

SYMPOSIUM GENOME SEQUENCING AND PRECISION MEDICINE FOR TYPE 2 DIABETES

17th-18th November 2022



Supported by:

SYMPOSIUM PROGRAM

17th Nov

- 15.30 IMPaCT, a Spanish Precision Medicine Initiative**
Chair: Dídac Mauricio
Welcome
Genomics in the IMPaCT Spanish Personalized Medicine Infrastructure
Mónica Bayes, CNAG
The IMPaCT-T2D initiative
Silvia Bonàs-Guarch, Center for Genomic Regulation

16.30 COFFEE BREAK

- 17.00 Keynote lectures**
Chair: Urko M. Marigorta
Human Genetics in T2D Drug Discovery and Development: A Tale from AstraZeneca's Genomics Initiative
Slavé Petrovski, Centre for Genomics Research, AstraZeneca
Embracing the complexity of complex disease
Marylyn D. Ritchie, University of Pennsylvania

18.00 End of first day

18th Nov

- 9.00 Genomic insights into Mendelian Diabetes**
Chair: Luis Castaño
Modeling non-coding genome defects underlying diabetes
Jorge Ferrer, Center for Genomic Regulation
Correctly identifying and treating pathogenic coding variants causing monogenic diabetes
Andrew Hattersley, University of Exeter
- 10.20 COFFEE BREAK**
- 10.50 Leveraging complex trait genetics for predictive medicine**
Chair: Arcadi Navarro
The genetic of human height, a model complex trait
Peter Visscher, University of Queensland
Translational genomics of cardiometabolic traits
Eleftheria Zeggini, Helmholtz Zentrum München
From GWAS to function – strategies to gain mechanistic disease insight
Danielle Posthuma, Vrije Universiteit Amsterdam
- 12.50 LUNCH**
- 14.00 Genomics and precision medicine for common forms of diabetes**
Chair: Ana M. Wägner
Type 1 diabetes genetic risk scores for improved classification and prediction of diabetes
Richard Oram, University of Exeter
Towards precision medicine in diabetes: using genetics to understand disease heterogeneity
José Florez, University of Harvard
- 15.20 Closing**
- 15.30 End of the Symposium**