Our popular Genomics of Rare Disease conference, now 13th in the series, offers an excellent multi-disciplinary forum for clinicians (consultants and trainees), research and clinical scientists, bioinformaticians and technology developers, to explore how genomic research translates into clinical care and informs our understanding of the biology of rare disease.

**KEY TOPICS**
- Solving the unsolved
- Informatics
- Developmental genetics
- Electronic health records
- Functional genomics
- Data sharing – legal and ethical issues
- Interaction of polygenic and monogenic disorders
- New therapeutic approaches

**KEYNOTE SPEAKERS**
- Nancy Cox Vanderbilt University School of Medicine, USA

**CONFIRMED SPEAKERS**
- William Gahl National Institutes of Health, USA
- Frank Kaplan University of Pennsylvania, USA
- Katrin Männik University of Lausanne, Switzerland
- Jennifer Posey Baylor College of Medicine, USA
- Heidi Rehm Massachusetts General Hospital, USA
- Malte Spielmann Max Planck Institute for Molecular Genetics, Germany
- Clara van Karnebeek Amsterdam University Medical Centres, The Netherlands
- Rosanna Weksberg The Hospital for Sick Children, Canada

**DEADLINES**
- Early bird: 02 January
- Bursaries: 16 January
- Abstracts: 30 January
- Registration: 27 February

More info and register:
bit.ly/GRD2019