



# PRAVARA INSTITUTE OF MEDICAL SCIENCES

## (DEEMED TO BE UNIVERSITY)

Loni, Tal. Rahata, Dist. Ahmednagar 413736  
NAAC Re-accredited with 'A' Grade

## SYLLABUS

### UG Programme- Pediatrics

(Competency Based Undergraduate Curriculum will be implemented from August 2019, i.e. MBBS batch admitted for first year in 2019)

### Course Content

(Based on Medical Council of India,  
Competency based Undergraduate curriculum for the Indian Medical Graduate, 2018. Vol. 2/ 3; page nos. 150-201)

1. Total Teaching hours: 105 hours (Lectures +Tutorials);  
15 hours (Self-directed learning);  
174hoursClinicalposting
2. A. Lectures (hours): 40 (20 hours each in III MBBS Part I & Part II)  
B. Self-directed learning (hours): 15 (5 hours in III MBBS Part I & 10 hours in III MBBS Part II)  
C. Clinical Postings (hours): 174 (2weeks/4weeks/4weeks)  
D. Small group teachings/tutorials/  
Integrated teaching/Practicals(hours): 65 hours (30 hours in III MBBS Part I and 35 hours in III MBBS Part II)
  - 8 symposia will be conducted from theory topics in
  - 15 hours of Self- directed Learning (3 in III MBBS (Part I) and 5inIIIMBBS(Part II))
  - Two(02)Full day workshops
  - IMNCI
  - NRP
  - Module 4.7 AETCOM Module will be covered in III MBBS (Part II) (05 hours)

**Tutorials/ Small Group Discussions III (PartI) MBBS (30 hours)**

| S. No | Topic   | Hours | Lectures (Competency No.)                                  | SLO  | Horizontal Integration |
|-------|---|-------|--|--|------------------------|
| 1     | Normal Growth and Development                     | 01    | Developmental milestones (PE 1.5, 1.6)                     | <ol style="list-style-type: none"> <li>1. Definition of Development</li> <li>2. Principals of development</li> <li>3. Factors affecting Development</li> <li>4. Domains of Development</li> <li>5. Milestones in various domains</li> <li>6. Developmental assessment</li> </ol>                       | Psychiatry             |
| 2     | Common problems related to growth                 | 02    | Failure to thrive (PE 2.1, 2.4)                            | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>1. Clinical Features</li> <li>2. Evaluation of a child with Failure to thrive</li> <li>3. Management</li> </ol>   |                        |
|       |   |       | Short stature (PE 2.6)                                     | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Clinical Features</li> <li>4. Evaluation of a child with Short stature</li> <li>5. Management</li> </ol>   |                        |
| 3.    | Care of the Normal Newborn, and High-risk Newborn | 02    | Care of normal newborn (PE 20.1, 20.2, 20.6,)              | <ol style="list-style-type: none"> <li>1. Define the common neonatal nomenclatures including the classification</li> <li>2. Describe the characteristics of a Normal Term Neonate and High-Risk Neonates.</li> <li>3. Explain the care of a normal neonate</li> </ol>                                  | Obs & Gynae            |
|       |   |       | Temperature regulation and Neonatal hypothermia (PE 20.12) | <ol style="list-style-type: none"> <li>1. Temperature regulation in neonates</li> <li>2. Disorders of temperature regulation</li> <li>3. Definition of hypothermia</li> <li>4. Prevention of hypothermia</li> <li>5. Clinical features of hypothermia</li> <li>6. Management of hypothermia</li> </ol> |                        |

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| 4. | To promote and support optimal Breast feeding for infants                              | 01 | Breast Feeding (PE 7.1, 7.2, 7.3, 7.4, 7.6)             | <ol style="list-style-type: none"> <li>1. Awareness on the cultural beliefs and practices of breast feeding.</li> <li>2. Enumerate advantages of breast feeding</li> <li>3. Explain the physiology of lactation.</li> <li>4. Technique of breast feeding</li> <li>5. Problems in breast feeding</li> <li>6. Enumerate the baby friendly hospital initiatives</li> <li>7. Describe the composition and types of breast milk</li> <li>8. Discuss the differences between cow's milk and Human milk.</li> <li>9. Discuss the advantages of breast milk.</li> <li>10. Overview about expressed breast milk</li> </ol>                             | Obs&G ynae |
| 5. | Complementary Feeding  | 01 | Complementary feeding and IYCF (PE 8.1, 8.2, 8.3)       | <ol style="list-style-type: none"> <li>1. Define the term Complementary Feeding.</li> <li>2. Discuss the principles, the initiation, attributes, frequency, techniques and hygiene related to Complementary Feeding</li> <li>3. IYCF</li> <li>4. Enumerate the common complimentary foods</li> </ol>  |            |
| 6. | Provide nutritional support, assessment and monitoring for common nutritional problems | 01 | Protein Energy Malnutrition (PE 10.1, 10.2, 10.4, 10.6) | <ol style="list-style-type: none"> <li>1. Define malnutrition</li> <li>2. Classify malnutrition including WHO classification,</li> <li>3. Describe the etio-pathogenesis, clinical features, complication of Severe Acute Malnourishment (SAM) and Moderate Acute Malnutrition (MAM).</li> <li>4. Differentiate between kwashiorkor and marasmus</li> <li>5. Outline the clinical approach to a child with SAM an MAM.</li> <li>6. Management of a child with SAM and MAM.</li> <li>7. Enumerate the role of locally prepared therapeutic diets and ready to use therapeutic diets.</li> <li>8. Strategies to prevent malnutrition</li> </ol> |            |

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| 7. | Obesity in Children  | 01 | Obesity (PE 11.1, 11.2, 11.6)  | <ol style="list-style-type: none"> <li>1. Define obesity</li> <li>2. Describe the common etiology, clinical features and management of obesity in children.</li> <li>3. Discuss the risk approach for obesity and criteria for referral</li> <li>4. Discuss the prevention strategies</li> </ol>   |  |
| 8. | <p>Micronutrients in health and disease 1: (Vitamins A,D,E,K,B Complex and C)</p> <p>Micronutrients in health and disease 2: Iron, Iodine, Calcium and Magnesium</p> | 04 | <p>Vitamin A<br/>Vitamin E, K(PE 12.1,12.2, 12.4, 12.5, 12.11,12.12, 12.13,12.14)Vitamin B, C and Iodine deficiency disorders (PE 12.15, 12.16,12.18, 12.19,12.20, 13.7, 13.8, 13.10, 13.10)</p> | <p><b>Vitamin A</b></p> <ol style="list-style-type: none"> <li>1. RDA, dietary sources of Vitamin A and the role in Health and disease.</li> <li>2. Describe the causes, clinical features, diagnosis and management of Deficiency/excess of Vitamin A.</li> <li>3. Discuss the Vitamin A prophylaxis program and the recommendations</li> </ol> <p><b>Vitamin E</b><br/>Discuss the RDA, dietary sources of Vitamin E and their role in health and disease. Describe the causes, clinical features, diagnosis and management of deficiency of Vitamin E.</p> <p><b>Vitamin K</b></p> <ol style="list-style-type: none"> <li>1. Discuss the RDA, dietary sources of Vitamin K and their role in health and disease.</li> <li>2. Describe the causes, clinical features, diagnosis management and</li> </ol> <ol style="list-style-type: none"> <li>1. Prevention of deficiency of Vitamin K</li> </ol> <p><b>Vitamin B</b></p> <ol style="list-style-type: none"> <li>1. Discuss the RDA, dietary sources of Vitamin B and their role in health and disease</li> <li>2. Describe the causes, clinical features, diagnosis and management of deficiency of B complex Vitamins.</li> </ol> <p><b>Vitamin C</b></p> <ol style="list-style-type: none"> <li>1. Discuss the RDA, dietary sources of Vitamin C and their role in Health and disease</li> <li>2. Describe the causes, clinical features, diagnosis and management of</li> </ol> |  |

