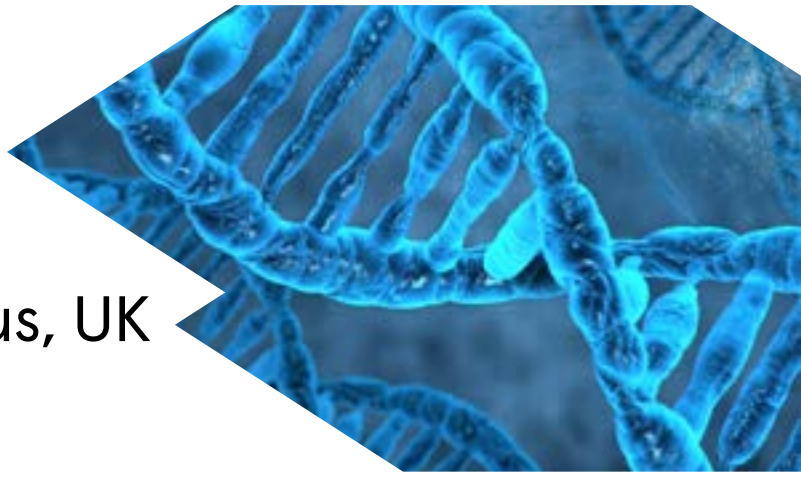
GENOMICS OF RARE DISEASE



27-29 March 2019
Wellcome Genome Campus, UK

Our popular Genomics of Rare Disease conference, now 13th in the series, offers a 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

DEADLINES

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

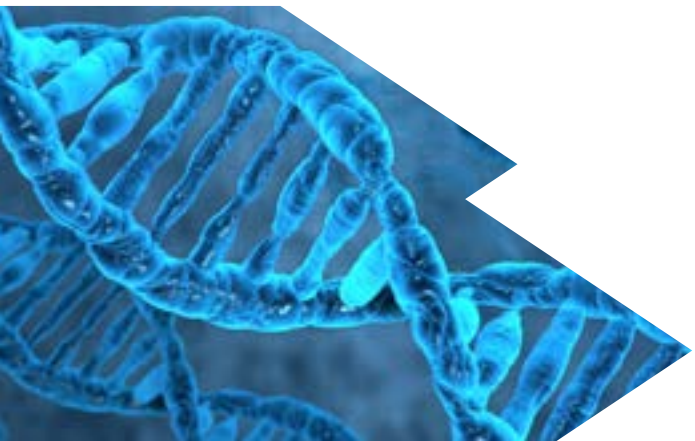
More info and register:
bit.ly/GRD2019

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



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