion that -	

- The above person has consumed alcohol and is under its influence. 1.
- 2. The above person has consumed alcohol and is not under its influence.
- 3.

Place :

Date :

Time :

The above person has not consumed alcohol.

Signature

(Dr. Designation & Seal

)

¢

### Form 'A'

ź	See	$\mathbf{p}_{\mathbf{r}}$	do.	No	3)
ŧ	ove	IX U	nc.	1101	31

(Certificat or has not	te by Registered Medical Practitior consumed an intoxicant)	her showing whether a person examined by him has		
Serial No	•	Name & location of the		
		Dispensary or Hospital		
Certified t	that Shri / Smt / Kum	Resident of		
was broug	ght to this Hospital / Dispensary by	· ·		
		(Here state the Name & Designation of the Officer)		
on	at	A.M. / P.M. & was examined by me		
	at			
A clinical	examination of the above person c	lisclosed the following :		
Age:	Years, Weight :	kg,Height :cm		
Breath :	Smelling / Not smelling of Alcol	nol / Ganja / Bhang.		
Speech :	Incoherent / Normal			
Gait :	Unsteady / steady			
Pupils	Dilated / Normal			
Additiona	al remarks, if any :			
I find that	t the above named person			
	HAS CONSUMED	Alcohol / Ganja / Bhang		

### HAS NOT CONSUMED ANY INTOXICANT

### I also find that he / she is not under the influence of alcohol.

(N.B. : Blood from the body of the above named was / was not collected by me for chemical examination)

"Certified that the procedure laid down under the rule (4) of Bombay Prohibition Medical Examination and Blood Test Rule 1959 has been followed."

Date :	
Time :	A.M. / P.M.

Signature Designation

Signature / Thumb impression of the Person examined.

Marks of identification of the person examined in case he refuses to give his signature or thumb impression
		Form "B"
		No
From,		
The Casualty N	Medical Officer, / Assis	tant Professor in Forensic Medicine
MGM Medica	l College and Hospital,	
Aurangabad	•	
To,		
The Director		
	ce Laboratory & Chem	ingl Another
	rashtra, Mumbai	
Govi, or manar	ashua, wumbai	Date :
Sir / Madam,		
I am forwardin	g herewith a parcel by	post / with Shri
01	containing	ml. of Blood and / or Urine sample collected by
	ai	A.M. / P.M. from the body of Shri / Shrimati / Kumari
:		of who
was produced b	efore me for medical e	xamination and/or collection of Blood and / or Urine from
ins / net bouy	бу	and parameter to that the
Blood and / or l	Urine and issue a certif	icate (in duplicate) regarding the result of the tests.

"Certified that the procedure laid down under the rule (4) of Bombay Prohibition Medical Examination Blood Test Rule 1959 has been followed".

Yours faithfully,

( Dr.

)

Casualty Medical Officer Assistant Professor in Forensic Medicine MGM Medical College and Hospital, Aurangabad

Facsimile of the Seal or Monogram used for Sealing the Phial containing Blood and/or Urine

# Examination of the Weapon

То				Annexure-34-c
The Investigating O	fficer,			
Police Station		11.11.11.11.11.11.11.11.11.11.11.11.11.		
Reference : Your let	ter No.		Dated	
Sir,				,
With reference to the with the injuries of	e above letter, I am se	ending the report about	weapon sen	t sealed in connection
		Kind of weap		
Description of the w				
Blade : Is of		, Texture :		
		, Thick		
		, Point :		
		, Hilt : Size :		
		, Texture :		
		readth / Circumference		
		nd it to C.A. for furthe		
Injuries possible :				, ,
Injuries impossible :				
Identification marks	if any on the weapon			
(Put the signature on	the weapon)			
The weapon packed,	sealed and handed or	ver to P.CN	lo	P.S
Place :				
Date & Time :				
Receipt of weapon &	report		Signature	
		(Dr.		)
		Dec	ionation & S	aal

Designation & Seal

# **Examination / Certification of the Injured (Injury Report/Certificate)** То Annexure-34-D The Investigating Officer. Police Station Reference : Your Letter No. _____ Dated _____ Sir. I am forwarding herewith the report of examination of : Name of Injured : ______ Son/Wife/Daughter/Widow of _____ Surname ______ resident of ______ Age: ______ Sex _____Occupation_____ Brought by PC ______ No. _____ P.S. _____ Consent for examination : Signature of Witness Signature of Examinee Identification marks: 1. 2.

