



Pre-meeting
Monday 29 June 2026

Annual Symposium
Tuesday 30 June -
Wednesday 1 July 2026



Poster List

Posters listed by presenting day then alphabetical by Presenting Author Surname.

All details are subject to change, including board numbers

Board Number	Poster	Theme	Presenting Day
T01	Bloodspot phenylalanine monitoring frequency in adults with phenylketonuria: A service improvement project Sarah Bailey ¹ ¹ All Wales Inherited Metabolic Disorder Service, Cardiff, Wales	Audit	Tuesday
T02	Two cases of taurine transporter (SLC6A6) deficiency presenting with retinopathy and severe hypotaurinaemia Roger Bramley ¹ , Dr Sergei Korenev ² , Robert Barski ¹ ¹ Leeds Teaching Hospitals NHS Trust, UK, ² Manchester University NHS Foundation Trust, UK	Clinical	Tuesday
T03	Elevated 2,3-dihydroxy-2-methylbutyric acid on urine organic acid analysis: a case series Roger Bramley ¹ , Robert Barski ¹ ¹ Leeds Teaching Hospitals NHS Trust, UK	Clinical	Tuesday
T04	Galactose-1-phosphate uridylyltransferase (GALT) testing for Classical Galactosaemia (CG): indications and impact on human milk feeding (HMF) Laura Byers ¹ , Dr Anupam Chakrapani ² , Dr James Davison ² , Katie Harvey ⁴ , Kelly Watson ¹ , Camilla Natale ³ , Marjorie Dixon ¹ ¹ Dietetics Department, Great Ormond Street Hospital, NHS Foundation Trust, London, UK, ² Department of Metabolic Medicine, Great Ormond Street Hospital, London, UK, ³ MSc Dietetics, UCL Division of Medicine, University College London, London, UK, ⁴ Chemical Pathology, Paediatric Laboratory Medicine, Great Ormond Street Hospital, London, UK	Audit	Tuesday
T05	Determinants of successful human milk feeding in inherited metabolic disorders (IMD) on dietary treatment Laura Byers ¹ , Ida Herlihy ¹ , Chloe Millington ¹ , Dr Anupam Chakrapani ² , Dr Spyros Batzios ² , Marjorie Dixon ¹ ¹ Dietetics Department, Great Ormond Street Hospital, London, UK, ² Department of Metabolic Medicine, Great Ormond Street Hospital, London, UK	Audit	Tuesday

T06	Protein Substitutes: challenges applying International Dysphagia Diet Standardisation Initiative standards (IDDSI) for an aging PKU community Lisa Gaff¹ , Simone Whiteway ² ¹ Cambridge University Hospitals NHS Foundation Trust, Cambridge, England, ² North Bristol NHS Trust, Westbury-on-Trym, England	Clinical	Tuesday
T07	An unusual presentation of Alpha-methylacyl-CoA racemase deficiency in a patient with elevated creatine kinase Dr Laura Hancox¹ , Dr Wycliffe Mbagaya ¹ , Dr Charlotte Dawson ² ¹ University Hospitals Bristol and Weston NHS Foundation Trust, UK, ² University Hospitals Birmingham NHS Foundation Trust, UK	Clinical	Tuesday
T08	PAH genetics and sapropterin responsiveness in paediatric NI patients Dr Caroline Hart¹ , Dr Siobhan O'Sullivan ¹ Royal Belfast Hospital for Sick Children, Northern Ireland	Audit	Tuesday
T09	followME Fabry Pathfinders registry: cardiac and renal effectiveness in a multi-national, multi-centre cohort of patients on migalastat treatment for ≥5 years Prof Derrallynn Hughes¹ , Joseph Giuliano ² , Prof Aleš Linhart ³ , Prof Antonio Pisani ⁴ , Prof Olga Azevedo ⁵ , Jennifer Hiros ⁶ , Vipul Jain ⁶ , Dr Biliانا Veleva-Rotse ⁶ , Dr Peter Nordbeck ⁷ ¹ Lysosomal Storage Disorders Unit, Royal Free London NHS Foundation Trust and University College London, UK, ² Rare Medical Affairs LLC, USA, ³ Second Department of Internal Cardiovascular Medicine, First Faculty of Medicine, Charles University and General University Hospital, Czechia, ⁴ Department of Public Health, Nephrology Unit, Federico II University Hospital, Italy, ⁵ Cardiology Department, Reference Center of Lysosomal Storage Disorders, MetabERN member, Hospital Senhora da Oliveira, Portugal, ⁶ Amicus Therapeutics, Inc., USA, ⁷ Internal Medicine I, University Hospital Würzburg, Germany	Clinical	Tuesday
T10	Hereditary Fructose Intolerance presenting with hepatic dysfunction and hypoglycaemia and prior to exposure to dietary fructose: Diagnostic Pitfalls Parveen Leghari¹ , Catherine Zephir ¹ , Bernd C Schwahn ^{1,2} ¹ Manchester Centre for Genomic Medicine, St Mary's Hospital, Manchester University NHS Foundation Trust, Health Innovation, UK, ² Division of Evolution & Genomic Sciences, School of Biological Sciences, Faculty of Biology, Medicine and Health, University of Manchester, Manchester, UK	Clinical	Tuesday
T11	Cardiac involvement in adults with Glycogen Storage Disease Type IIIa: an observational cohort study Dr Eamon Mccarron¹ , Dr Adam Lomas ¹ , Dr Godfrey Gillett ¹ , Andrew Oldham ² , Dr Alison Woodhall ² , Dr Rob Cooper ³ , Dr Oliver Watson ⁴ , Dr Ana Jovanovic ² , Dr Reena Sharma ² , Dr Karolina Stepień ² ¹ Adult Inherited Metabolic Disorders, Sheffield Teaching Hospitals NHS Foundation Trust, UK, ² Adult Inherited Metabolic Disorders, Salford Care Organisation, Northern Care Alliance NHS Foundation Trust, UK, ³ Cardiology Department, Liverpool Heart and Chest Hospital NHS Foundation Trust, UK, ⁴ Cardiology Department, Northern General Hospital, Sheffield Teaching Hospitals NHS Foundation Trust, UK	Research	Tuesday

