

PC01**Introduction of Neurometabolic Diseases in Infants and Children**

- 08:30-09:00 Overview of Neurometabolic Diseases in Infants
Ingrid Tein (The Hospital for Sick Children, Canada)
- 09:00-09:25 Mitochondrial Diseases: Infantile Onset Epilepsy from Genetic Testing to Precision Management
Ching-Shiang Chi (Tungs' Taichung MetroHarbor Hospital, Taiwan)
- 09:25-09:50 Investigations in Pediatric Neurometabolic Diseases
Yann-Jang Chen (Taipei Veterans General Hospital, Taiwan)
- 09:50-10:10 Neurophysiological Monitoring on Neurometabolic Disorders
Douglas R. Nordli (University of Chicago, USA)

PC02**Mitochondrial Encephalopathy**

- 10:40-11:10 Approach to Mitochondrial Disorders in Children
Haluk Topaloglu (Hacettepe University School of Medicine, Turkey)
- 11:10-11:35 Mitochondrial Encephalopathy: Clinical and Genetic Features
Kei Murayama (Chiba Children's Hospital, Japan)
- 11:35-12:00 Molecular Basis of Neurometabolic and Neurogenetic Diseases
Henrike O. Heyne (University of Helsinki, Finland)

PC03**Amino Acid Metabolic Diseases**

- 13:30-14:00 Overview of Amino Acid Metabolic Diseases: Clinical Presentations and Genetic Roles
Cheuk Wing Fung (Hong Kong Children's Hospital, Hong Kong)
- 14:00-14:30 MSUD and Related Disorders: Different Presentations in Infants and Children
Sylvia Estrada (Philippine General Hospital, Phillipines)
- 14:30-15:00 Sulfite Oxidase Deficiency and Related Disorders: Neuroimaging Findings and Genetic Roles
Syuan-Yu Hong (China Medical University, Taiwan)

PC04**Miscellaneous Etiologies and Treatment**

- 15:40-16:05 Artificial Intelligence Application in EEG Analysis: A New Strategy for the Neurophysiological Management of Neurometabolic Disorders
Noboru Yoshida (Juntendo University Nerima Hospital, Japan)
- 16:05-16:35 Peroxisomal Disorders in Infants and Children
Hsi Chang (Taipei Medical University Hospital, Taiwan)
- 16:35-17:00 Metabolic Imbalances in Fatty Acid Oxidation Disorders. Implications for Pathophysiology and Treatment
Jerry Vockley (University of Pittsburgh, USA)
- 17:00-17:30 Lysosomal Diseases in Infancy and Children: Diagnostic Approach
Jonathan Mink (University of Rochester Medical Center, USA)

BS01**Update in Neurophysiology**

- 07:15-07:45 EEG Monitoring Approaches in Critically Ill Infants and Children
Nicholas S. Abend (University of Pennsylvania, USA)
- 07:45-08:15 Update Applications of Neurophysiological Monitoring in Neonates
Ronit Pressler (Great Ormond Street Hospital, UK)

K01**Keynote Speech I**

- 08:40-09:20 Recent Advance in Neurometabolic Diseases: The Genetic Role in Modern Era
Ingrid Tein (The Hospital for Sick Children, Canada)

K02**Keynote Speech II**

- 09:20-10:00 Recent Advance in the Treatment of Neurometabolic Diseases
Phillip Pearl (Boston Children's Hospital, USA)

S01**Neurometabolic Diseases and Epilepsy (I)**

- 10:30-10:50 Ion-Channel Disorders V.S. Neurometabolic Disorders in Newborns
Inn-Chi Lee (Chung Shan Medical University Hospital, Taiwan)
- 10:50-11:10 Novel Treatment of Neurometabolic Disease Presenting with Epileptic Encephalopathy
Kazuhiro Muramatsu (Jichi Medical University, Japan)
- 11:10-11:30 Diet Therapy for Infants with Neurometabolic Disorders and Genetic Epilepsy
Heung Dong Kim (Yonsei University College of Medicine, Korea)
- 11:30-12:00 Precision Medicine in Infantile Seizures
Henrike O. Heyne (University of Helsinki, Finland)