#### **History** :

Sr. No.	Type of injury	Size of injury	Situation over the body	Nature of injury	Probable weapon	Age of injury	Advice

Remark

Place :

Date :

Signature

)

**Designation & Seal** 

(Dr

Receipt

#### Form No. 4

(For hospital in patient death, not to be used for still birth)

## MEDICAL CERTIFICATE OF CAUSE OF DEATH

(To be sent to Registrar of Births and Deaths along with Death Report form no. 2)

Name of Hospital :

I do hereby certify that the person whose particulars are given below died in Hospital in Ward No.

on ______ at _____ A.M. / P.M.

Name of the deceased :

Addres	s of norm	al Residenc	e:				sta	ntistical	office	
				Occupation	Religion		Age at Death			Detailed list
	yrs	Bírth	status S, M, W or D			If und yea Months			ler 24 urs Min.	code
				4	Cause of D	eath		terval be iset and		vorox
1. Imm	ediate C	ause :		a)					uono, up	.p.o.d
State th	e disease	, injury or c	omplication		Due to :					
			ode of dying	or as a c	onsequence	e of				
		lure, asthen	ia, etc.							
	dent cau			b)						
			ving rise to the		Due to :					
above c	ause, sta	ing underly	ring condition la							
2 Othe	r sionific	ant conditio	ne	C)	F					
			of related to the							
		ion causing								
		Natural /	Accident / Suic	ide / Homici	de (specify	) : How di	d the ini	ury occu	ur?	
IF DFC	TEASED	WAS A FF				· · · · · · · · · · · · · · · · · · ·				
			with pregnancy	· ?		Yes/N	0			
	ere a del		in pregnancy	•	Yes/No					
		**								
Name oi	rubber-st	amp of instit	ution :	Serial Nu	Serial Number of institution				Date of report	
Date an	d Time :					Signat	ture and	address	of	
				(	Dr.				)	
			· · · · · · · · · · · · · · · · · · ·			Desig	nation &	Seal		
			be detached a			lative of th	ie deceas			*****
Certified that Shri / Smt/Kum							,	Reside	ent of	
			was admi							
Date Ti	me :						Signati	are		
				(	Dr.		~		)	
						Desig	nation &	Seal	,	

For use by

#### **EXAMINATION OFA CASE FOR DETERMINATION OF POTENCY**

	FM No/		/20	
	Date : /	/	/ 20	
To,		entrained.	Annexy	re-34-F
Defense a Very letter / order no	Datad		or CP and the Reservation and Country All Country and	
Reference : Your letter / order no.				
Name of the individual				
Age as stated:, Sex: Ma	rital status (If married, du	uratior	ı)	
Address :				
Occupation :				
Brouught by (Name, signature & designation	.)			
Date, place & time of examination :				
Light arrangement				
Consent :				
Q - Are you willing to be examined by me examination will include physical examination assessment. The examination by dept of U to evaluate your potency. You have right	nation, laboratory investi Jrology would also inclu	gation de adr	ns and psych ministration	ological of drugs

court of law.

Answer given - Yes / No

Name, signature of the person giving consent with Date -

Witness to the consent - Name, signature & Date -

Identification marks-

1.

2.

### History

1. Do you have erectile dysfunction ? - Yes / No

If yes

a. Since how long have you noticed the erectile dysfunction?

b. Did the problem being abruptly or insidiously?

- c. Do you have inability to achieve or maintain an erection or both ?
- d. Are you able to penetrate or not?
- e. Whether partial penetration or ejaculation before penetration?
- f. Do you ever get normal or near normal erection (During masturbation with other partner, early morning)
- 2. H/o any major illness HT / DM / TB / Vascular disease / Endocrinal diseases etc.
- 3. H/o STD -
- 4. H/o mental illness -
- 5. Any stress-
- 6. Family environment-
- 7. Any history of medication / for what ailment / duration of medication
- 8. H/o Drug abuse Nicotine / Ganja /Alcohol / other
- 9. H/o any head injury / spinal injury / any operation on genitals -
- 10. H/o aversion dislike / dejection / for any particular sex partner