T12	From Guidance to Practice: An Audit of PKU Management in Northern Ireland Dr Kerri Munn-bookless¹, Dr Niamh McCool² , Dr Caroline Hart ¹ ¹ Paediatric Metabolic Medicine, RBHSC, Northern Ireland, ² Paediatric Haematology Oncology, RBHSC, Northern Ireland	Audit	Tuesday
T13	Impact on biochemical markers following reduced dose olipudase alfa in adults with acid sphingomyelinase deficiency Dr Elaine Murphy¹ , Kirsty White ¹ , Ana Amado Fondo ¹ , Dr Robin Lachmann ¹ ¹ University College London NHS Foundation Trust, UK	Clinical	Tuesday
T14	7.5-year efficacy outcomes with cipaglucosidase alfa plus miglustat in adults with Pompe disease in ATB200-02, an open-label Phase I/II study Prof Mark Roberts ¹ , Dr Drago Bratkovic ² , Dr Barry J Byrne ³ , Prof Paula R Clemens ⁴ , Dr Ozlem Goker-Alpan ⁵ , Prof Priya S Kishnani ⁶ , Prof Tahseen Mozaffar ⁷ , Dr Nadine AME van der Beek ⁸ , Farah Amon ⁹ , Dr Brian Fox ⁹ , Dr Fred Holdbrook ⁹ , Vipul Jain ⁹ , Prof Benedikt Schoser ¹⁰ , Dr Samreen Safdar¹¹ ¹ Salford Royal NHS Foundation Trust, UK, ² PARC Research Clinic, Royal Adelaide Hospital, Australia, ³ University of Florida, USA, ⁴ Department of Neurology, University of Pittsburgh School of Medicine and VA Pittsburgh Healthcare System, USA, ⁵ Lysosomal and Rare Disorders Research and Treatment Center, USA, ⁶ Duke University Medical Center, USA, ⁷ Department of Neurology, University of California, USA, ⁸ Erasmus MC University Medical Center, Netherlands, ⁹ Amicus Therapeutics, Inc., USA, ¹⁰ Friedrich-Baur-Institute at the Department of Neurology, LMU University Hospital, LMU Munich, Germany, ¹¹ Northern Care Alliance NHS Foundation Trust, UK	Clinical	Tuesday
T15	Long-term complications in methylmalonic acidemia in adults Dr Samreen Safdar¹ , Mairead Jones ¹ , Alison Woodall ¹ , Priscilla Sia ¹ , Faiza Adrees ¹ , Dr John Bassett ¹ , Dr Jonathan Meyer ¹ , Dr Karolina M Stepień ¹ , Dr Ana Jovanovic ¹ , Dr Reena Sharma ¹ ¹ Salford Royal Hospital, Northern Care Alliance NHS Foundation Trust, UK	Clinical	Tuesday
T16	Multifactorial Progressive Osteoporosis in Adults with PMM2-CDG: A Five-Patient Case Series Dr Samreen Safdar¹ , Andrew Oldham ¹ , Janet Gorton ¹ , Dr Jonathan Meyer ¹ , Dr John Bassett ¹ , Dr Karolina Stepień ¹ , Dr Ana Jovanovic ¹ , Dr Reena Sharma ¹ ¹ Salford Royal Hospital, Northern Care Alliance NHS Foundation Trust, Salford, UK,	Clinical	Tuesday
T17	Intrafamilial phenotypic variability in m.3243A>G mitochondrial disease: Implications for cascade testing and multisystem surveillance Dr Samreen Safdar¹ , Andrew Oldham ¹ , Faiza Adrees ¹ ¹ Salford Royal Hospital, Northern Care Alliance NHS Foundation Trust, UK	Clinical	Tuesday
T18	Factors affecting long-term outcomes in a cohort of adults with LC-FAODs Zona Ismail ¹ , Dr John Bassett ¹ , Andrew Oldham ¹ , Alison Woodall ¹ , Dr Karolina M Stepień ¹ , Dr Ana Jovanovic ¹ , Dr Samreen Safdar¹ , Dr Reena Sharma ¹ ¹ Adult Inherited Metabolic Diseases, Salford Care Organisation, Northern Care Alliance NHS Foundation Trust, Salford, Greater Manchester, Salford, UK	Clinical	Tuesday

