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410 Kinderklinik

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Psychosomatische Tagesklinik, Sportmedizin,
ukfit, Gästehaus

500 Medizinische Klinik

510 Blutspendezentrale

520 Konferenzzentrum
Casino, Cafeteria

530 Nuklearmedizin

600 Hals-Nasen-Ohrenklinik

610 Medizinische Mikrobiologie, Medizinische Virologie

620 Augenklinik

University Clinic Tübingen
Kliniken Berg CRONA
Lecture Hall, 4th floor, room 200
Hoppe-Seyler-Str. 3
72076 Tübingen

Conference link for online participants/speakers:

<https://med-uni-tuebingen-de.zoom-x.de/j/69452016734?pwd=FpVvfMyx2KqEa9374SZ7CS9cHPW58.1>

Meeting-ID: 694 5201 6734

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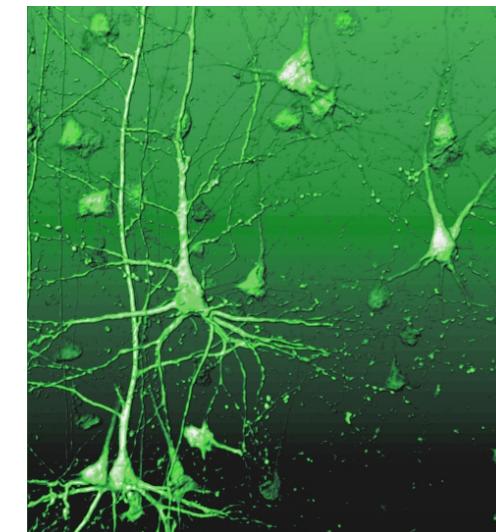
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3rd Channelopathy Meeting Tübingen

Genetic epilepsies and other
neuronal ion channel disorders:
Mechanisms and therapeutic
perspectives



Program

Wednesday, 24th September 2025

8:30-8:40 am	Introduction: Holger Lerche (Tübingen)
8:40-10:15 am	Session 1: Ion channel mechanisms I Chair: Dirk Isbrandt (Cologne) & Snezana Maljevic (Melbourne) Maurizio Taglialatela (Naples): KCNQ modulators: from an atomistic view of channel gating Marisol Sampedro Castaneda (London): CDKL5 and its interaction with Ca _v 2.3 Massimo Mantegazza (Valbonne-Sophia Antipolis): Mechanisms of SCN2A variants associated with Autism Spectrum Disorder Ahmed Eltokhi (Columbus): Gating pore current in Na _v 1.2 mutations: Implications for autism and epilepsy Daniil Kirianov (Cologne): Unravelling the seizure initiation and progression through the neonatal Scn2a (p.A263V) hippocampus
10:15-10:40 am	Coffee Break
10:40 am-12:00 pm	Session 2: Immunological and other epileptogenic mechanisms Chair: Albert Becker (Bonn) & Christian Geis (Jena) Harald Prüß (Berlin): Antibody-mediated channelopathies - new disease concepts Julika Pitsch (Bonn): Microstructural correlates of neuro-immune dysregulation in autoimmune encephalitis Michael Wenzel (Bonn): Hippocampal spreading depolarization as a key epilepsy disease factor Christian Geis (Jena): Effects of NMDAR autoimmunity on receptor function and hippocampal circuits
12:00-1:05 pm	Session 3: Progress in genetic mechanisms and prediction algorithms Chair: Rikke Møller (Dianalund) & Yvonne Weber (Aachen) Alex Hoischen/Holm Graeßner (Nijmegen/Tübingen): Rare diseases – boosting diagnostic yield by data re-analysis and long-read genome sequencing Josua Kegele (Tübingen): Short- and long-read genome sequencing in early-onset DEE: results and insights Henrike Heyne (Cambridge): Predicting functional effects of genetic variants in ion channels with methods of deep learning
1:05-2:15 pm	Lunch Break
2:15-3:25 pm	Session 4: Gene therapy Chair: Marius Ueffing (Tübingen) & Gaia Colasante (Milan) Keynote lecture Steven Gray (Dallas): Gene therapy for neurological disorders: the example of SLC6A1 deficiency Dirk Grimm (Heidelberg): Parvoviral vector-mediated gene therapy in the CNS (and beyond): A no-brainer? Elvir Becirovic (Zürich): AAV vector-mediated delivery of large genes for retinal gene therapy and beyond
3:25-4:30 pm	Poster Session (with coffee)
4:30-6:30 pm	Session 5: Clinical trials and molecular therapeutic board (case reports) Chair: Holger Lerche (Tübingen) & Steve Petrou (Praxis Precision Medicines, Boston) Victoria Ruschil (Tübingen): Familial episodic pain syndrome in a family with an unknown SCN11A-variant

