

9.30 - 9.45am Welcome address

Dr Chris Gyngell

9.45 - 11.15am Session 1: Rapid Genomic Sequencing in Critically III 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

Session 2: Additional Findings in Paediatrics 11.45 - 1.00pm

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 Massie
Prof Julian Savulescu

Prof Ainsley Newson Dr Meg Wall

3.25 - 3.30pm Closing remarks

Dr Chris Gyngell