



**Universität
Zürich** ^{UZH}



10th Rare Disease Summer School

Kartause Ittingen, Warth
July 9th to 12th 2024

PROGRAM

The program is subject to change without notice.

Please refer to the Summer School website for updates.





TUESDAY, JULY 9th 2024

13:00 – 14:00 ARRIVAL OF THE PARTICIPANTS AND REGISTRATION

14:00 – 14:30 **Welcome and introduction round**, Prof. Matthias Baumgartner

14:30 – 15:15 Prof. Nine Knoers, Department of Genetics, University Medical Center
Groningen, the Netherlands

Latest developments in exome and genome sequencing: impact for rare diseases

15:15 – 15:45 COFFEE BREAK

15:45 – 16:30 Dr. Jasmin Barman-Aksözen, Scientific advisor, Swiss society for porphyria, Switzerland

Let the sunshine in! Development, approval and benefit assessment of afamelanotide, the first drug for treating erythropoietic protoporphyria.

16:30 – 17:30 **Oral presentations** by selected participants

17:30 – 18:30 Check-in time

18:30 – 20:15 DINNER

20:15 – **Poster viewing**



WEDNESDAY, JULY 10th 2024

07:45 – 08:45 BREAKFAST

08:45 – 09:30 Prof. Marshall Summar, Professor of Paediatrics George Washington University / CEO: Uncommon Cures, USA

Rare diseases today

09:30 – 10:15 Prof. Stephan Neuhaus, Department of Molecular Science, University of Zurich, Switzerland

Studying rare diseases in the (not so rare) Zebrafish model organism

10:15 – 10:45 COFFEE BREAK

10:45 – 12:15 Prof. Nine Knoers, Department of Genetics, University Medical Center Groningen, the Netherlands

Workshop: Next generation sequencing

12:15 – 13:45 LUNCH

13:45 – 14:30 Kelly Ormond, ABGC and EBMG Registered Genetic Counselor Lecturer and Research Scientist, ETH-Zurich, Switzerland

Ethics of Genetic Testing and Treatments of Genetic Conditions

14:30 – 15:15 Dr. Fabienne Hartmann-Fritsch, University of Zurich, CUTISS AG, Switzerland

Regulatory framework for ATMPs on the basis of a case study

15:15 – 15:45 COFFEE BREAK

15:45 – 16:30 "Molecular underpinnings of rare, extreme memory performance"

Keynote Lecture: Andreas Papassotiropoulos

16:30 – 17:30 Networking/Free time

17:30 – 18:30 Visit to the Kartause Museum

18:30 – 20:00 DINNER

20:00 – **Poster viewing**



THURSDAY, JULY 11th 2024

07:45 – 08:45 BREAKFAST

08:45 – 10:30 Dr. Jürg Streuli, Institute of Biomedical Ethics, University of Zurich,
Switzerland

Interactive workshop: Ethical considerations (case study)

10:30 – 11:00 COFFEE BREAK

11:00 – 11:45 Prof. Isaac Canals, Department of Experimental Medical Science,
Lund University, Sweden

**iPSC and genome editing as tools to generate human in vitro
models of neuronopathic lysosomal storage disorders**

11:45 – 12:45 **Oral presentations** by selected participants

12:45 – 14:15 LUNCH

14:15 – 15:00 Prof. Olivier Devuyst, Institute of Physiology, University of Zurich,
Switzerland

**Genetic architecture driving therapeutic targets in kidney
diseases**

15:00 – 15:45 Prof. Elvir Becirovic, Ophthalmology, University Hospital Zurich,
Switzerland

Treatment of inherited retinal diseases: chances and obstacles

15:45– 16:15 COFFEE BREAK

16:15 – 17:45 Prof. Marshall Summar, Children's National Medical Center, George
Washington University, USA

Workshop: Designing rare disease clinical trials

17:45 – 18:30 Free time

18:30 – 21:00 GRILL



FRIDAY, JULY 12th 2023

07:45 – 08:45 BREAKFAST

08:45 – 09:30 **Take-home messages** by participants

09:30 – 10:15 Prof. Günter Schwarz, University of Cologne, Germany
Regulatory approval of a first treatment for the ultra-rare molybdenum cofactor deficiency

10:15 – 10:45 COFFEE BREAK

10:45 – 11:30 Prof. Britta George, University Hospital Zurich, Switzerland

11:30 – 12:15 Hansruedi Silberschmidt, patient with rare disease
Coughing prohibited – living with a rare disease

12:15 – 12:30 Award of poster prizes and concluding remarks

12:30 **END OF THE SUMMER SCHOOL AND DEPARTURE OF THE PARTICIPANTS**

SPEAKER (*in alphabetical order*)

DR. JASMIN BARMAN-AKSÖZEN



Jasmin Barman-Aksözen is a patient with the ultra-rare inborn error of metabolism erythropoietic protoporphyria (EPP). She studied molecular biology and biochemistry at the University of Heidelberg and obtained her PhD and Venia Legendi from the University of Zurich. Her research topics cover basic science and drug development in EPP and related diseases (porphyrias), as well as aspects concerning the regulatory assessment and benefit evaluation of orphan drugs. During her PhD, she was involved in the development of the first drug for treating EPP and later became a patient representative for the respective approval proceedings at the European Medicines Agency. To promote access to therapy for all patients with EPP and other porphyrias, she co-founded the International Porphyria Patient Network, a group of dedicated patient advocates with a professional background in science, medicine, and other relevant expertise. She is working at the Institute for Laboratory Medicine at the Municipal Hospital Zurich, which is the Swiss Reference Center for the Porphyrias, and is currently studying health economics and methods of health technology assessment at the University of Applied Sciences in Winterthur.