T19	When a Simple Test Saves a Life: Neonatal Galactosemia Diagnosed by Benedict Test in a Resource Limited Setting Dr Pavithra Samarakoon ¹ , Dr Harsha Minasinghe ² , Dr Thayani Arooran ¹ , Dr Y Hemika ¹ , Dr MFS Mihika ¹ , Dr KD Rammuthupura ¹ , Dr E Jasinghe ¹ ¹ Department of chemical Pathology, Colombo, Sri Lanka, ² Department of Paediatrics, Colombo, Sri Lanka	Clinical	Tuesday
T20	Diagnostic Delay and Divergent Phenotypes in Metachromatic Leukodystrophy: Real-World Evidence Supporting Newborn Screening in the Era of Gene Therapy Dr Berna Seker Yilmaz ^{1,2} , Dr Anupam Chakrapani ³ , Dr James Davison ³ , Dr Helen Mundy ² , Dr Preeya Rehsi ³ , Dr Roshni Vara ² , Prof Paul Gissen ^{1,3}	Clinical	Tuesday
T21	Rare Disease Awareness in Young People in the UK: Insights from a Survey Gabriel Tipper ¹ , Zeynep Ada Yilmaz ² , Prof Paul Gissen ^{3,4} , Dr Berna Seker Yilmaz ^{3,5} ¹ Department of Biomedical Sciences, University of Exeter, UK, ² School of Film, Media, and Creative Technologies, University of Portsmouth, UK, ³ Genetics and Genomic Medicine Department, Great Ormond Street Institute of Child Health, University College London, UK, ⁴ Department of Paediatric Metabolic Medicine, Great Ormond Street Hospital for Children NHS Foundation Trust, UK, ⁵ Department of Paediatric Inherited Metabolic Disease, Evelina Children's Hospital, Guy's, and St Thomas' NHS Foundation Trust, UK	Research	Tuesday
T22	Review of prescribing practice in Urea Cycle Disorders- one adult centre experience Dr Karolina Stepien ¹ , Dr John Bassett, Dr Jonathan Meyer, Dr Reena Sharma, Dr Ana Jovanovic, Dr Samreen Safdar, Dr Alison Woodall, Faiza Adrees ¹ Salford Royal Hospital, Northern Care Alliance NHS Foundation Trust, Salford, UK	Clinical	Tuesday
T23	Pancreatitis in Two Adolescents with Argininosuccinic Aciduria: Expanding the Phenotypic Spectrum Chern Yan Tan ¹ , Dr Andrew Morris ¹ ¹ Willink Metabolic Unit, Manchester, UK	Clinical	Tuesday
T24	Natural History and Comorbidities of Childhood Ketotic Hypoglycaemia: Experience from a Regional Specialist Centre Chern Yan Tan ¹ , Dr Sameera Auckburally ² , Prof Indi Banerjee ² , Dr Arunabha Ghosh ¹ ¹ Willink Metabolic Unit, Manchester, UK, ² Department of Paediatric Endocrinology, Royal Manchester Children's Hospital	Clinical	Tuesday
T25	Timeliness in PKU Care: A National Audit into Turnaround Time of Dried Blood Spots and Frequency of appointments for Adults with PKU in the UK Simon Tapley ¹ , Benjamin Hayes ¹ , Natalie Shaw ² , Bailey Sarah ³ , Cameron Arbuckle ⁴ , Louise Robertson ⁵ , Lisa Gaff ⁶ , Imogen Hall ⁷ , Katie Yeung ⁷ , Nicola McStravick ⁸ , Alison McKee ⁸ ¹ University Hospitals Bristol and Weston, UK, ² Newcastle Upon Tyne Hospitals NHS Foundation Trust, UK, ³ University Hospital of Wales, UK, ⁴ University College London Hospitals, UK, ⁵ University Hospitals Birmingham, UK, ⁶ Cambridge University Hospitals, UK, ⁷ Guy's and St Thomas NHS Foundations Trust, UK, ⁸ Belfast Health and Social Care Trust, Northern Ireland	Audit	Tuesday

