



SSMG

KINDERKLINIK

Annual Meeting 2024

WITH YOUNG INVESTIGATOR DAY

May 16th and 17th, 2024

Bern University Hospital, Auditorium Ettore Rossi

Conference Committee



Christiane Zweier
University Hospital of Bern



André Schaller
University Hospital of Bern

Keynote Speakers 2024



Thomas Bourgeron
Université Paris Cité:
Phenotypic effects of genetic
variants associated with autism



Christa Flück
University Hospital of Bern:
Differences of Sex Development
(DSD) – a clinical and genetic
overview



Nadine Hornig
University of Kiel:
Molecular insights into androgen
insensitivity



Anne Gregor
University Hospital of Bern:
Disease modeling in
neurodevelopmental disorders

ORGANIZERS

Christiane Zweier
University Hospital of Bern

André Schaller
University Hospital of Bern

CREDIT POINTS

SGMG
8 CME points
Thursday/Friday

FAMH
8 CME points
Thursday/Friday

REGISTRATION

Medworld AG
Sennweidstrasse 46
6312 Steinhausen



<https://sgmg.ch/de/meeting-2024>



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Thursday, May 16th, 2024 – SSMG Business Meeting & General Assembly

Chairpersons: Isabel Filges, Basel & Naomi A. Porret, Bern

From 08:30	Registration open
09:45 – 12:15	Business Meeting of the Swiss Society of Medical Genetics
12:15 – 13:30	Lunch, Poster and Exhibition Viewing
13:30 – 14:00	Lunch Symposium 1 organized by AstraZeneca

Thursday, May 16th, 2024 – Young Investigator Day

PART I & II – PRESENTATIONS OF THE FINALISTS: YOUNG INVESTIGATOR AWARD 2024

Chairpersons: Christiane Zweier, Bern & Sven Cichon, Basel

14:05 – 14:15	Welcome address Christiane Zweier, Bern & Sven Cichon, Basel
14:15 – 15:55	SSMG Young Investigator Presentations with Q&A (20 min each)
15:55 – 16:25	Short Break
16:25 – 17:45	SSMG Young Investigator Presentations with Q&A (20 min each)

PART III – EVENING LECTURE

Chairpersons: André Schaller, Bern

17:45 – 18:30	Phenotypic effects of genetic variants associated with autism Thomas Bourgeron, Paris
18:30 – 18:35	Closing Remarks Day 1 André Schaller, Bern
From 18:35	Apéro at the University Hospital of Bern
From 20:00	Dinner at Kornhauskeller in Bern

Friday, May 17th, 2024- Scientific Meeting Day

Chairperson: Christiane Zweier, Bern

08:20 – 08:50	Satellite Symposium 2 organized by Oxford Nanopore Technologies
09:00 – 09:15	Opening and Welcome address Christiane Zweier, Bern & André Schaller, Bern
09:15 – 09:25	Announcement of the Winner of the SSMG Young Investigator Awards 2024 Sven Cichon, Basel
09:25 – 10:10	Differences of Sex Development (DSD) – a clinical and genetic overview Christa Flück, Bern
10:10 – 10:55	Molecular insights into androgen insensitivity Nadine Hornig, Kiel
10:55 – 11:15	Break
11:15 – 12:00	Disease modeling in neurodevelopmental disorders Anne Gregor, Bern
12:00 – 12:30	Satellite Symposium 3 organized by PacBio
12:30 – 12:40	Closing Remarks Day 2 Christiane Zweier, Bern & André Schaller, Bern
From 12:40	Farewell Lunch