4:40 pm	Walid Fazeli (Bonn)/Matias Wagner (Munich)/Steve Petrou (Praxis Precision Medicines, Boston): Towards disease-modification in SCN2A DEE: insights into antisense-oligonucleotide
5:05 pm	Oleg Vinogradov/Elena Kuster (Tübingen)/Henning Steinhagen (Lario, Edinburgh): Towards a targeted therapy for CACNA1E-associated DEE: Functional insights and natural history
5:30 pm	Lidia Carotenuto (Antwerp): The fast-dissociating D2 antagonist antipsychotic JNJ-378 22681 is a neuronal K _v 7-channel opener: potential repurposing for epilepsy treatment
5:40 pm	Rikke Steensbjerre Møller (Dianalund): From disease-causing variants to targeted therapy in GABA-A receptor related epilepsies
5:50 pm	Francesco Miceli (Naples): <i>In vitro</i> pharmacological characterization of a potent blocker of the epilepsy-associated K _v 7.2 channel
6:00 pm, online	Kris Kahlig (Praxis Precision Medicines, Boston): Relutrigine and Vormatrigine: Functional-State Sodium Channel Modulators Advancing Therapies for Rare and
7:30 pm	Dinner Restaurant „Liquid“

Thursday, 25th September 2025

8:30-9:35 am	Session 6: Advanced techniques and model systems Chair: Ulrike Hedrich (Tübingen) & Michael Wenzel (Bonn) Simon Musall/Viviana Rincon Montes (Jülich): High-Density Neurophysiology: From Rigid Arrays to Flexible Interfaces Karen van Loo (Aachen): Gene modulation in epilepsy: using organotypic brain slice cultures to analyze functional and molecular changes Yvonne Weber (Aachen): Drosophila as a model system for genetic epileptic and developmental epilepsies
9:35 am-1:00 pm	Session 7: Ion channel mechanisms II Chair: Thomas Wuttke (Tübingen) & Gabriele Lignani (London) Evangelos Kiskinis (Chicago): Advanced iPSC models on KCNQ2 and KCNH1 associated epilepsy Franck Kalume (Seattle): Interneuron-specific dual-AAV SCN1A gene replacement for Dravet syndrome: A preclinical update Gaia Colasante (Milan): Enhancing Na _v 1.1 translation by gene editing to treat Dravet Syndrome
10:35-11:00 am	Coffee Break
11:00 am	Gabriele Lignani (London): Prenatal sodium channel dysfunction in Dravet syndrome alters cortical development
11:20 am	Snezana Maljevic (Melbourne): Outside the Spotlight: The Unexpected Roles of Ion Channels in Early Brain Development
11:40 am	JP Gilbert (Xenon, Burnaby): Selective potentiation of Na _v 1.1 channels by XPC-A in Dravet mice suppresses spontaneous seizures, prevents SUDEP, and increases long term potentiation
12:00 pm	Alison Obergrussberger (Nanion, Munich): Exploring the role of lysosomal ion channels and transporters in neurodegenerative diseases
12:20 pm	Daniela Miely (Tübingen): Molecular and cellular basis of episodic ataxia 6
12:40 pm	Gaëtan Lesca (Lyon): Involvement of CACNA2D2 in developmental and epileptic encephalopathy through disruption of calcium channel functionality and synaptic function
1:00-1:30 pm	Farewell Lunch with free discussions