T26	Short-term use of a slow-release Phenylalanine-free microtablet protein substitute in children and adolescents with Phenylketonuria Dr Martina Tosi¹ , Dr Anne Daly ¹ , Dr Sharon Evans ¹ , Catherine Ashmore ¹ , Dr Alex Pinto ¹ , Prof Anita MacDonald ¹ ¹ Birmingham Women's and Children's Hospital, UK	Research	Tuesday
T27	Defining the inherited metabolic disorder landscape in numbers using structured databases: proportion and mortality across EU-5 Dr Elise Wilby¹ , James Stanway ² , Laura Smith Van Carroll¹ ¹ Metabolic Support, England, ² Keele University, England	Research	Tuesday
T28	Two-Year Sapropterin Outcomes in Phenylketonuria: Longitudinal Study of Metabolic, Dietary, Psychosocial Effects Dr Ozlem Yilmaz Nas^{1,2} , Catherine Ashmore ¹ , Maria Ines Gama ¹ , Dr Anne Daly ¹ , Dr Sharon Evans ¹ , Dr Alex Pinto ¹ , Prof Yahya Ozdogan ² , Prof Anita MacDonald ¹ ¹ Birmingham Women's and Children's NHS Foundation Trust, UK, ² Department of Nutrition and Dietetics, Ankara Yildirim Beyazit University, 06760, Turkey	Research	Tuesday

W01	Use of skin testing to advise therapy options in Pompe patients following suspected IgE mediated infusion associated reactions (IAR) Ana Amado Fondo¹ , Anna Katherina Cuarentas ¹ , Kirsty White ¹ , Tigisti Beyene ¹ , Dr Joanna Lukawska, Dr Elaine Murphy, Dr Robin Lachmann ¹ UCLH, London, UK	Clinical	Wednesday
W02	Review of genotype-phenotype correlation regarding sapropterin responsiveness in adult patients living with PKU Dr Kylie Beale¹ , Simon Tapley ¹ , Suzanne Ford ¹ , Dr Wycliffe Mbagaya ¹ , Dr Nathan Cantley ¹ ¹ Bristol NHS Group, UK	Clinical	Wednesday
W03	A single-centre patient experience survey of parents and carers of children on low protein diets Lucy Blackstone¹ , Victoria Holmes ¹ , Zixin Luo ¹ , Leah Paulden ¹ , Emma Simpson ¹ , Karen Van Wyk ¹ , Georgina Wood ¹ , Catherine Zephir ¹ ¹ Willink Biochemical Unit, Manchester, England	Research	Wednesday
W04	Variability in dietary management in Isovaleric Acidaemia (IVA): A retrospective longitudinal cohort study Sarah Cawtherley¹ , Louise Van Dorp ¹ , Cameron Arbuckle ⁴ , Dr Martina Messina ^{2,3} , Dr Spyros Batzios ² , Dr Emma Footitt ² , Dr Anupam Chakrapani ² , Rachel Skeath ¹ , Dr Julein Baruteau ² , Dr James Davison ² , Dr Stephanie Grunewald ² , Dr Alexander Broomfield ² , Dr Preeya Rehsi ² , Marjorie Dixon ¹ ¹ Dietetics, Great Ormond Street Hospital for Children, NHS Foundation Trust, London, UK, London, UK, ² Department of Paediatric Metabolic Medicine, Great Ormond Street Hospital for Children, NHS Foundation Trust, London, UK, London, UK, ³ Genetics and Genomic Medicine Department, GOS Institute of Child Health, University College London, London, UK, London, UK, ⁴ Charles Dent Metabolic Unit, University College London Hospitals, London, UK, London, UK	Audit	Wednesday

