9.30 - 9.45am  Welcome address
Dr Chris Gyngell

9.45 - 11.15am  Session 1: Rapid Genomic Sequencing in Critically Ill Children
Chair: Dr Chris Gyngell
Prof Zornitza Stark  Overview of the Acute Care Genomics Program
Katie Arkell  Health professionals’ experiences with making treatment decisions using RGS
Kirsten Boggs  Case studies of using RGS to make treatment decisions
Dr Julian Koplin  Ethical analysis of using RGS to make treatment decisions

11.15 - 11.45am  Morning tea

11.45 - 1.00pm  Session 2: Additional Findings in Paediatrics
Chair: Dr Danya Vears
Prof Clara Gaff  Offering additional findings in the Acute Care Genomics Program
Dr Lilian Downie  Offering genomic sequencing and additional findings to children with hearing loss
Dr Danya Vears  Genetic health professionals’ and parents’ perspectives on offering additional findings to children with hearing loss

1.00 - 2.00pm  Lunch

2.00 - 3.25pm  Session 3: Panel discussion: Are we ready for Genomic Newborn Screening?
Prof Lynn Gillam (Chair)  Prof John Christodoulou
Prof John Massie  Prof Julian Savulescu
Prof Ainsley Newson  Dr Meg Wall

3.25 - 3.30pm  Closing remarks
Dr Chris Gyngell