PROF. ELVIR BECIROVIC



Elvir Becirovic studied biology at the University of Cologne from 2001-2006. He received his PhD at the Ludwig-Maximilians-University of Munich in 2010. He then taught and conducted research there as a postdoctoral fellow and, since 2014, as a junior research group leader on the pathophysiology of retinal diseases. In 2016, he was enrolled as a postdoctoral fellow and completed the habilitation process in 2019. Since 2016, his research focus has been on gene therapy of retinal diseases. In June 2022, he was appointed Assistant Professor of Experimental and Translational Ophthalmology at the University of Zurich. In addition to his academic activities, he works as Chief Technology Advisor for ViGeneron GmbH and has several patents as lead inventor.

PROF. ISAAC CANALS



Isaac Canals studied biology at University of Barcelona and graduated in 2006. He received his PhD in Genetics in 2015 by the University of Barcelona on a work on Sanfilippo syndrome, a rare lysosomal storage disorder in which he generated novel iPSC-derived cellular models. During his postdoctoral training at Lund University, he specialized in neuroscience and develop a method to differentiate iPSCs towards astrocytes, technology that he combined with genome editing and methods to generate neurons for modeling neurodegenerative disorders. He established his own research group in 2021 at Lund

University. In 2023 he joined the University Children's Hospital of Zurich as an Assistant Professor. The aim of his research is to generate different iPSC-based 2D and 3D in vitro models to understand disease mechanisms of lysosomal storage disorders, the contribution of different brain cells to the pathology and what are the neurodevelopmental impairments in patients, as well as to develop new therapeutic approaches based on genome editing technologies.

PROF. OLIVIER DEVUYST



Olivier Devuyt, M.D., Ph.D., graduated from UCLouvain Medical School in Brussels (Belgium) and trained at the Technion Institute (Haifa, Israel) and the Johns Hopkins Medical School (Baltimore, USA). He is Full Professor at the University of Zurich, where he co-chairs the University priority program on rare diseases and has a joint clinical appointment at Saint-Luc Academic Hospital, UCLouvain. Devuyt and his group investigate the mechanisms of epithelial transport and the genetic architecture of kidney diseases, paving the way for novel therapeutic approaches. In parallel, they demonstrated fundamental mechanisms of osmosis and the crucial role of water channels (aquaporins) in dialysis. O. Devuyt has published more than 300 original studies, cited >48,000 times. He served as President and Board Member in the Belgian and Swiss societies of nephrology, coordinated several EU-funded research networks and established the Working Groups on Inherited Kidney Disorders of the European Renal Association and the Swiss Society for Nephrology.

DR. FABIENNE HARTMANN-FRITSCH



Fabienne Hartmann-Fritsch received her MSc in Microbiology and Immunology from the ETH Zurich in 2008 and her PhD in Biology from the University of Zurich in 2013. During her postdoctoral studies at the Tissue Biology Research Unit of the University Children's Hospital Zurich, Dr. Hartmann-Fritsch specialized on GMP-production of human skin grafts as well as on clinical trial management and regulatory affairs. In 2017, she co-founded the UZH spin-off company CUTISS AG, with the vision to provide patients who suffer from large and deep skin defects with the first personalized and automated skin tissue therapy that is safe, effective, and accessible for children and adults.

PROF. NINE KNOERS



Nine Knoers is the chair of the Department of Genetics at the University Medical Centre Groningen, The Netherlands. She received her M.D. (1986) and Ph.D. (cum laude, 1990) from the Catholic University Nijmegen, The Netherlands. She was trained and certified as Clinical Geneticist at the Radboud University Medical Centre Nijmegen. From 2011 until 2018, she was chair of the Department of Genetics at the University Medical Centre Utrecht, The Netherlands. Her major research effort focuses on the identification and further characterization of genes for inherited renal disorders and on their pathophysiology. The ultimate aim of her studies is to find clues for treatments for these disorders. Prof. Knoers is member of several European Consortia on genetic renal disorders and coordinated the Dutch Scientific Consortium on renal ciliopathies “Kouncil”. She has published over 280 scientific papers and in the last 10 years delivered >250 invited lectures.