S02**Investigations in Neurometabolic Disease**

- 13:30-14:00 Clinical Utility of Rapid Whole Genome Sequencing in Infants with Neurometabolic Diseases
Pui-Yan Kwok (Academia Sinica, Taiwan)
- 14:00-14:30 Pyridoxine-Dependent Epilepsy (PDE-ALDH7A1): Implications for Newborn Screening
Laura Tseng (BC Children's Hospital, Netherlands)
- 14:30-15:00 Neuroimaging Features in Infants with Neurometabolic Diseases
Kshitij Mankad (Great Ormond Street Hospital / University College London Hospital, UK)

S03**Infantile Epilepsy in Mitochondrial Disorders**

- 16:00-16:30 Mitochondrial Transplantation in MELAS Disease
Chin-San Liu (Changhua Christian Hospital, Taiwan)
- 16:30-16:50 Neurophysiologic Monitoring in Infants with Mitochondrial Diseases
Ronit Pressler (Great Ormond Street Hospital, UK)
- 16:50-17:10 Infantile Onset Epilepsy in Mitochondrial Disorder: Clinical and Genetic Insights
Jong Hee Chae (Seoul National University College of Medicine, Korea)
- 17:10-17:30 Paradigm Changes in the Genetics of the Infantile Epilepsies – Understanding the Exome and Beyond
Ingo Helbig (Children's Hospital of Philadelphia, USA)

BS02**Progressive Myoclonic Epilepsy in Infants and Children**

- 07:15-07:45 Overview of Progressive Myoclonic Epilepsy: Genetic Roles
Nicola Specchio (Bambino Gesù Children's Hospital, Italy)
- 07:45-08:15 Update Treatment of PME
Jorge Vidaurre (Nationwide Children's Hospital, USA)

K03**Keynote Speech III**

- 08:30-09:10 Metabolic Epilepsy in Infancy: The Role of Genes
Pratibha Singhi (Medanta, India)

S04**Neurometabolic Diseases and Epilepsy (II)**

- 10:30-10:55 Clinical, Genetic Role, and Treatment Strategies toward SSADHD
Phillip Pearl (Boston Children's Hospital, USA)
- 10:55-11:20 Genetic Investigation in Lipid Metabolism with Infantile Epilepsy
Ting-Rong Hsu (Taipei Veterans General Hospital, Taiwan)
- 11:20-11:45 The Role of Autophagy in the Pathomechanism and Treatment of Leukodystrophy
Dar-Shong Lin (Taipei Veterans General Hospital, Taiwan)
- 11:45-12:10 Update Treatment for Leukodystrophy and Its Associated Epilepsy in Infants
Wang-Tso Lee (National Taiwan University Hospital, Taiwan)

S05**Neurometabolic Diseases and Epilepsy (III)**

- 13:30-13:55 Congenital Disorders of Glycosylation and Infantile Epilepsy
Hsiu-Fen Lee (Taichung Veterans General Hospital, Taiwan)
- 13:55-14:20 Infantile Epilepsy and Carnitine Inborn Errors of Metabolism: The Role of Genes
Shinichi Hirose (Fukuoka University, Japan)
- 14:20-14:40 Organic Acid and Infantile Epilepsy: The Role of Genes
Yi Wang (Fudan University Children Hospital, China)
- 14:40-15:00 Diagnosis of Neurometabolic Disorders Involving Basal Ganglia based on Neuroimages
Shekeeb Mohammad (The Children's Hospital at Westmead, Australia)

S06**Treatable Neurometabolic Diseases and Epilepsy**

- 15:30-15:55 Implication of Glucose Transporter 1 Deficiency in Neurologic Disease
Shin Nabatame (Osaka University, Japan)
- 15:55-16:20 Pyridoxine-Responsive and Dependent Epilepsy
Huei-Shyong Wang (Linkou Chang Gung Memorial Hospital, Taiwan)
- 16:20-16:45 Biotinidase Deficiency and Infantile Seizures
I-Ching Chou (China Medical University Hospital, Taiwan)
- 16:45-17:10 Novel Treatment of Mitochondrial Disorders
Tsu-Kung Lin (Kaohsiung Chang-Gung Memorial Hospital, Taiwan)
- 17:10-17:35 Creatine Transporter Deficiency and Epilepsy
Ming-Tao Yang (Far Eastern Memorial Hospital, Taiwan)