

## 3<sup>rd</sup> Channelopathy Meeting Tübingen

Genetic epilepsies and other neuronal ion channel disorders: Mechanisms and therapeutic perspectives 23<sup>rd</sup>-25<sup>th</sup> September 2025, Tübingen, Germany

## **Preliminary Program**

Tuesday, 23 September 2025		
7:30 pm	Get-Together Schloss Hohentübingen	
	Wednesday, 24 September 2025	
8:30 am	Welcome and introduction: Holger Lerche (Tübingen)	
8:40-10:00 am	Session 1: Ion channel mechanisms I	
	Chair: Dirk Isbrandt & Snezana Maljevic	
8:40	Maurizio Taglialatela (Neapel): KCNQ modulators: from an atomistic view of channel	
	gating to personalized therapies	
9:00	Marisol Sampedro Castaneda (London): CDKL5 and its interaction with Cav2.3	
9:20	Massimo Mantegazza (Valbonne-Sophia Antipolis): Mechanisms of SCN2A variants	
	associated with Autism Spectrum Disorder	
9:40	Ahmed Eltokhi (Columbus): Gating Pore Current in Na $_{\rm V}$ 1.2 Mutations: Implications for	
	Autism and Epilepsy	
9:55	Daniil Kirianov (Cologne): Unravelling the seizure initiation and progression through the	
	neonatal <i>Scn2a</i> (p.A263V) hippocampus	
10:15-10:40 am	Coffee Break	
10:40-12:00 am	Session 2: Immunological and other epileptogenic mechanisms	
	Chair: Albert Becker & Christian Geis	
10:40	Harald Prüß (Berlin): Antibody-mediated channelopathies - new disease concepts	
11:00	Julika Pitsch (Bonn): Microstructural correlates of neuro-immune dysregulation in	
	autoimmune encephalitis	
11:20	Michael Wenzel (Bonn): Hippocampal spreading depolarization as a key epilepsy disease	
11.40	factor	
11:40	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	
12.00-1.05 pm	Chair: Rikke Møller & Yvonne Weber	
12:00	Alex Hoischen/Holm Graeßner (Nijmegen/Tübingen): Rare diseases – boosting diagnostic	
	yield by data re-analysis and long-read genome sequencing	
12:30	Josua Kegele (Tübingen): Results in the novel-omics sequencing project	
12:45	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 3: Gene therapy	
	Chair: Marius Ueffing & Gaia Colasante	
2:15, virtuell	Keynote lecture	
	Steven Gray (Dallas): Gene therapy for neurological disorders: the example of SLC6A1	
	deficiency	
2:45	Dirk Grimm (Heidelberg): Parvoviral vector-mediated gene therapy in the CNS (and	
	beyond): A no-brainer?	
3:05	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 4: Clinical trials and Molecular therapeutic board (case reports)	
	Chair: Holger Lerche & Steve Petrou	

4:30	Walid Fazeli (Bonn)/Matias Wagner (Munich)/Steve Petrou (PraxisMedicine, Boston): tbd
4:55	Oleg Vinogradov/Elena Kuster (Tübingen)/Henning Steinhagen (Edinburgh): tbd
5:20	Lidia Carotenuto (Antwerp): The fast-dissociating D2 antagonist antipsychotic JNJ-
	37822681 is a neuronal K <sub>v</sub> 7 channel opener: potential repurposing for epilepsy treatment
5:30	Victoria Ruschil (Tübingen): Familial episodic pain syndrome in a family with an unknown
	SCN11A-variant
5:40	Rikke Steensbjerre Møller (Dianalund): From disease-causing variants to targeted
	therapy in GABA-A receptor related epilepsies
5:50	Francesco Miceli (Naples): In vitro pharmacological characterization of a potent blocker
	of the epilepsy-associated K <sub>v</sub> 7.2 channel
6:00-6:30	Further cases
8:00 pm	Dinner Restaurant "Liquid"

Thursday, 25 September 2025		
8:30-9:35 am	Session 6: Advanced techniques and model systems	
	Chair: Ulrike Hedrich & Michael Wenzel	
8:30	Simon Musall/Viviana Rincon Montes (Jülich): High-Density Neurophysiology: From Rigid	
	Arrays to Flexible Interfaces	
8:55	Karen van Loo (Aachen): Gene modulation in epilepsy: using organotypic brain slice	
	cultures to analyze functional and molecular changes	
9:15	Yvonne Weber (Aachen): Drosophila as a model system for genetic epileptic and	
	developmental epilepsies	
9:35-1:00 pm	Session 7: Ion channel mechanisms II	
	Chair: Thomas Wuttke & Gabriele Lignani	
9:35	Franck Kalume (Seattle): Interneuron-specific dual-AAV SCN1A gene replacement for	
	Dravet syndrome: A preclinical update	
9:55	Gaia Colasante (Milan): Enhancing Na $_{\rm V}$ 1.1 Translation by Gene Editing to Treat Dravet	
	Syndrome	
10:15	Gabriele Lignani (London): Prenatal sodium channel dysfunction in Dravet syndrome	
	alters cortical development	
10:35-11:00 am	Coffee Break	
11:00	Snezana Maljevic (Melbourne): Outside the Spotlight: The Unexpected Roles of Ion	
	Channels in Early Brain Development	
11:20	GP Gilbert (Xenon, Burnaby): Selective potentiation of Na $_{ m V}$ 1.1 channels by XPC-A in	
	Dravet mice suppresses spontaneous seizures, prevents SUDEP, and increases long	
	term potentiation	
11:40	Christoph Fahlke (Jülich)/Daniela Miely (Tübingen): Molecular and cellular basis of	
	episodic ataxia 6	
12:00	Alison Obergrussberger (Nanion, Munich): Exploring the role of lysosomal ion channels	
	and transporters in neurodegenerative diseases	
12:20	Myriad Essid (Lyon): Involvement of CACNA2D2 in developmental and epileptic	
	encephalopathy through disruption of calcium channel functionality and synaptic	
	function	
12:40	Evangelos Kiskinis (Chicago): Advanced iPSC models on KCNQ2 and KCNH1 associated	
	epilepsy	
1:00-1:30 pm	Farewell Lunch with free discussions	

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