Thursday, May 16th, 2024 – Oral presentation

▶ On site meeting

14:15 – 14:35 h	Oral Presentation 1 Rare copy-number variants as modulators of common disease susceptibility Chiara Auwerx, University of Lausanne
14:35 – 14:55 h	Oral Presentation 2 Missense variants in ANO4 cause sporadic encephalopathic or familial epilepsy with evidence for a dominant-negative effect Anais Begemann, University of Zurich
14:55 – 15:15 h	Oral Presentation 3 Deleterious ZNRF3 germline variants as a novel cause of neurodevelopmental disorders with mirror brain phenotypes due to distinct domain-specific effects on Wnt/β-catenin signaling Paranchai Boonsawat, University of Zurich
15:15 – 15:35 h	Oral Presentation 4 Deregulated of ion channels contribute to RHOTB2 associated developmental and epileptic encephalopathy Franziska Langhammer, University of Bern
15:35 – 15:55 h	Oral Presentation 5 Bi-allelic variants in BRF2 are associated with perinatal death and craniofacial anomalies Francesca Mattioli, University of Lausanne
15:55 – 16:25 h	<i>Short Break</i>
16:25 – 16:45 h	Oral Presentation 6 Novel insights into tumorigenesis revealed by molecular analysis of Lynch syndrome cases with multiple colorectal tumors Alisa Olkinuora, University Hospital Basel
16:45 – 17:05 h	Oral Presentation 7 Biallelic missense variants in CSMD2 are associated with a neurodevelopmental disease and epilepsy Clara Pailler-Pradeau, University of Lausanne
17:05 – 17:25 h	Oral Presentation 8 Further characterization of BRSK2-associated neurodevelopmental disorder Palak Singhal, University of Bern
17:25 – 17:45	Oral Presentation 9 Long-read whole-genome sequencing analysis of rare structural risk variants in families with bipolar disorder Priyadarshini Thirunavukkarasu, University Hospital Basel



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Thursday, May 16th, 2024 – Poster presentation

▶ On site meeting

POSTER 1	Poster presentation 1 Structural and functional insights from single cell transcriptional profiles of pituitary tumors Maxime Brunner, CHUV Lausanne
POSTER 2	Poster presentation 2 Identification of microbiota components correlated with host lifestyle, molecular, biochemical, immunophenotypic measurements and genotype in a deeply phenotyped Sardinian cohort Maria Antonietta Diana, CHUV Lausanne
POSTER 3	Poster presentation 3 Current Genetic Counselling Practice in Paediatric Oncology in Switzerland A qualitative Study Corinne Gemperle, University Children's Hospital Zurich
POSTER 4	Poster presentation 4 Search for modifier genes influencing disease severity in Hereditary Angioedema Asensio Gonzales, University Hospital of Basel
POSTER 5	Poster presentation 5 Health care transition to adulthood in young patients with rare genetic diseases: exploring the contributions of genetic counselling and genetic counsellors Sabrina Marti, Cantonal hospital Aarau
POSTER 6	Poster presentation 6 Long-read sequencing in SV calling and variant phasing Janine Meienberg, Centre for Cardiovascular Genetics, Schlieren
POSTER 7	Poster presentation 7 Genetic host factors in variation on viral control in HIV-infected individuals Sergey Oreshkov, CHUV Lausanne
POSTER 8	Poster presentation 8 Reclassification of variants of uncertain clinical significance in mendelian cardiomyopathies and arrhythmias using targeted blood RNA analysis Thomas Rio Frio, University Hospital of Geneva
POSTER 9	Poster presentation 9 Further delineation of SCAF4 associated neurodevelopmental disorder Cosima Schmid, University of Bern
POSTER 10	Poster presentation 10 Molecular trajectories of embryonic GnRH neurons associate with distinct biological processes and gene sets linked to human reproduction Yassine Zouaghi, CHUV Lausanne

Sponsors

EXHIBITORS

- | | |
|-----------------|------------------------------|
| ADS Biotec | Nostos Genomics |
| AmpliTech | Oxford Nanopore Technologies |
| Bencard GmbH | PacBio |
| Devyser GmbH | Roche |
| Illumina | Sysmex |
| Life & Brain | ThermoFischer |
| Macrogen Europe | |

SATELLITES

- AstraZeneca
- Oxford Nanopore Technologies
- PacBio

Industry partner