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|    |                                   |    |   | <p>deficiency of Vitamin C(scurvy)</p> <p><b>Iodine deficiency Disorder</b></p> <ol style="list-style-type: none"> <li>1. Discuss the RDA , dietary sources of Iodine and their role in Health and disease.</li> <li>2. Describe the causes, clinical features, diagnosis and management of deficiency of Iodine.</li> <li>3. Discuss the National Goiter Control program and their recommendations.</li> </ol>  |  |
|    |                                   |    | Iron deficiency anemia (PE 13.1, 13.2, 13.5, 13.6)  | <ol style="list-style-type: none"> <li>1. Discuss the RDA, dietary sources of Iron and their role in health and disease'</li> <li>2. Describe the causes, clinical features, diagnosis and management of Fe deficiency</li> <li>3. Discuss the National Anemia control program and its recommendations.</li> </ol>   |  |
|    |                                   |    | Vitamin D and Calcium & Magnesium deficiency (PE 12.6, 12.7, 12.9, 12.10, 13.11,13.12, 13.13,13.14) | <p><b>Vitamin D/Ca/Mg</b></p> <ol style="list-style-type: none"> <li>1. Discuss the RDA, dietary sources of Vitamin D and their role in health and disease.</li> <li>2. Describe the causes, clinical features,</li> </ol>   |  |
|    |                                   |    |   | <ol style="list-style-type: none"> <li>3. diagnosis and management of Deficiency/ excess of Vitamin D (Rickets and Hype vitaminosis D).</li> <li>3. Discuss the role of screening for Vitamin D deficiency</li> <li>4. Discuss the RDA ,dietary sources of Calcium and their role in health and disease</li> <li>5. Describe the causes, clinical features, diagnosis and management of Ca Deficiency</li> <li>6. Discuss the RDA, dietary sources of Magnesium and heirrolein health and disease.</li> <li>7. Describe the causes, clinical features, diagnosis and management of Magnesium Deficiency</li> </ol> |  |
| 29 | Anemia and other Hemato-oncologic | 02 | Anemia (PE 29.1)  | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiopathogenesis</li> <li>3. Classification</li> </ol>  |  |

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|    | disorders in children                                    |    |  | 4. Approach to a child with anemia  |  |
|    |  |    | Nutritional anemia (PE 29.2, 29.3, 29.5)                               | <b>Iron def anemia/ Megaloblastic anemia</b><br>1. Etiopathogenesis<br>2. Clinical features<br>3. Lab investigations<br>4. Management<br>5. Discuss the National Anemia Control Program   |  |
| 9. | Fluid and electrolyte balance                            | 01 | Fluid and electrolytes (PE 15.1, 15.2)                                 | 1. Composition of body fluids<br>2. Water balance and Osmolality<br>3. Normal maintenance fluid and electrolyte requirements<br>4. Sodium balance and its disorders<br>5. Potassium balance and its disorders<br>6. Overview of Acid-Base disorders   |  |
| 10 | National Programs, RCH - Universal Immunizations program | 02 | Vaccines in children (PE 19.1, 19.2, 19.3, 19.4)                       | 1. Components of the Universal Immunization Program and the National Immunization program.<br>2. Epidemiology of Vaccine preventable diseases<br>3. Vaccine description with regard to classification of vaccines, strain used, dose, route, schedule, risks, benefits and side effects, indications and contraindications. (BCG, OPV, IPV HepB, DPT, Hib, MMR)   |  |
|    |  |    | Immunization in special situations and newer vaccines (PE 19.5, 19.16) | 4. Define cold chain and discuss the methods of safe storage and handling of vaccines<br>1. Immunization in special situations- HIV positive children, immune deficiency, pre-term, organ transplants, those who received blood and blood products, splenectomised children, adolescents, travelers.<br>2. Enumerate available newer vaccines and their indications including pentavalent pneumococcal, rotavirus, JE, typhoid IPV & HPV.<br>3. Combination vaccines<br>4. AEFI |  |

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| 11 | Respiratory system                          | 02 | RTI GEM – I<br>(PE 28.1, 28.2, 28.3, 28.4, 28.5, 28.6, 28.7, 28.8))   | <b>Nasopharyngitis/Pharyngo Tonsillitis/ Acute Otitis Media (AOM)</b><br>1. Etio-pathogenesis<br>2. Clinical features<br>3. Management<br>4. Complications                     |  |
|    |   |    |   | <b>Stridor/ Epiglottitis/Acute laryngotracheobronchitis/Foreign Body Aspiration</b><br>1. Etiopathogenesis<br>2. Clinical features<br>3. Management                            |  |
|    |   |    | RTI GEM -II<br>(PE 28.18)   | <b>Bronchiolitis and wheeze associated LRTI/ Empyema/Lung Abscess</b><br>1. Etio-pathogenesis<br>2. Clinical features<br>3. Diagnosis<br>4. Management<br>5. Prevention        |  |
| 12 | Vaccine preventable Diseases & Tuberculosis | 02 | Fever & Exanthematous Fever<br>(PE 34.14, 34.15)  | 1. Enumerate the common causes of fever<br>2. Etiopathogenesis<br>3. Clinical features<br>4. Complications<br>5. Management<br>6. Approach to a child with Exanthematous Fever |  |
|    |   |    | Measles, Mumps, Rubella & Chicken pox<br>(PE 34.15)   | 1. Etiopathogenesis<br>2. Clinical features<br>3. Complications<br>4. Management<br>5. Prevention<br>6. Measles, Mumps, Rubella & Chicken poxvaccines                          |  |
| 13 | Chromosomal Abnormalities                   | 01 | Down syndrome, Turner & Klinefelters syndrome<br>(PE 32.1, 32.3, 32.4, 32.5, 32.6, 32.8, 32.9, 32.10, 32.11, 32.13) | 1. Geneticbasis<br>2. Riskfactors<br>3. Clinical features<br>4. Complications<br>5. Prenatal diagnosis<br>6. Management<br>7. Genetic counseling.                              | General Medicine - PE 32.3, 32.9<br>Obs & Gynae- PE 32.9 |

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| 14 | Diarrheal diseases and Dehydration                                  | 01 | Diarrheal diseases & dehydration incl Persistent diarrhea, Chronic diarrhea and dysentery (PE 24.1, 24.2, 24.3, 24.4, 24.5, 24.6, 24.7, 24.8, 24.14) | <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Classification</li> <li>3. Clinical presentation</li> <li>4. Management</li> <li>5. Physiological basis of ORT</li> <li>6. Types of ORS</li> <li>7. Composition of various types of ORS</li> <li>8. Classification and clinical presentation of various types of diarrheal dehydration</li> <li>9. Types of fluid used in Pediatric diarrheal diseases and their composition</li> <li>10. Role of antibiotics, antispasmodics, anti-secretory drugs, probiotics, anti-emetics in acute diarrheal diseases</li> </ol> |                  |
| 15 | Pediatric Emergencies - Common Pediatric Emergencies                | 02 | Poisoning (PE 27.8, 14.1, 14.2, 14.3, 14.4)  | <ol style="list-style-type: none"> <li>1. Clinical approach to a child with suspected poisoning</li> <li><b>2. Common poisonings - Hydrocarbon/OP/PCM/Lead/Envenomation</b></li> <li>3. Etiopathogenesis</li> <li>4. Clinical features</li> <li>5. Lab investigations</li> <li>6. Management</li> </ol>  | General Medicine |
|    |   |    | Child abuse (PE 27.29)   | <ol style="list-style-type: none"> <li>1. Causes</li> <li>2. Clinical presentation</li> </ol> <p>Medico-legal implications</p>   |                  |
| 16 | Allergic Rhinitis, Atopic Dermatitis,                               | 01 | Allergy in children (PE 31.1, 31.3, 31.12)   | <b>Allergic Rhinitis/Atopic Dermatitis/Urticaria Angioedema</b> <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Clinical features</li> </ol>  |                  |
|    | Bronchial Asthma, Urticaria Angioedema                              |    |  | <ol style="list-style-type: none"> <li>3. Management</li> <li>4. Complications</li> <li>5. Prevention</li> </ol>   |                  |
| 17 | Adolescent health and common problems related to Adolescent Health. | 01 | Adolescence & Puberty (PE 6.10, 6.11)  | 1. Visit to the Adolescent Clinic. Discuss the objectives and functions of AFHS (Adolescent Friendly Health Services) and the referral criteria.   | Psychiatry       |



