CORE 21123- Paediatrics

Status	Core
No of Hours	45 hours
No of Credits	3
Learning Outcomes	 Outline the main ways in which children's health and illness is different from adults and say how health professionals monitor growth, nutrition and development in Sri Lanka. List the most common childhood illnesses and be aware of their causes and consequences. Demonstrate understanding of elementary aspects of genetics and show awareness of the most common genetic conditions which cause disabilities. List the principle types of child abuse and outline a general procedure to follow if abuse is suspected. Describe briefly the possible effects of poor health or nutrition on children's health and development and show awareness of the link between poverty and nutrition. Show awareness of the link between poverty and nutrition. Demonstrate understanding of the structure and function of the central nervous system, especially as it relates to cognitive and language functioning and motor skills. Outline types and major causes of neurodevelopmental disorders and show understanding of the professionals how the multi- disciplinary professionals will collaborate from different sectors to provide services for these children and their families.
Methods of Teaching	 Discuss the incidence and prevalence of childhood disabilities Lectures(30 hours) and clinical lecture demonstrations
and Learning	
Module content	Unit 1: Growth
	 Nutrition screening and monitoring in Sri Lanka
	 Growth measurements and charts
	 Impact of nutrition and poverty on developmental and health
	 Breast feeding and complementary feeding
	 Unit 2: Nutritional disorders in children Protein energy malnutrition Water soluble vitamins Trace elements Biochemical basis of nutrition
	 Unit 3: Early identification of perinatal pediatric disorders leading to childhood disabilities Prematurity Birth asphyxia Other neonatal/ infant high risk conditions
	Unit 4: Child development
	 Normal development 0-5 years
	 Common neurodevelopmental disorders

Early detection and intervention

Unit 5: Childhood disabilities

- Congenital disorders
- Neuro developmental disorders
- Developmental delay
- Early detection and intervention of developmental disorders
- Common syndromes including Down Syndrome
- Cerebral palsy
- Epilepsy

Unit 6: Medical management of paediatric conditions

 Common pharmacological agents used in children with disabilities including medications for conditions such as Epilepsy, ADHD, Cerebral palsy and the effect on cognition, learning and development of communication

Unit 7: Infections of the Central Nervous system

- The microbiological basis for infections and common infections: Meningitis and encephalitis
- Congenital infections
- Vaccine prenvetable conditions and immunization schedule

Unit 8 : Child abuse and neglect

- Types of child abuse
- Policies for child protection in Sri Lanka
- Implementation of child protection policies in Sri Lanka

Unit 9: Child and adolescent mental health

- Autism
- Learning disorders
- Attention deficit hyperactive disorder

Unit 10: Genetics-Principles of genetics

- a. Principals of genetics
- Genes
- Human chromosome
- Cytogenetics
- Mitosis and Meiosis
- Numerical aberrations
- Sex chromosome anomalies
- Symbols used in pedigree construction
- Traits
- Environment and genetic interactions influencing fetus

b. Genetic component of impairment

- Cognitive impairment
- Pervasive developmental disorder
- Dyslexia
- Specific reading disabilities
- Stuttering

Assessment	 Role of the paediatrician in multidisciplinary management of paediatric conditions. Role of the SLT and Audiologist in multidisciplinary management of paediatric conditions. Other professionals involved Referral process
	 Environment syndromes Unit 11: Multi-disciplinary team management of paediatric conditions Multi-disciplinary team management Role of the paediatrician in multidisciplinary management of
	 Sporadic syndromes Genetic counseling Environment syndromes
	Polygenic – multifactorial syndromes
	 d. Syndromes Chromosomal syndromes Single gene syndromes
	 c. Genetic component of hearing impairment Use of gene libraries in the study of the molecular genetics of auditory system Epidemiology of genetic hearing impairment Genetic hearing loss