PROF. STEPHAN NEUHAUSS



Stephan Neuhauss studied biology in Tübingen (D) and Eugene (Oregon, USA) doing his graduate work at Harvard Medical School (Massachusetts General Hospital). He then returned to Europe for a postdoctoral fellowship to the Max-Planck Institute for Developmental Biology (Tübingen) before accepting a senior lecture ship of the ETH Zurich at the Brain Research Institute. After stints as an assistant professor at the ETH, he became associate and later full professor at the Department of Molecular Life Sciences of the University of Zurich. He is also Vice-Dean for Research at the Faculty of Science and an editor of various open access journals.

KELLY ORMOND



Kelly Ormond is a genetic counselor (US ABGC certified and EBMG registered) and ELSI researcher. She received her MS in Genetic Counseling from Northwestern University (1994) and a post-graduate certificate in Clinical Medical Ethics from the MacLean Center at the University of Chicago (2001). She joined the Health Ethics and Policy Lab as a Senior Scientist in February 2021, and is an Adjunct Professor in the Department of Genetics at Stanford School of Medicine, Stanford University, California, USA. Prior to joining ETH, Kelly practiced as a clinical genetic counselor in the United States for >25 years, gaining experience in reproductive genetics, teratology, pediatric/medical genetics, adult neurogenetics, cancer genetics and primary care applications of genetic testing. Kelly's recent research focuses on ethical issues related to genetics and personalized medicine, including patient decision making, the role of uncertainty in



decision making, consent and disclosure of genetic test results, gene editing, and the interface between genetics and disability.

PROF. ANDREAS PAPASSOTIROPOULOS



Andreas Papassotiropoulos is board certified psychiatrist and psychotherapist and, since 2007, full Professor of Molecular Neuroscience at the University of Basel, Switzerland. His areas of expertise include the investigation of the molecular basis of human cognition and the development of improved therapies for neuropsychiatric disorders. His research combines targeted and genome-wide genetic, epigenetic and transcriptomic analyses with functional brain imaging in healthy and diseased human populations. He is recipient of numerous honors and awards, amongst others the Robert-Bing-Prize, The Weizmann Lecture, and the Cloetta Prize.

PROF. GÜNTER SCHWARZ



Günter Schwarz graduated his studies in Plant Biology and Biochemistry and habilitated *Venia legendi* in Biochemistry and Molecular Biology at the Technical University of Braunschweig (TU), Germany.

He is full Professor (W3) for Biochemistry at the University of Cologne, Germany. He and his team are interested to understand protein structure-function relations in different biological processes ranging from basic metabolism to molecular neuroscience. Their research is based on a variety of methods including molecular biology, protein biochemistry, enzymology, structural biology (protein crystallography), cell biology, bioinorganic chemistry and biotechnology.

DR. JÜRIG STREULI



Jürg Streuli is head of the Pediatric Palliative and Advanced Care Team at the Children's Hospital of Easter Switzerland, Senior Researcher and Research group leader for Pediatric Bioethics at the Institute of Biomedical Ethics at the University of Zurich. Dr. Streuli studied medicine (state examination 2005) and specialized in pediatric and adolescent medicine at the Triemli City Hospital (2006-2008) and at the University Children's Hospital Zurich (2012-2015). In 2014, he obtained his doctoral degree in biomedical ethics and law from the University of Zurich. In his research and clinical work, Dr. Streuli aims to assess, develop and support integrated and universal health care services that enables both local accessibility for support and management, and access to specialist services for the care of complex medical conditions.

In this context, he has been the new head of the Interdisciplinary Institute for Ethics in Healthcare at the "Stiftung Dialog Ethik" since March 2024.

HANSRUEDI SILBERSCHMIDT



Hansruedi Silberschmidt is a patient with the primary ciliary dyskinesia (PCD). From the first day of his life he had health problems (several respiratory arrests, infections). Many hospital stays followed during his childhood. In his autobiography «husten verboten» (coughing prohibited), he describes his life as a child and young adult with an undiagnosed disease and his path of suffering until he was finally diagnosed at the age of 39 at the University Hospital in Zurich.

He completed the teacher training and studied psychology at the IAP, Zurich. He worked as the executive director of andante Winterthur - a non-profit, social enterprise for the benefit of people with a cognitive and/or cerebral impairment as well as for people with a brain injury.

PROF. MARSHALL SUMMAR



Marshall Summar is the CEO of Uncommon Cures and Professor of Pediatrics George Washington University. Prof. Summar received his MD degree from the University of Tennessee, completed a residency and internship in pediatrics and medical/ biochemical genetics at Vanderbilt University. He is board-certified in pediatrics, clinical genetics and medical biochemical genetics. His research is focused on devices and treatments for patients with genetic and biochemical diseases and adapting knowledge from rare diseases to mainstream medicine. His work has resulted in new drugs undergoing Food and Drug Administration (FDA) trials for patients with congenital heart disease and organic acidemias. Clinically, he launched, and leads the first clinical Rare Disease Institute (RDI) at Children's National Hospital. He is involved in the WHO's effort to globally expand care and diagnosis for rare disease patients and in public policy around rare diseases and orphan drug development in the U.S.