### **Obsevations**

## **General examination**

Gen	eral built and appearance	· •					
Weig	ght : kg	Height :	cm				
Teet	h :	Total	1 No. :				
Seco	ondary sexual characters	:					
Bear	rd :		Moustache :				
Axil	lary hairs :		Pubic hairs :				
Brea	ast development / Gynae	comastia if any :					
Any	marks of injury / scar of	the body :					
Loca	al examination : (Along	with Urology department) d	one in ward no				
a.	Penis :						
	Circumcised / Non-Circumcised :						
	Stretched penile length -						
	Length when erect -						
	Circumference (flaccid & erect) :						
	Disease / deformity / in	ury (if any) :					
	Sensation over glans penis :						
	Foreskin (Retractable / Non-retractable) :						
	Dorsal penile pulsation :						
	Any Discharge :						
	Smegma :		1				
	Hygine :						

Scrotum : b. Pendulous or not : Developmental defects : Deformities : Cremasteric reflex : Testes : e. Whether present in scrotum or not : Size : Consistency : Prostate (Per rectal examination) : đ. Bulbocavernous reflex : e. Any evidence of S.T.D £ Effect of administration of ______in _____dose _____After _____minutes g. Result :

## SYSTEMIC EXAMINATION

- C.N.S. :
- R. S. :
- C. V. S. Pulse : BP:

Femoral artery :

Dorsalispedis artery :

• G.I.T. :

# Laboratory Investigations (If required)

- 1. CBC :
- 2. Hb:
- 3. BSL (Fasting & PP) :
- 4. Sr. FSH :
- 5. Sr. LH :
- 6. Sr. testosterone & Oestrogen :
- 7. Sr. prolactin :
- 8. VDRL :
- 9. USG/Colour doppler :
- 10. TFT (TSH, T3, T4):
- 11. LFT:
- 12. HbA1C:

**Opinion** :After detailed examination i.e. based on physical examination, psychiatric evaluation and examination by urologist, we are of the following opinion". There is nothing to suggest that the above examined person is incapable to perform sexual intercourse ". / The person is in capable of performing sexual intercourse due to.....

Place :

Date _____

Signature Name & Qualification : Designation Registration No. :

# MEDICAL SICKNESS / UNDER TREATMENT CERTIFICATE

Signature of the applicant(Ge	Annexyse-34-1
(Go	overnment servant / Private)
I Dr	after careful
	rtify that Mr. / Mrs./ Ms
	whose signature is given above was suffering
	and was under my treatment for the same as
	hat a period of absence from duty of
with effect from	n is absolutely necessary for restoration
of his / her health	source in the second seco
He / She was advised rest for a period of	days
Identification marks:	
1)	
2)	
Hospital No.	
Date:	Authorised Medical Attendant Seal & Reg. No.

# MEDICAL FITNESS CERTIFICATE

Signature of the applicant	
(Gove	rnment servant / Private)
1 Dr	after careful
personal examination of the case hereby certi	fy that Mr. / Mrs. / Ms
	whose signature is given above was suffering and was under my treatment for the same.
He / She recovered completely from the illness	and he/she is fit to resume his / her duty with effect
from	
Identification marks:	
1)	
2)	
1	
Hospital No.	
	Authorised Medical Attendant Seal & Reg. No.

.

X

# **Certificate of Physical Fitness**

This is to Certify that I have examined Shri / Sm	nt/Kum. Annexyre-34-H
	who signed below in my presence and who
is a candidate for employment for the post of	in
the department / office	at
I could not discover that he / she has any diseas	se (communicable or otherwise) constitutional
weakness or bodily infirmity, except	I do consider / do not consider
this is a disqualification for such an employment.	
He / she	age is according to his / her own
statement years and by appearance ab	out years.
Identification marks:	
1)	
2)	
Signature of the applicant :	
(Government se	rvant / Private)
Houmital No.	
Hospital No.	
Date: Aut	thorised Medical Attendant
Sea	l & Reg. No.

#### Resolution No. 4.13 of BOM-55/2018: Resolved as follows:-

- (i) Slow learners must be re-designated as potential learners.
- (ii) Students scoring less than 35% marks in a particular subjects/course in the 1st formative exam are to be listed as potential learners. These learners must be constantly encouraged to perform better with the help of various remedial measures.
- (iii) Students scoring more than 75% marks in a particular subjects/course in the 1st formative exam are to be listed as advanced learners. These learners must be constantly encouraged to participate in various scholarly activities.

