Pathways to the Clinical Implementation of Genome-Wide Sequencing for Rare Diseases:
Community Consultation on Genome Canada’s Precision Health Strategy

Genome Canada will host a panel discussion on the impact, challenges and opportunities for the implementation of genome-wide sequencing (GWS) as standard of care for rare diseases.

**Moderator**
- **Peter Goodhand**
  CEO, The Global Alliance for Genomics and Health (GA4GH)

**Panel Members**
- **Cindy Bell**
  Executive Vice President, Corporate Development, Genome Canada
- **William T. Gibson**
  Professor of Medical Genetics, University of British Columbia
  Senior Clinician Scientist, Laboratory for Obesity Genetics and Indirect Calorimetry (LOGIC), BC Children’s Hospital Research Institute-Investigator, BC Children’s Hospital
- **Eriskay Liston**
  Genetic Counsellor, The Hospital for Sick Children
- **Christian R. Marshall**
  Clinical Laboratory Director
  Genome Diagnostics, Department of Paediatric Laboratory Medicine
  The Hospital for Sick Children
- **Wyeth Wasserman**
  Executive Director, BC Children’s Hospital Research Institute
  Associate Dean for Research, Faculty of Medicine, University of British Columbia
- **Durhane Wong-Rieger**
  President and CEO, Canadian Organization for Rare Disorders (CORD)