Invited Speakers（Tentative）

*As of January 19. Information will be updated as needed.

Edward C. Cooper
Neurology, Neuroscience, and Molecular and Human Genetics, Baylor College of Medicine, USA
"The title will be announced at a later date."

Yuwu Jiang
Children's Medical Center, Pediatric Epilepsy Center, Peking University First Hospital, China
"Identification of Novel Genes in Rare Genetic Epilepsies"

Katty (Jing-Qiong) Kang
Department of Neurology & Pharmacology, School of Medicine, Vanderbilt University, Vanderbilt University Medical Center, USA
"GABAergic Pathway Dysfunction in Epilepsy: From Mechanisms to Therapeutic Rescue"

Hoon-Chul Kang
Yonsei University, Korea
"Molecular therapy for channelopathy"

Wang-Tso Lee
National Taiwan University Hospital Taipei, Taiwan
"The title will be announced at a later date."

Naomichi Matsumoto
Department of Human Genetics, Yokohama City University Graduate School of Medicine, Japan
"A Comprehensive Genetic Analysis of Rare Neurological Disorders Including Epilepsy"

Jeffrey L. Noebels
Cullen Trust for Health Care Endowed Chair, Professor of Neurology, Neuroscience, and Molecular and Human Genetics, Vice Chair for Research, USA
"Channelopathy during Early Brain Development"

Ingrid Scheffer
The University of Melbourne, Australia
"The genetics of the epilepsies: an ever expanding landscape"

Pratibha Singh
The International Child Neurology Association, Head Department of Pediatric Neurology Amrita Hospital Faridabad, India
"The Molecular Pathomechanisms and Therapeutic Strategies of Rare and Intractable Epilepsies"

Nicola Specchio

Neurology, Epilepsy and Movement Disorders Unit, Director of Research Unit on Neurological and Neurosurgical Diseases, Bambino Gesù Children's Hospital, IRCCS, Italy

"The title will be announced at a later date."

Kazuhiro Yamakawa

Department of Neurodevelopmental Disorder Genetics, Institute of Brain Science,
Nagoya City University Graduate School of Medical Sciences, Japan

"Molecular pathomechanism of epileptic sodium channelopathy"