#### Resolution No. 3.1.4.2 of BOM-57/2019:

- i. Resolved to include "Gender Sensitization" into UG (from new batch 2019-2020) and PG (from existing batches) curricula. [Annexure-21]
- **ii.** Resolved to align the module of "Gender Sensitization" with MCI CBME pattern for MBBS students.
- iii. Resolved that Dr. Swati Shiradkar, Prof., Dept. of OBGY., MGM Medical College, Aurangabad will coordinate this activity at both campuses.

# Annexure - 21

Gender sensitization for UG (2nd, 3rd, 8th semesters) and PG (3 hours)

## **INCLUSION OF "GENDER SENSATIZATION" IN CURRICULUM**

### Introduction :

The health care provider should have a healthy gender attitude, so that discrimination, stigmatization, bias while providing health care will be avoided. The health care provider should also be aware of certain medico legal issues related with sex & gender.

Society particularly youth & adolescents need medically accurate, culturally & agewise appropriate knowledge about sex, gender & sexuality. So we can train the trainers for the same. It is need of the hour to prevent sexual harassment & abuse .

To fulfill these objectives, some suggestions are there for approval of BOS.

# <u>Outline</u>

1)For undergraduates :- Three sessions of two hours each, one in  $2^{nd}$  term, one in  $3^{rd}$  term & one in  $8^{th}$  term.

2)For Faculties and postgraduates :- One session of two hrs .

3)For those want to be trainers or interested for their ownself, value added course, which is optional about sex, gender, sexuality & related issues.

## **Responsibility**

ICC of MGM, MCHA , with necessary support from IQAC & respective departments.

### **Details of undergraduate sessions**

### 1)First session in 2nd term

Aim – To make Students aware about the concept of sexuality & gender.

To check accuracy of knowledge they have,

To make them comfortable with their own gender identify & related issues.

To make them aware about ICC & it is functioning.

**Mode** – Brain storming , Interactive power point presentation experience sharing.

**Duration** – Around two hours

**Evaluation** – Feedback from participants.

# 2)Second session in 3rd / 4th term

**Aim** – To ensure healthy gender attitude in these students as now they start interacting with patients.

To ensure that the maintain dignity privacy while interacting with patients and relatives, particularly gender related.

To make them aware about importance of confidentiality related with gender issues.

--2--

To encourage them to note gender related issues affecting health care & seek solutions.

Mode – focused group discussions on case studies, Role plays & discussion.

--3--

Duration – Around two hours.

Evaluation – Feedback from participants.

Third session in 8th term.

**Aim** – To understand effect of gender attitudes on health care in various subjects.

To develop healthy gender attitude while dealing with these issues.

**Mode** – Suggested PBL by departments individually. (In collaboration with ICC till faculty sensitization is complete)

**Evaluation** – Feedback

****

--4--

# FOR POSTGRADUATES

Session of 2-3 hrs preferably in induction program.

- **Aim** To introduce medically accurate concept of gender, sex, gender role & sex role.
- To ensure healthy gender attitude at workplace.

To understand gender associated concepts on health related issues & avoid such bias wile providing health care.

To make them aware about ICC & it's functioning.

Mode – Interactive PPT

Role plays & discussion

Duration – 2 to 3 hrs

**Evaluation** – Feedback.

## --5--

## FOR FACULTIES

Session of 2 hours may be during combined activities.

**Aim** – To ensure clarity of concept abut gender & sex.

To discuss effect of these concept on health related issues.

To identify such gender & sex related issues in indivual subject specialties.

To discuss methodology like PBL for under graduate students when whey are in  $7^{\text{th}}-8^{\text{th}}$  semester.

Mode – Role play

Focused group discussion

**Case studies** 

**Evaluation** – Feed back.

*****



# MGM INSTITUTE OF HEALTH SCIENCES

(Deemed to be University u/s 3 of UGC Act, 1956) Grade 'A' Accredited by NAAC Sector-01, Kamothe, Navi Mumbai - 410209 Tel 022-27432471, 022-27432994, Fax 022-27431094 E-mail- registrar@mgmuhs.com Website : www.mgmuhs.com