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| 18           | Common problems related to Development-1 (Developmental delay, Cerebral palsy)                         | 01        | Developmental delay (PE 3.5,3.6, 3.7)   | 1. Visit a Child Developmental Unit and observe its functioning. Discuss the role of the child developmental unit in management of developmental delay. Discuss their feral criteria for children with developmental delay |  |
| 19           | Common problems related to Development-2 (Scholastic backwardness, Learning disabilities, Autism ADHD) | 01        | Scholastic backwardness and Learning Disabilities (LD) (PE 4.5, 4.6, 5.10,5.11) | 1. Visit to child guidance linic. Discuss the role of Child Guidance clinicin children with Developmental problems & Behavioral problems.  |  |
| <b>TOTAL</b> |  | <b>30</b> |   |  |  |

**Theory III (Part I) MBBS (20 hours)**

| S. No | Topic  | Hours | Lectures (Competency No)  | SLO  | Horizontal Integration             |
|-------|--|-------|---|--|------------------------------------|
| 1.    | Normal Growth and Development  | 01    | Growth & Development (PE1.1,1.2,1.3, 1.5)                           | <ol style="list-style-type: none"> <li>1. Definition of Growth</li> <li>2. Definition of Development</li> <li>3. Physiology of Growth &amp; Development</li> <li>4. Normal Growth-Somatic and physical</li> <li>5. Assessment of Growth</li> </ol>   | Psychiatry                         |
|       |  |       |   | <ol style="list-style-type: none"> <li>6. parameters; Growth charts</li> <li>7. Factors affecting Growth &amp; Development</li> <li>8. Overview of disorders related to Growth &amp; Development</li> </ol>  |                                    |
| 2.    | Common problems related to Development-1 (Developmental delay, Cerebral palsy)                         | 02    | Developmental delay (PE 3.1,3.2, 30.10)                             | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Developmental delay vs Intellectual disability</li> <li>3. Etiology</li> <li>4. Clinical Features</li> <li>5. Approach to developmental delay and ID</li> <li>6. Prevention and management</li> </ol>                                     |                                    |
|       |  |       | Cerebral palsy (PE 3.8, 30.11)                                      | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiopathogenesis</li> <li>3. Types of CP</li> <li>4. Evaluation of a child with CP</li> <li>5. Prevention and management</li> </ol>   | Physical Medicine & Rehabilitation |
| 3.    | Common problems related to Development-2 (Scholastic backwardness, Learning disabilities, Autism ADHD) | 02    | Scholastic backwardness and Learning Disabilities (LD) (PE 4.1,4.2) | <ol style="list-style-type: none"> <li>1. Causes of Scholastic backwardness</li> <li>2. Approach to a child with Scholastic backwardness</li> <li>3. Definition of LD</li> <li>4. Types of LD and clinical features</li> <li>5. Etiology</li> <li>6. Approach to a child with LD and Management</li> </ol> |                                    |

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|    |   |    | ADHD and Autism (PE 4.3,4.4)   | <ol style="list-style-type: none"> <li>1. Etiology of ADHD</li> <li>2. Clinical features of ADHD</li> <li>3. Diagnosis and management of ADHD</li> <li>4. Etiology of Autism</li> <li>5. Clinical features of Autism</li> <li>6. Diagnosis and management of Autism</li> </ol>  |            |
| 4. | Common problems related to behavior                                 | 01 | Behavioral problems of children include Enuresis & Encopresis (PE5.1,5.2,5.3, 5.4,5.5,5.6,5.7, 5.8, 5.9) | <ol style="list-style-type: none"> <li>1. Describe the clinical features, diagnosis and management of common behavioral problems like <ul style="list-style-type: none"> <li>• Thumb sucking,</li> <li>• Feeding problems,</li> <li>• Nail biting</li> <li>• Breath Holding spells,</li> <li>• Pica,</li> <li>• Fussy infant</li> </ul> </li> <li>2. Definition of enuresis and encopresis</li> <li>3. Differentiate between primary and secondary enuresis</li> <li>4. Maturation of bowel and bladder control</li> <li>5. Etiology of Enuresis and Encopresis</li> <li>6. Clinical features of Enuresis and Encopresis</li> <li>7. Management of Enuresis and Encopresis</li> </ol> | Psychiatry |
| 5. | Adolescent health and common problems related to Adolescent Health. | 01 | Adolescence & Puberty (PE6.1,6.2,6.3, 6.4,6.5,6.6,6.7, 6.12, 6.13)                                       | <ol style="list-style-type: none"> <li>1. Define Adolescence</li> <li>2. Stages of adolescence and SMR</li> <li>3. Describe the physical, physiological and psychological changes during adolescence and Puberty.</li> <li>4. Outline the general health problems during adolescence.</li> <li>5. Describe adolescent sexuality and common problems related to it.</li> <li>6. Explain the Adolescent Nutrition and common</li> </ol>   | Psychiatry |

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|    |  |    |   | <p>nutritional problems.</p> <ol style="list-style-type: none"> <li>Outline the common Adolescent eating disorders (Anorexia Nervosa, Bulimia).</li> <li>Describe the common mental health problems during adolescence.</li> <li>Enumerate the importance of obesity and other NCD in adolescents.</li> <li>Enumerate the prevalence and the importance of recognition of sexual drug abuse in adolescents and children.</li> </ol>   |   |
| 6. | Normal nutrition, assessment and monitoring. | 01 | Normal Nutrition (PE9.1,9.2,9.3, 9.7)   | <ol style="list-style-type: none"> <li>Describe the age-related nutritional needs of infants, children and adolescents including micronutrients and vitamins</li> <li>Concept of RDA and balanced diet.</li> <li>Describe the tools and methods for assessment and classification of nutritional status of infants, children and adolescents.</li> <li>Explains the Calorific value of common Indian foods</li> </ol>   |   |
| 7. | Vaccine preventable Diseases & Tuberculosis  | 8  | <p>Tuberculosis in children (PE 34.1, 34.2, 34.12, 34.13)</p> <p>Management of tuberculosis (PE 34.3, 34.4)</p> | <ol style="list-style-type: none"> <li>Epidemiology</li> <li>Clinical features and clinical types</li> <li>Complications of Tuberculosis</li> <li>Diagnostic tools for childhood tuberculosis.</li> <li>Indications and discuss the limitations of methods of culturing M. Tuberculosis.</li> <li>Newer diagnostic tools for Tuberculosis including BACTEC CBNAAT and their indications</li> </ol> <ol style="list-style-type: none"> <li>Various regimens for management of Tuberculosis as per National Guidelines.</li> <li>Preventive strategies adopted and the objectives and outcome of the National Tuberculosis Control Programme</li> </ol> | <p>Respiratory Medicine</p> <p>Respiratory Medicine</p> |

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|    |  |    | Diphtheria, Pertussis, Tetanus (PE 34.16)                  | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Complications</li> <li>4. Management</li> <li>5. Prevention</li> <li>6. Diphtheria, Pertussis, Tetanus vaccines</li> </ol> |  |
|    |  |    | Enteric fever (PE 34.17)                                   | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Complications</li> <li>4. Management</li> <li>5. Prevention</li> <li>6. Typhoid vaccines</li> </ol>                        |  |
|    |  |    | Rickettsial diseases(PE 34.20)                             | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Complications</li> <li>4. Management</li> <li>5. Prevention</li> </ol>   |  |
|    |  |    | Parasitic infections (PE 34.19)                            | <b>Common Parasitic infections - leishmaniasis, filariasis, helminthes infestations, amebiasis, giardiasis</b>  |  |
|    |  |    | Malaria (PE 34.19)   | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Complications</li> <li>4. Management</li> <li>5. Prevention</li> <li>6. National Malaria Eradication Programme</li> </ol>  |  |
|    |  |    | Dengue Fever (PE 34.18)                                    | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Complications</li> <li>4. Management</li> <li>5. Prevention</li> <li>6. Overview of Chikungunya</li> </ol>                 |  |
| 8. | Systemic Pediatrics-Central Nervous system | 01 | Acute Flaccid Paralysis (AFP) and Poliomyelitis (PE 30.13) | <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Approach to a child with</li> <li>3. AFP Evaluation</li> <li>4. Management</li> <li>5. AFP Surveillance</li> </ol>   |  |

