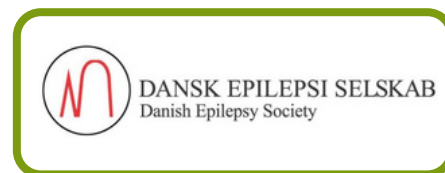


FILADELFIA

6th Dianalund International Conference on Epilepsy

Overlapping clinical phenotypes
in monogenic epilepsies –
common molecular pathways?



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MAY 02 - 03, 2024

Køge, Denmark

Pre-Conference Meeting Agenda

Wednesday 1 May

Chairs: Sebastian Ortiz, Francesca Furia & Sopio Gverdtsiteli

16:30 **Welcome**

16:30 - 16:40 **Helene Meinild** (Jazz Pharmaceuticals): Treatment complexity in patients with epilepsy and severe intellectual disability requires a comprehensive approach

16:40 - 16:50 **Kate Baker** (Cambridge, UK): The neurodevelopmental spectrum of synaptic vesicle cycling disorders

16:50 - 17:00 **Nathan Routledge** (London, UK): Lessons learned from a genotype-first analysis of voltage-gated ion channels

17:00 - 17:10 **Charlotte Andersen** (Angelini Pharma): Cenobamate - recent updates

17:10 - 17:20 **Antonietta Coppola** (Naples, IT): Clinical features and genotype-phenotype correlations in epilepsy patients with de-novo DYNC1H1 variants

17:20 - 17:30 **Kristine Sloth Vilhelmsen** (Copenhagen, DK): Development of the first GABAAR δ subunit NAM

17:30 - 17:40 **Troels Wesenberg Kjær** (UNEEG medical): Ultra long-term monitoring in epilepsy with a subcutaneous EEG device

Break

17:40 - 17:50 Coffee break



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Wednesday 1 May (continued)

- 17:50 - 18:00 Roberto Previtali** (Milan, IT): The use of ketogenic diet as target therapy in a cohort of Italian patients affected by GLUT1 deficiency syndrome
- 18:00 - 18:10 Susan Lin** (Sydney, AU): If a variant is predicted to be “benign/tolerated” by silico tools and seen in gnomAD. Is it benign?
- 18:10 - 18:20 Konrad Werhahn** (UCB): Precision-based treatment within epilepsy
- 18:20 - 18:30 Allan Bayat** (Dianalund, DK): Natural history of adults with KBG syndrome: a physicians reported experience
- 18:30 - 18:40 Leonardo Affronte** (Dianalund, DK): ESES/DEE-SWAS in p.(Pro405Leu) KCNA2 pathological variant
- 18:40 - 18:50 Baharak Saeedi** (Orion Pharma): Long-term treatment with ganaxolone for seizures associated with cyclin-dependent kinase-like 5 deficiency disorder (CDD): 2-year open-label extension follow-up
- 18:50 - 19:00 Sopio Gverdtsiteli** (Dianalund, DK): Neonatal Developmental and Epileptic Encephalopathy with movement disorders and arthrogyriposis (NDEEMA) – common phenotype across brain-expressed sodium channelopathies



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