

Day 1: Monday 24th of November

09:30–10:00: Registration, coffee and sandwiches

SESSION 1: Basics and background

10:00–10:15 Introduction of course and participants

10:15–10:45 Why does epileptology need genetics and precision medicine?

Tommy Stöddberg, MD, PhD

10:45–11:20 Genetics: Basics and methods

Kaja Selmer, MD, PhD

11:20–12:00 Genetic evaluation, diagnosis, and counselling

Inger-Lise Mero, Clinical Geneticist, MD, PhD

12:00–13:00 Lunch break

SESSION 2: Epilepsy genetics

13:00–13:45 State of the art of genetic investigations in epilepsy in 2025

Rikke Steensbjerre Møller, Professor, MSc, PhD

13:45–14:30 A genotype-phenotype clinical interpretation approach

Elena Gardella, Prof., MD PhD and Rikke Steensbjerre Møller, Prof, MSc PhD

14:30–14:45 Mini break – leg stretch

14:45–15:45 *In silico* prediction tools and online portals for variant interpretation

Christian Bosselmann, MD, PhD

15:45–16:15 Mini break – with snacks

SESSION 3: Workshop and hands on variant evaluation

16:15–16:45 Introduction to case workshops on variant evaluation

16:45–17:45 Case work in groups

17:45–18:15 Round-up in plenum

18:15–18:30 Summary of day 1

1930 Dinner

Day 2: Tuesday 25th of November, 2025

SESSION 4: Precision treatment

09:00–09:30 Patient perspectives

09:30–10:00 Somatic mutations in focal cortical dysplasia

Fridny Heimisdottir, Pediatrician, MD

10:00–10:40 Principles of clinical pharmacology and pharmacogenetics

Cecilie Johannessen Landmark, Professor, MScPharm, PhD

10:40–11:10 Drug repurposing in epilepsy

Allan Bayat, Ass. Professor, MD, PhD

11:10–11:30 Break

11:30–12:30 Precision medicine – state of the art and future perspectives

Amy McTague, MD, PhD

12:30–13:30 Lunch break

SESSION 5: Clinical case discussions

13:30–15:00 Plenary seminar: Cases from participants

15:00–16:00 Clinical discussions, evaluations, conclusions

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