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| 9.           | Endocrinology | 03        | Hypothyroidism<br>(PE 33.1)                                 | <ol style="list-style-type: none"> <li>1. Physiology of thyroidal</li> <li>2. and Thyroid function</li> <li>3. test Etiology</li> <li>4. Congenital vs</li> <li>5. Acquired Clinical</li> <li>6. features Evaluation</li> <li>7. Management</li> <li>8. New-born Screening</li> </ol> |  |
|              |               |           | Diabetes<br>mellitus in<br>children and<br>DKA<br>(PE 33.4) | <ol style="list-style-type: none"> <li>1. EtiopathogenesisD</li> <li>2. iagnostic criteria</li> <li>3. Classification</li> <li>4. Clinical features</li> <li>5. Management</li> <li>6. Complications incl DKA</li> </ol>  |  |
|              |               |           | Disorders of<br>puberty<br>(PE 33.8)                        | <p><b>Precocious and delayed Puberty</b></p> <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Clinical Features</li> <li>4. Evaluation</li> <li>5. Management</li> </ol>   |  |
| <b>TOTAL</b> |               | <b>20</b> |   |   |  |

**Self-Directed Learning III (Part I) MBBS (05 hours)**

| S. No        | Topic  | Hours     | Lectures (Competency No.)   | SLO   | Horizontal Integration |
|--------------|--|-----------|---|---|------------------------|
| 1.           | The National Health Programs, NHM<br>The National Health Programs, RCH | 02        | National programs pertaining to maternal & child health, child survival & safe motherhood (PE 17.1, 17.2, 18.1, 18.2) | <ol style="list-style-type: none"> <li>1. State the vision and outline the goals, strategies and plan of action of NHM and other important national programs pertaining to maternal and child health including RMNCH A+, RBSK, RKSK, JSSK mission India Dhanush and ICDS.</li> <li>2. List and explain the components, plan, outcome of Reproductive Child Health(RCH)program and appraise its monitoring and evaluation</li> <li>3. Explain preventive interventions for child survival and safe motherhood</li> </ol> | Obs&G<br>ynaec         |
| <b>TOTAL</b> |  | <b>02</b> |   |   |                        |

**Tutorials/Small Group Discussions III (PartII) MBBS (35 hours)**

| S. No | Topic             | Hours | Domain (Competency No.)  | SLO  | Horizontal Integration |
|-------|-------------------|-------|--|--|------------------------|
| 1     | Group Discussions | 01    | Fluids & Electrolytes, Nutrition<br>(PE 15.3, 15.4, 15.5, 9.5) | <ol style="list-style-type: none"> <li>1. Calculate fluid and electrolyte imbalance, Interpret electrolyte report,</li> <li>2. Calculate the fluid and electrolyte requirement in health</li> <li>3. Plan an appropriate diet in health &amp; disease</li> </ol>   |                        |
|       |                   | 01    | Cardiac Failure<br>(PE23.11,23.16,23.17, 23.18)                | <ol style="list-style-type: none"> <li>1. Develop treatment plan and prescribe appropriate drugs including fluids in cardiac diseases, anti -failure drugs, and intropicagents.</li> <li>2. Discuss the indications and limitations of Cardiac catheterization.</li> <li>3. Enumerate some common cardiac surgeries like BT shunt, Potts and Waterston's and corrective surgeries</li> <li>4. Demonstrate empathy while dealing with cardiac disease.</li> </ol> |                        |
|       |                   | 01    | Oxygen Therapy<br>(PE 27.9, 27.10, 14.5)                       | <ol style="list-style-type: none"> <li>1. Discuss oxygen therapy in Pediatric emergencies and modes of administration.</li> <li>2. Observe the various methods of administering Oxygen.</li> <li>3. Discuss oxygen toxicity and free radical injury</li> </ol>   |                        |



|    |           |    |  |   |  |
|----|-----------|----|--|---|--|
|    |           | 01 | Counseling<br>(PE2.3,3.4,8.5,27.32,<br>27.33, 28.20)                                   | <ol style="list-style-type: none"> <li>1. Counseling apparent with failing to thrive child</li> <li>2. Counselling a parent with developmental delay</li> <li>3. Counsel &amp; educate mothers on the best practices in complimentary feeding</li> <li>4. Obtain Informed Consent.</li> </ol> |  |
|    |           |    |  | <ol style="list-style-type: none"> <li>5. Counsel parents of dangerously ill/terminally ill child to break bad news</li> <li>6. Counsel the child with asthma on the correct use of inhalers in a simulated environment</li> </ol>  |  |
|    |           | 01 | Hemat<br>(PE 29.18,29.20)  | <ol style="list-style-type: none"> <li>1. Enumerate the referral criteria for Hematological conditions.</li> <li>2. Enumerate the indications for splenectomy and precautions</li> </ol>  |  |
| 2. | Radiology | 01 | X-Ray/USG/Neuroimaging<br>(PE 21.12,21.13,<br>23.12, 26.9,28.17,<br>30.21, 30.22,31.9, | <ol style="list-style-type: none"> <li>1. Interprêt report of Plain X Ray of KUB</li> <li>2. Enumerate the indications for and Interpret the written report of Ultra</li> </ol>   |  |

|    |                             |    |   |   |  |
|----|-----------------------------|----|---|---|--|
|    |                             |    | 34.8)   | <p>sonogram of KUB</p> <ol style="list-style-type: none"> <li>3. Interpret a chest X ray and recognize Cardiomegaly</li> <li>4. Interpret Liver USG</li> <li>5. Interpret X-ray of the paranasal sinuses and mastoid; and /or use written report in case of management</li> <li>6. Interpret CX R in foreign body aspiration and lower respiratory tract infection, understand the significance of thymic shadow in pediatric chest X-rays</li> <li>7. Enumerate the indication and limitations &amp; Interpret the reports of CT, MRI Brain &amp; Spine</li> <li>8. Interpret CX Ray in Asthma</li> <li>9. Interpret a Chest Radiographn pediatric TB</li> </ol> |  |
| 3. | Cards (Case Scenario based) | 01 | (PE21.11,23.13,23.14. 24.13, 26.9,26.11, 28.16, 29.14, 19.15, 29.16, 30.20, 30.21, 30.22,33.3,33.6,33.9, 34.9, 34.10) | <ol style="list-style-type: none"> <li>1. Interpret Hemogram and Iron Panel</li> <li>2. interpret the common analyses in a Urine examination</li> <li>3. Interpret Pediatric ECG</li> <li>4. Choose and Interpret blood reports in Cardiac illness</li> <li>5. Interpret RFT and electrolyte report</li> <li>6. Interpret Liver Function Tests,</li> </ol>  |  |

