



14th Glasgow Paediatric Research Day Friday 3rd November 2023 QEUH & RHC Teaching & Learning Centre

0900 0930 – 0935	Registration Welcome Dr Helen McDevitt, Consultant in Neonatology and Leonard Gow Lecturer in Child Health
Session 1 Chair	Short Oral Presentations Dr Martina Rodie, Consultant in Neonatology, Honorary Senior Lecturer and Lead for the Office for Rare Conditions, Glasgow
0935 - 0941	Differences between necrotising enterocolitis and focal intestinal perforation in a cohort of preterm infants: Judith Simpson, NICU, Royal Hospital for Children, Glasgow
0941 - 0947	Modelling Barth Syndrome: understanding clinical signs and identifying therapeutic targets: Freya Nye, Cancer Research UK Beatson Institute
0947 - 0953	Mepolizumab in children and adolescents with severe eosinophilic asthma not eligible for Omalizumab: a single centre experience: Yu Ting Lim, Department of Respiratory Paediatrics, Royal Hospital for Children, Glasgow
0953 - 0959	Blood pressure and eGFR in boys with hypospadias: a single centre experience : Iman Malik, School of Medicine, Dentistry & Nursing, University of Glasgow
0959 – 1005	ROS scavengers may improve genital skin healing in boys with hypospadias: Angela Lucas-Herald, Developmental Endocrinology Research Group, University of Glasgow
Session 2 Chair	Short Oral Presentations Mr Stuart O'Toole, Consultant in Paediatric Urology, Associate Clinical Professor and Barclay Lecturer in Child Health, Glasgow
1005 - 1011	Evaluating novel mutant-selective PDE4D PROTACs for the treatment of Acrodysostosis Type 2: <i>George Baillie, School of Cardiovascular and Medical Science, University of Glasgow</i>
1011 - 1017	Longitudinal change in external masculinization score in boys with xy disorders of sex development (DSD): Malika Alimussina, Developmental Endocrinology Research Group, University of Glasgow
1017 - 1023	Is MRI Molecular imaging accurate: Alexander Fletcher, School of Cardiovascular and Metabolic Health, University of Glasgow





1023 - 1029	Dependency upon SCD1 in CNS leukaemia reveals vulnerability to ferroptosis: Nikolai Gajic, Halsey Group, School of Cancer Sciences, University of Glasgow
1029 - 1035	Late mortality in children with oesophageal atresia (OA) in a unique 45-year cohort: Lewis Campbell, Neonatal Unit, Royal Hospital for Children, Glasgow
1035 - 1100	Coffee break
Session 3	Invited Speakers' Session
Chair	Dr Ruth McGowan, Consultant in Clinical Genetics, Honorary Clinical Associate Professor, Glasgow
1100 - 1140	The DDD study: Genomic Diagnosis of Rare Paediatric Disease in the United Kingdom and Ireland Prof Helen Firth, Addenbrooke's Hospital, Cambridge
1140 - 1200	Diagnostic trio-based whole exome sequencing for severe developmental disorders in Scotland Dr Morad Ansari, South-East Scotland Genetic Service, Western General Hospital, Edinburgh
1200 - 1220	Questions
Session 4	Child Health Prizes Professor S. Faisal Ahmed, Samson Gemmell Chair of Child Health, Glasgow
1220 - 1225	Child Health Prizes & Awards for 2023
1225 - 1230	The failure of agencies to protect Matilda Wormwood from harm : A case study Freya Semple: 2023 Yorkhill Essay Prize Winner
1230 - 1330	Lunch & Posters
Session 5 Chair	Invited Speakers' Session Professor Daniela Pilz, Consultant Clinical Geneticist, West of Scotland Regional Genetics Service
1330 - 1410	Rapid genetic testing in neonates to avoid antibiotic hearing loss Professor William Newman, Manchester Centre for Genomic Testing & University of Manchester
1410 - 1440	Rapid exome sequencing for prenatal diagnosis: lessons learnt from the English NHS Genomic Medicine Service Dr Rhiannon Mellis, West of Scotland Centre for Genomic Medicine, Royal Hospital for Children, Glasgow
1440 - 1500	Questions
1500 - 1515	Coffee break





Session 6 Chair	Short Oral Presentations Dr Ross Langley, Consultant in Paediatric Respiratory Medicine, Honorary Senior Lecturer and NRS Fellow, Glasgow
1515 - 1521	A novel brain-penetrant kinase inhibitor targeting diffused intrinsic pontine glioma: Aditi Atmasidha, Division of Cellular and Systems Medicine, University of Dundee
1521 - 1527	Genotype: Phenotype relationships in Myhre syndrome: Madeline Pearson, School of Medicine, University of Dundee
1527 - 1533	An international study of the association between local health care resources and acute adrenal insufficiency events in children with congenital adrenal hyperplasia: Xanthippi Tseretopoulou, Developmental Endocrinology research Group, University of Glasgow
1533 - 1539	Acute adrenal insufficiency related adverse events in children with congenital adrenal hyperplasia (CAH): Changes during the period 2019-2022 in I-CAH: Xanthippi Tseretopoulou, Developmental Endocrinology Research Group, University of Glasgow
1539 - 1545	Development and validation of short versions of quality of life questionnaires for parents of children with disorders/differences of sex development: Salma Ali, Developmental Endocrinology Research Group, Royal Hospital for Children, Glasgow
1545 -1555	Short break
1555 - 1600	Prize winners / feedback survey / close of meeting Dr Helen McDevitt https://link.webropol.com/s/GPRD2023Feedback

The Glasgow Paediatric Research Day is kindly supported by The University of Glasgow and NHS Greater Glasgow & Clyde