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|--|--|--|--|--|--|
|  |  |  |  | <p>viral markers.</p> <ol style="list-style-type: none"> <li>7. Enumerate indications of UGI Endoscopy</li> <li>8. Interpret blood tests relevant to upper respiratory problems.</li> <li>9. Interpret CBC, LFT in anemia</li> <li>10. Perform and interpret peripherals mear</li> <li>11. Discuss the indications for Hemoglobin electrophoresis and interpret report</li> <li>12. Interpret and explain the findings in a CSF analysis</li> <li>13. Interpret and explain neonatal thyroid screening report</li> <li>14. Perform and interpret Urine Dip Stick for Sugar. Interpret Blood sugar reports and explain the diagnostic criteria for Type 1 Diabetes</li> <li>15. Interpret the reports of EEG</li> <li>16. Perform Sexual Maturity Rating (SMR) and interpret</li> <li>17. Interpret blood tests in the context of laboratory evidence for tuberculosis. Discuss the various samples for demonstrating the organism e.g. Gastric Aspirate, Sputum, CSF, FNAC.</li> </ol> |  |
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|----|------------|----|--|---|-------------------|
| 4. | Skills Lab | 02 | (PE 15.6, 15.7, 19.9, 19.13, 20.3,24.15, 24.16, 24.17,26.10, 27.20, 29.17,30.23) | <ol style="list-style-type: none"> <li>1. Demonstrate the steps of inserting an IV cannula in a model</li> <li>2. Demonstrate the steps of inserting an interosseousline in amannequin</li> <li>3. Demonstrate the correct administration of different vaccines in a mannequin.</li> <li>4. Describe the components of safe vaccine practice-Patient education/ counselling; adverse events following immunization, safe injection practices, documentation and Medico-legal implications</li> <li>5. Perform Neonatal resuscitation</li> </ol> | AETCOM<br>-PE19.9 |
|    |            |    |  | <ol style="list-style-type: none"> <li>in a manikin</li> <li>6. Perform NG tube insertion in a manikin</li> <li>7. Perform IV cannulation in a model</li> <li>8. Demonstrate the technique of liver biopsy or perform Liver Biopsy in a simulated environment.</li> <li>9. Demonstrate performance of bone marrow aspiration in manikin</li> <li>10. Perform in a mannequin lumbar puncture. Discuss the indications, contraindication of the procedure</li> </ol>  |                   |

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|----|---|----|--|---|--|
| 5. | Genit<br>o-<br>Urinary<br>system        | 02 | Hypertension in children (PE 21.17)                          | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiopathogenesis</li> <li>3. Grading</li> <li>4. Clinical features</li> <li>5. Management</li> <li>6. Complications</li> <li>7. Acute severe hypertension</li> </ol>   |  |
|    |   |    | Voiding Disorders (PE 21.15)                                 | <ol style="list-style-type: none"> <li>1. Discuss &amp; Enumerate the referral criteria for children with genitor urinary disorder</li> <li>2. Counsel &amp; educate patients regarding referral</li> </ol>   |  |
| 6. | Cardiovascular system:<br>Heart disease | 04 | Congestive cardiac failure in infants and children (PE 23.3) | <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Pathogenesis</li> <li>3. Clinical presentation</li> <li>4. Management</li> </ol>   |  |
|    |   |    | Acyanotic congenital heart diseases (PE 23.1)                | <b>VSD, ASD and PDA</b> <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Hemodynamic changes</li> <li>3. Clinical features</li> <li>4. Investigations</li> <li>5. Management</li> </ol>   |  |
|    |   |    | Cyanotic congenital heart diseases (PE 23.2)                 | <ol style="list-style-type: none"> <li>1. Classify Cyanotic congenital heart disease</li> </ol> <b>Fallot's Physiology</b> <ol style="list-style-type: none"> <li>2. Etiology</li> <li>3. Hemodynamic changes</li> <li>4. Clinical features</li> <li>5. Investigations</li> </ol>   |  |
|    |   |    |  | 6. Management   |  |
|    |   |    | Acquired Heart Disease (PE 23.4, 23.5, 23.6)                 | <b>Infective endocarditis</b> <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Diagnosis</li> <li>4. Management</li> </ol> <b>Acute rheumatic fever</b> <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Diagnosis</li> <li>4. Management and prevention</li> <li>5. Complications</li> </ol> |  |

|    |  |    |  |  |  |
|----|--|----|--|--|--|
| 7. | Pediatric Emergencies - Common Pediatric Emergencies | 03 | Shock in children (PE 27.5)                            | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. BP regulation</li> <li>3. Pathophysiology</li> <li>4. Classification</li> <li>5. Monitoring</li> <li>6. Management</li> </ol>   |  |
|    |  |    | Status epilepticus(PE 27.6, 30.9)                      | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Approach to a child with status epilepticus</li> <li>4. Evaluation</li> <li>5. Management</li> </ol>   |  |
|    |  |    | Unconscious child and Coma (PE 27.8)                   | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiopathogenesis</li> <li>3. Evaluation</li> <li>4. Management</li> <li>5. Brain death</li> </ol>   |  |
| 8. | Care of the Normal Newborn, and High-risk Newborn    | 04 | Care of low birth weight (LBW) babies (PE 20.11)       | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Explain the terminologies- IUGR/SGA</li> <li>4. Clinical features</li> <li>5. Issues in LBW care</li> <li>6. Feeding in LBW babies</li> <li>7. Management of LBW babies</li> <li>8. Growth monitoring of LBW babies</li> </ol> |  |
|    |  |    | Neonatal hypoglycemia & hypocalcaemia (PE 20.13,20.14) | <b>Hypoglycemia and hypocalcaemia</b> <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Clinical features</li> <li>4. Management</li> </ol>  |  |
|    |  |    | Neonatal Seizures (PE 20.15)                           | <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Clinical features</li> <li>3. Management</li> </ol>   |  |
|    |  |    | Perinatal infections (PE 20.17)                        | TORCH/Tuberculosis/HepB/Va ricella <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Transmission</li> <li>3. Clinical features</li> <li>4. Management</li> </ol>   |  |

|     |   |    |  |  |  |
|-----|---|----|--|--|--|
| 9.  | Anemia and other Hemato-oncologic disorders in children | 02 | Hemolytic anemia (PE 29.4)   | <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Classification</li> <li>3. Approach to a child with hemolytic anemia</li> <li>4. Management</li> <li>5. Overview of HS, AIHA and HUS</li> </ol>   |  |
|     |   |    | Thalassemia and Sickle Cell Anemia (PE 29.4)                           | <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Clinical features</li> <li>3. Lab investigations</li> <li>4. Management in cl Iron Chelation therapy</li> <li>5. Complications</li> </ol>   |  |
| 10. | Acute and chronic liver disorders                       | 02 | Acute liver disease & Fulminant hepatic failure (PE 26.1, 26.2)        | <b>Acute hepatitis in children-Viral (Hep A,B,C), Autoimmune and Wilsons disease</b> <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Management</li> </ol> <b>Fulminant Hepatic Failure in children</b> <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Management</li> </ol>  |  |
|     |   |    | Chronic liver disease & Portal hypertension (PE26.3,26.4,26.11, 26.12) | <b>Chronic liver diseases in children</b> <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Evaluation</li> <li>4. Complications - hepatic encephalopathy and ascites</li> <li>5. management</li> </ol> <b>Portal Hypertension in children</b> <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Management</li> <li>4. Complications</li> </ol> |  |
| 11. | Respiratory system                                      | 01 | Pneumonia and ARDS (PE 27.3,27.4)                                      | <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical features</li> <li>3. Diagnosis</li> <li>4. Management</li> </ol>  |  |
|     |   |    |  | 5. Prevention  |  |

|              |                |           |                          |  |  |
|--------------|----------------|-----------|--------------------------|--|--|
| 4.           | Mal absorption | 01        | Mal absorption (PE 25.1) | <ol style="list-style-type: none"> <li>1. Etio-pathogenesis</li> <li>2. Clinical presentation</li> <li>3. Management</li> <li>4. Overview of celiac disease</li> </ol> |  |
| <b>TOTAL</b> |                | <b>28</b> |                          |  |  |

### Theory III (Part II) MBBS (20 hours)

| S. No | Topic   | Hours | Lectures (Competency No.)  | SLO   | Horizontal Integration |
|-------|---|-------|--|---|------------------------|
| 1.    | Care of the Normal Newborn, and High-risk Newborn | 05    | Birth asphyxia (PE 20.7)   | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Clinical features</li> <li>4. Management</li> <li>5. Prevention</li> </ol>  |                        |
|       |   |       | Respiratory distress in newborn (PE 20.8)                              | <b>RDS/TTNB/MAS</b> <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Clinical features inclscoring systems</li> <li>3. Management</li> </ol>  |                        |
|       |   |       | Birth injuries & Hemorrhagic disease of newborn (HDN) (PE 20.9, 20.10) | <b>Birth Injuries</b> <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Clinical features</li> <li>3. Management</li> </ol> <b>HDN</b> <ol style="list-style-type: none"> <li>1. Definition and classification</li> <li>2. Etiology</li> <li>3. Clinical features</li> <li>4. Management</li> <li>5. Prevention</li> </ol> |                        |
|       |   |       | Neonatal Sepsis (PE 20.16)   | <ol style="list-style-type: none"> <li>1. Classification</li> <li>2. Etiology</li> <li>3. Clinical features</li> <li>4. Investigations</li> <li>5. Management</li> </ol>  |                        |



|    |  |    |  |   |  |
|----|--|----|--|---|--|
|    |  |    | Surgical conditions in newborn (PE 20.20)                                  | TEF, esophageal atresia, anal atresia, cleft lip and palate, congenital diaphragmatic hernia<br>1. Etiology<br>2. Clinical presentation<br>3. Management<br>4. Causes of acute abdomen in neonates  |  |
| 2. | Genito-Urinary system  | 03 | UTI (PE 21.1)  | 1. Etiology and predisposing factors<br>2. Clinical features<br>3. Diagnosis<br>4. Management<br>5. VUR   |  |
|    |  |    | Approach to hematuria & Acute glomerulonephritis (PE 21.2, 21.4)           | Hematuria<br>1. Definition<br>2. Diagnostic evaluation<br>3. Referral criteria<br>Acute Glomerulonephritis<br>1. Definition<br>2. Etiology<br>3. Clinical features of PSGN<br>4. Management of PSGN<br>5. Complications   |  |
|    |  |    | Acute kidney injury (AKI) and Chronic kidney disease (CKD) (PE 21.5, 21.6) | 1. Definition and classification<br>2. Etiology and pathophysiology<br>3. Approach to a child with AKI<br>4. Management<br>5. Complications<br>6. Renal replacement therapy   |  |
| 3. | Approach to and recognition of a child with possible rheumatologic problem | 02 | Approach to Rheumatological Problems incl JIA and SLE (PE 22.1)            | 1. Enumerate the common Rheumatological problems in children.<br>2. Approach to a child with arthritis<br>3. Referral criteria for a child with possible rheumatologic problem<br><b>JIA/SLE</b><br>1. Definition<br>2. Etiopathogenesis<br>3. Clinical sub types/Clinical features |  |

|    |                                 |    |   |   |  |
|----|---------------------------------|----|---|---|--|
|    |                                 |    |   | 4. Diagnosis<br>5. Management   |  |
|    |                                 |    | Vasculitic disorders in children (PE 22.3)                              | Enumerate common Vasculitic disorders in children and its classification<br><b>Kawasaki disease/HSP</b><br>1. Etiology<br>2. Clinical features<br>3. Diagnosis<br>4. Management |  |
| 4. | Anemia and other Hemato-        | 02 | Thrombocytopenia and Hemophilia (PE 29.6, 29.7)                         | <b>Thrombocytopenia</b><br>1. Causes of thrombocytopenia<br>2. Etiology of ITP<br>3. Clinical features and management of ITP  |  |
|    | oncologic disorders in children |    |   | <b>Hemophilia</b><br>1. Approach to a child with bleeding disorder<br>2. Etiology and types of hemophilia<br>3. Clinical features and management of hemophilia                  |  |
|    |                                 |    | Leukemia, Lymphomas and Solid Tumors in children (PE 29.8, 29.9, 21.17) | <b>ALL/Lymphoma/Wilm's Tumor</b><br>1. Etiology<br>2. Clinical features<br>3. Management  |  |

|    |   |    |  |   |  |
|----|---|----|--|---|--|
| 5. | Systemic Pediatrics- Central Nervous system | 08 | Meningitis in children (PE 30.1, 30.2)               | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Lab investigations</li> <li>4. Management</li> <li>5. Prevention</li> <li>6. Differentiate between Bacterial, Viral and TB Meningitis</li> <li>7. Approach to a child with acute febrile encephalopathy</li> </ol> |  |
|    |   |    | Hydrocephalus (PE 30.3)                              | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Investigations</li> <li>4. Complications</li> <li>5. Management</li> <li>6. Overview of IIH</li> </ol>   |  |
|    |   |    | Microcephaly and Neural tube defects (PE 30.4, 30.5) | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Classification/Types</li> <li>3. Clinical features</li> <li>4. Complications</li> <li>5. Management</li> </ol>   |  |
|    |   |    | Infantile hemiplegia/ Stroke (PE 30.6)               | <ol style="list-style-type: none"> <li>1. Etiopathogenesis</li> <li>2. Clinical features</li> <li>3. Investigations</li> <li>4. Management</li> </ol>   |  |
|    |   |    | Epilepsy in children (PE 30.8)                       | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Pathogenesis</li> <li>3. Types of Epilepsy</li> <li>4. Clinical presentation</li> <li>5. Management</li> <li>6. Overview of status epilepticus</li> </ol>  |  |
|    |   |    | Muscular dystrophy (PE 30.14)                        | <b>DMD/BMD</b>  |  |
|    |   |    | Ataxia in children(PE 30.15)                         | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Etiology</li> <li>3. Clinical features</li> <li>4. Differential Diagnosis</li> <li>5. Management</li> </ol>  |  |

|              |  |           |   |   |  |
|--------------|--|-----------|---|---|--|
|              |  |           | Approach to headache in children (PE 30.16) | <ol style="list-style-type: none"> <li>1. Pathophysiology of headache</li> <li>2. Approach to a child with headache</li> <li>3. Types of Headache</li> <li>4. Management</li> </ol> |  |
| <b>TOTAL</b> |  | <b>20</b> |   |   |  |

### **Self-Directed Learning III (Part II) MBBS (10 hours)**

| S. No | Topic   | Hours | Lectures (Competency No.)              | SLO   | Horizontal Integration |
|-------|---|-------|--|---|------------------------|
| 1.    | Systemic Pediatrics- Central Nervous system       | 04    | Floppy infant (PE 30.12)               | <ol style="list-style-type: none"> <li>1. Etiology</li> <li>2. Clinical features</li> <li>3. Differential diagnosis</li> <li>4. Evaluation</li> <li>5. Management</li> </ol>  |                        |
|       |   |       | Febrile seizures (PE 30.7)             | <ol style="list-style-type: none"> <li>1. Definition</li> <li>2. Types</li> <li>3. Etio pathogenesis</li> <li>4. Clinical features</li> <li>5. Investigations</li> <li>6. Complications</li> <li>7. Management</li> </ol>                   |                        |
| 2.    | Care of the Normal Newborn, and High-risk Newborn | 02    | Neonatal Hyperbilirubinemia (PE 20.19) | <ol style="list-style-type: none"> <li>1. Physiological vs pathological jaundice</li> <li>2. Etiology</li> <li>3. Clinical features</li> <li>4. Approach to a neonate with jaundice</li> <li>5. Management</li> <li>6. Follow-up</li> </ol> |                        |

|              |                       |           |   |   |  |
|--------------|-----------------------|-----------|---|---|--|
| 3.           | Genito-Urinary system | 02        | Approach to Proteinuria & Nephrotic syndrome (PE 21.3)        | Proteinuria<br>1. Definition<br>2. Diagnostic evaluation<br><br>3. Referral criteria<br>Nephrotic Syndrome<br>1. Definition<br>2. Etiology<br>3. Terminologies - Remission/Relapse/Steroid dependence/Steroid resistance<br>4. Clinical features<br>5. Management<br>6. Complications<br>7. SDNS/SRNS/Congenital nephritic syndrome |  |
| 4.           | Respiratory system    | 02        | Asthma in children (PE 28.19, 28.20, 31.5, 31.7, 31.8, 31.10) | 1. Pathophysiology incl Triggers<br>2. Clinical features<br>3. Diagnosis and differential diagnosis<br>4. Management<br>5. Inhalational therapy<br>6. Monitoring and modification of treatment<br>8. Management of acute exacerbation of bronchial asthma   |  |
| <b>TOTAL</b> |                       | <b>10</b> |   |   |  |

**Internal Assessment****Subject - Pediatrics**

**Applicable w.e.f October 2020 onwards examination for batches admitted from June 2019 onwards**

| Phase       |        |   |
|-------------|--------|---|
|             | Theory | Practical   |
| Second MBBS | -      | EOP Practical Examination may be conducted. However, these marks shall not be added to the Internal Assessment. |

| <b>3<sup>rd</sup> Year (III MBBS, PART I)</b> |        |           |             |         |           |             |
|---|--------|-----------|-------------|---------|-----------|-------------|
| Phase   | I-Exam |           |             | II-Exam |           |             |
|   | Theory | Practical | Total Marks | Theory  | Practical | Total Marks |
| III/I MBBS                                    | 50     | 50        | 100         | 50      | 50        | 100         |

| <b>4<sup>th</sup> Year (III MBBS, PART II)</b>                                      |          |           |             |                                   |           |             |
|---|----------|-----------|-------------|-----------------------------------|-----------|-------------|
| <b>Clinical posting- 4weeks</b>   |          |           |             |                                   |           |             |
| <b>Theory- lectures- 20, tutorials- 35, self-directed learning-10. Total 65 hrs</b> |          |           |             |                                   |           |             |
| Phase   | III-Exam |           |             | IV-Exam (Preliminary examination) |           |             |
|   | Theory   | Practical | Total Marks | Theory                            | Practical | Total Marks |
| III/II MBBS   | 50       | 50        | 100         | 100                               | 100       | 200         |

**Assessment in CBME is ON GOING PROCESS,  
No Preparatory leave is permitted.**

1. There shall be 4 internal assessment examinations in Pediatrics including Prelim.
2. The suggested pattern of question paper for internal assessment examinations, except prelim examination is attached at the end. Pattern of the prelims examinations should be similar to the University examinations.
3. Internal assessment marks for theory and practical will be converted to out of 25 (theory) + 25 (practical). Internal assessment marks, after conversion, should be submitted to university within the stipulated time as per directives from the University. **Conversion Formula for calculation of marks in internal assessment examinations.**

|  | Theory                                     | Practical                                     |
|--|--|---|
| <b>Phase II</b>                              | -  | -   |
| <b>Phase III/I</b>                           | 100  | 100   |
| <b>Phase III/II</b>                          | 150  | 150   |
| <b>Total</b>                                 | 250  | 250   |
| <b>Conversion out of</b>                     | 25   | 25  |
| <b>Conversion formula</b>                    | Total marks in 4 IA theory examinations/10 | Total marks in 4 IA Practical examinations/10 |
| <b>Eligibility criteria after conversion</b> | <b>10</b>                                  | <b>10</b>                                     |
|  | <b>Combined theory + Practical=25</b>      |   |

1. While preparing Final Marks of Internal Assessment, the rounding-off marks shall done as illustrated in following table.

| Total Internal Assessment Marks | Final rounded marks |
|---------------------------------|---------------------|
| 13.01 to 13.49                  | 13                  |
| 13.50 to 13.99                  | 14                  |

2. Students must secure atleast 50% marks of the total marks (combined in theory and practical / clinical; not less than 40 % marks in theory and practical separately) assigned for internal assessment in order to be eligible for appearing at the final University examination of that subject.
3. Internal assessment marks will not to be added to marks of the University examinations and will be shown separately in marklist.
4. Remedial measures

#### **A. Remedial measures for non - eligible students**

- i) At the end of each internal assessment examination, student securing less than 50% marks shall be identified. Such students should be counseled at the earliest and periodically. Extra classes for such students may be arranged.
- ii) If majority of the students found to be weak in a particular area then extra classes must be scheduled for all such students. Even after these measures, if a student is failed to secure 50% marks combined in theory and practical (40% separately in theory and practical) after prelim examination, the student shall not be eligible for final examination.
- iii) Non eligible candidates are offered to reappear for repeat internal assessment examination/s, which must be conducted 2 months before next University examination. The pattern for this repeat internal assessment examination shall be similar to the final University examination. The marks in this examination shall be considered for deciding the eligibility criteria. Following conversion formula shall be used for converting the marks.



|  | <b>Theory</b>                                   | <b>Practical</b>                                   |
|--|---|--|
| <b>Remedial examination</b>                  | <b>100</b>                                      | <b>100</b>   |
| <b>Conversion out of</b>                     | <b>25</b>                                       | <b>25</b>  |
| <b>Conversion formula</b>                    | <b>Marks in remedial theory examinations /4</b> | <b>Marks in remedial Practical examinations /4</b> |
| <b>Eligibility criteria after conversion</b> | <b>10</b>                                       | <b>10</b>  |
|  | <b>Combined theory + Practical = 25</b>         |  |

### **B. Remedial measures for absent students:**

- If any of the students is absent for any of the 4IA examinations due to any reasons, following measures shall be taken.
- i. The student is asked to apply to the academic committee of the college for reexamination, through HOD, to ascertain the genuineness of the reason for absentee.
  - ii. If permitted by academic committee, an additional examination for such students is to be conducted after prelims examination. Marks for such additional examination shall be equal to the missed examination.
  - iii. Even if a student has missed more than one IA examination, he/she can appear for only one additional IA examination. In such scenario, eligibility should be determined by marks obtained in internal assessment examinations for which the candidate has appeared, without changing the denominator.

**Internal Assessment Practical Examinations**  
**Pediatrics**  
**Internal Assessment Practical - I, II and III**

| Subject: Pediatrics Practical (IA - I, II and III) |        |        |                    |                       |
|--|--------|--------|--------------------|-----------------------|
| Case   | OSCE 1 | OSCE 2 | Journal & log book | Practical Total marks |
| 20   | 10     | 10     | 10                 | 50                    |
|  |        |        |                    |                       |

#OSCE Stations to include Signs of General examinations, Local examinations, Psychomotor skills and Communication skills., history taking of a particular symptom; nutrition history, developmental history, immunization history.

**Prelim Practical**

| Subject: Pediatrics Practical (Prelims)                |  |                                      |      |                    |                       |
|--|--|--------------------------------------|------|--------------------|-----------------------|
| Long Case<br>(Including clinical skills demonstration) | Short Case<br>(Including communication skills) | OSCE<br>(4 stations x 10 marks each) | Viva | Journal & log book | Practical Total marks |
| 25   | 15   | 40                                   | 10   | 10                 | 100                   |
|  |  |                                      |      |                    |                       |

OSCE1-ClinicalSkills, OSCE2-Anthropometry assessment ,OSCE3-Certifiable procedural skills , OSCE 4 - AETCOM related skills

**MUHS Final Practical**

| Subject: Pediatrics Practical (Prelims)                |  |                                      |      |                       |
|--|--|--------------------------------------|------|-----------------------|
| Long Case<br>(Including clinical skills demonstration) | Short Case<br>(Including communication skills) | OSCE<br>(4 stations x 10 marks each) | Viva | Practical Total marks |
| 30   | 20   | 40                                   | 10   | 100                   |
|  |  |                                      |      |                       |

OSCE1-ClinicalSkills, OSCE2-Anthropometry assessment, OSCE3-Certifiable procedural skills , OSCE 4 - AETCOM related skills

**Internal Assessment Examination (I, II and III) Pediatrics****Question SECTION "A" MCQ (10 Marks)****No.**

1. Multiple Choice Questions (Total-10 MCQ of One mark each from General Medicine) (10x1=10)

a)    b)    c)    d)    e)    f)    g)    h)    i)    j)

**SECTION "B" (40 Marks)**

2. Short Answer Questions (Five marks each) (5x5= 25 )  
(Any 5 out of 6)

a)    b)    c)    d)    e)    f)

3. Long answer question (1x15= 15)  
a)

**Final Theory Examination  
FORMAT / SKELETON OF QUESTION PAPER**

**Question SECTION "A" MCQ 20 Marks)**

**No.**

1. Multiple Choice Questions (Total-20 MCQ of One mark each) (20x1=20)

a)      b)      c)      d)      e)      f)      g)      h)      i)      j)

l)      m)      n)      o)      p)      q)      r)      s)      t)      u)

**SECTION "B" (45 Marks)**

2. Long Answer Questions (any 2 out of 3) (Structured clinical questions) (2x15= 30 )

a)      b)      c)

3. Short Answer Questions (All 3) (including 1 on ARTCOM) (3x5= 15 )

a)      b)      c)

**SECTION "C"- Allied (35 Marks)**

4. Long answer questions (1x15= 15)  
a)

5. Short Answer Questions (4x5= 20)  
(Any 4 out of 5) (Clinical reasoning)

a)      b)      c)      e)      f)

**Annexure- 1**  
**Course Content Phase II( October 2020)**  
**Subject: PAEDIATRICS**  
**Theory / Practical**

(Based on National Medical Council of India, Competency based Undergraduate curriculum for the Indian Medical Graduate, 2018. Vol. 2 / 3.)

1. Total Teaching hours:
  - A. Lectures(hours): **No**
  - B. Self-directed learning(hours):
  - C. Clinical Postings(hours):
    - Weeks- 2wks
    - Hours perweek-15
    - Monday to Friday- 3 hours per day.
  - D. Small group teachings/tutorials/Integrated teaching/Practical(hours):No

**Tentative Clinical posting schedule-**

| Day | Topic   | Day | Topic                                    |
|-----|---|-----|--|
| 1   | Round to Paediatric ward, Maternal ward, Kangaroo Mother Care, PICU, NICU, Labour room, OPD, Immunization room etc. | 6   | Systemic examination of child- CVS n     |
| 2   | History taking in Paediatrics   | 7   | Systemic examination of child- RS and PA |
| 3   | Assessment of growth and development  | 8   | Neonatal examination                     |
| 4   | General examination of child.   | 9   | Elicitation of neonatal reflexes         |
| 5   | Systemic examination of child- CNS  | 10  | Posting ending exam                      |

| Competency Nos. | Topics, Subtopics and Lectures |
|-----------------|--------------------------------|
|                 |                                |

## Annexure- 2.

### Course Content Phase III-I( October 2020)

#### Subject: PAEDIATRICS (Theory/Practical)

(Based on National Medical Council of India, Competency based Undergraduate curriculum for the Indian Medical Graduate, 2018. Vol. 2 / 3.)

Total Teaching hours:

- A. Lectures (hours):20
- B. Self-directed learning (hours): 5
- C. Clinical Postings(hours):
  - Weeks-4
  - Hours per week-15
  - Monday to Friday- 3 hours per day.
- D. Small group teachings/tutorials/Integrated teaching/Practical (hours): 30

#### Tentative Clinical posting schedule-

| Day | Topic   | Day | Topic                            |
|-----|---|-----|----------------------------------|
| 1   | Round to Paediatric ward, Maternal ward, Kangaroo Mother Care, PICU, NICU, Labour room, OPD, Immunization room etc. | 11  | Elicitation of neonatal reflexes |
| 2   | History taking in Paediatrics   | 12  | Immunization clinic              |
| 3   | Assessment of growth and development  | 13  | Immunization clinic              |
| 4   | General examination of child.   | 14  | Immunization clinic              |
| 5   | Systemic examination of child- CNS  | 15  | Immunization clinic              |
| 6   | Systemic examination of child- CNS  | 16  | Paediatric Emergencies           |
| 7   | Systemic examination of child- RS   | 17  | Paediatric Emergencies           |
| 8   | Systemic examination of child- Per Abdomen  | 18  | Paediatric Emergencies           |
| 9   | Systemic examination of child- CVS  | 19  | Paediatric Emergencies           |
| 10  | Neonatal case taking and examination.   | 20  | Posting ending exam              |

| Competency Nos. | Topics, Subtopics and Lectures |
|-----------------|--------------------------------|
|                 |                                |

### Annexure- 3.

**Course Content Phase: III-II (October 2020)**

**Subject: PAEDIATRICS (Theory / Practical )**

(Based on National Medical Council of India, Competency based Undergraduate curriculum for the Indian Medical Graduate, 2018. Vol. 2 / 3.)

Total Teaching hours:

A. Lectures(hours): 20

B. Self-directed learning (hours): 10

C. Clinical Postings(hours):

- Weeks-4
- Hours per week-15
- Monday to Friday- 3 hours per day.

D. Seminars/Small group teachings/tutorials/Integrated teaching/Practical (hours):35

#### Tentative Clinical posting schedule-

| Day | Topic  | Day | Topic  |
|-----|--|-----|--|
| 1   | History taking and General examination of child.                       | 11  | Neonatal case taking, examination and Elicitation of neonatal reflexes |
| 2   | Systemic examination of child- CNS                                     | 12  | Demonstration of Common procedures related to Paediatrics              |
| 3   | Systemic examination of child- CNS                                     | 13  | Demonstration of Common procedures related to Paediatrics              |
| 4   | Systemic examination of child- RS                                      | 14  | Common Drugs used in Paediatrics                                       |
| 5   | Systemic examination of child-Per Abdomen                              | 15  | Common Drugs used in Paediatrics                                       |
| 6   | Systemic examination of child- CVS                                     | 16  | Common Instruments used in Paediatrics                                 |
| 7   | Systemic examination of child- CVS                                     | 17  | X-Ray film reading related to Paediatrics.                             |
| 8   | Short case discussion  | 18  | Nutrition  |
| 9   | Neonatal case taking, examination and Elicitation of neonatal reflexes | 19  | Nutrition  |
| 10  | Neonatal case taking, examination and Elicitation of neonatal reflexes | 20  | Posting ending exam  |

| Competency Nos. | Topics, Subtopics and Lectures |
|-----------------|--------------------------------|
|-----------------|--------------------------------|

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## **Annexure- 4.**

### **Exam Pattern - Paediatrics**

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#### **Theory Paper (100 marks)**

- Section A-MCQ-:
- Section B-
- Section C-

#### **Practical exam (100 marks)**

- Long case-
- Short case/ Newborn-
- Table viva- (Drugs, Instruments, Nutrition, Vaccines and X-Rays-
- OSCE-

#### **Internal Assessment:**

- 50% combined in theory and practical (not less than 40% in each) for eligibility for appearing for University

#### **University Examination**

- Mandatory 50% marks separately in theory and practical (practical = practical/ clinical+ viva)
-



**Annexure- 5**  
**Distribution of journal marks**  
**Total- 10 marks**

| Parameter       | Total                                    | Marks | Phase                       |
|-----------------|--|-------|-----------------------------|
| Long cases      | -  | -     | Phase: II (Second year)     |
|                 | 6 (CNS-2, RS-1, PA-1, CVS-2)             | 1     | Phase: III-1 (Third Minor)  |
|                 | 66 (CNS-2, RS-1, PA-1, CVS-2)            | 1     | Phase: III-II (Third Major) |
| Short cases     | 3  | 1/2   | Phase: II (Second year)     |
|                 | 3  | 1/2   | Phase: III-1 (Third Minor)  |
|                 | 3  | 1/2   | Phase: III-II (Third Major) |
| Newborns        | 3  | 1/2   | Phase: II (Second year)     |
|                 | 3  | 1/2   | Phase: III-1 (Third Minor)  |
|                 | 3  | 1/2   | Phase: III-II (Third Major) |
| Emergencies     | 5  | 1     | Phase: III-1 (Third Minor)  |
| Procedures      | 5  | 1     | Phase: III-II (Third Major) |
| Vaccines        | All vaccines as per Government of India. | 1     | Phase: III-I                |
| Drugs           | 10                                       | 1     | Phase: III-II               |
| Instruments     | 10                                       | 1/2   | Phase: III-II               |
| Nutrition       | 10                                       | 1/2   | Phase: III-II               |
| Total- 10 marks |  |       |                             |



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