W05	Cholestatic Jaundice with Hypocalcaemia: One Pathway and a Unifying Diagnosis Sharon Colyer¹ , Claire Hart ¹ , Katherine Wright ¹ , Dr Elizabeth Griffiths ² ¹ Department of Clinical Chemistry, South Yorkshire & Bassetlaw Pathology, Sheffield Children's Hospital, UK, ² Department of Paediatric Gastroenterology, Sheffield Children's Hospital, UK	Clinical	Wednesday
W06	The holistic management of Late Onset Pompe Disease through nurse and physio led clinics-the Birmingham experience Sarah Steeds ¹ , Nicola Condon¹ , Rachel Curran¹ , Dr Charlotte Dawson ¹ ¹ University Hospitals Birmingham, UK, ² University Hospitals Birmingham, UK, ³ University Hospitals Birmingham, UK, ⁴ University Hospitals Birmingham, UK	Clinical	Wednesday
W07	Sleep Disturbance in Mucopolysaccharidosis Type III (Sanfilippo Syndrome): A Service Review of Pharmacological Management Efficacy and Safety Dr Daniela Castillo Garcia ¹ , Clare Hamilton-Kay ¹ , Sarah Phillippo ¹ , Rachel Tetlow ¹ , Dr Sarah Hulley ¹ , Dr Arubnabha Ghosh ¹ , Dr Richard Curnock ¹ , Dr Maria De Castro Lopez¹ ¹ St. Mary's Hospital, Manchester NHS Foundation Trust, Manchester, UK	Audit	Wednesday
W08	Peri-operative Management Guide for Inherited Metabolic Diseases Eva Fox¹ , Dr Caroline Hart² ¹ Ulster University, Northern Ireland, ² Royal Belfast Hospital for Sick Children, Northern Ireland	Clinical	Wednesday
W09	Development of an erythrocyte debrancher enzyme assay to biochemically confirm a diagnosis of GSD IIIb Rohit Hirachan¹ , Dr Ralph Wigley ¹ , Dr Alistair Horman ¹ ¹ Enzymes Unit, Chemical Pathology, Great Ormond Street Hospital for Children NHS Foundation Trust	Research	Wednesday
W10	Complications of Central Venous Access in Metabolic Patients Undergoing Hemofiltration Dr Nazreen Banu Kamarus Jaman¹ , Dr James Davison ¹ , Dr Preeya Rehsi ¹ , Megan Dorman ¹ , Dr Emma Footitt ¹ , Dr Rola Zaatra ¹ ¹ Great Ormond Street Hospital, UK	Audit	Wednesday
W11	Development of a selective HPPD inhibitor without an off-target liability Domen Kampjut¹ , Petros Tyrakis ¹ , Jonathan Lindström ¹ , Nigel Ramsden ² , Oliver Maddocks ¹ ¹ Faeth Therapeutics, Cambridge, UK, ² Rxcelerate, Cambridge, UK	Research	Wednesday
W12	Sleep study severity as a predictor of intraoperative airway difficulty in Mucopolysaccharidosis patients: a single-centre experience Dr Christos Lazaridis¹ , Dr Lois Mackley ² , Dr Marie-Claire Raphael ² , Dr Riona Tully ² , Dr Elena Targa ² , Mathilda Antonini ¹ , Rebecca Whiteley ¹ , Megan Keating ¹ , Dr Preeya Rehsi ¹ , Dr James Davison ¹ , Dr Anupam Chakrapani ¹ , Dr Emma Footitt ¹ , Dr Elaine Chan ³ , Dr Amaki Sogbodjor ² , Prof Paul Gissen ^{1,4} , Dr Alexander Broomfield ¹ ¹ Department of Paediatric Metabolic Medicine, Great Ormond Street Hospital for Children NHS Foundation Trust, UK, ² Department of Paediatric Anaesthetic Medicine, Great Ormond Street Hospital for Children NHS Foundation Trust, UK, ³ Department of Paediatric Respiratory Medicine, Great Ormond Street Hospital for Children NHS Foundation Trust, UK, ⁴ UCL Great Ormond Street Institute of Child Health, UK	Research	Wednesday

W13	Continuous glucose monitoring in assessment of susceptibility to ketotic hypoglycaemia in children diagnosed with spinal muscular atrophy in infancy on disease-modifying therapies Dr Sophie Manoy¹, Dr Marita Smith¹, Kate Munro², Antonia Wright², Dr Anita Cairns², Dr Matthew Lynch^{1,2} ¹Queensland Lifespan Metabolic Medicine Service, Queensland Children's Hospital, Australia, ²Department of Neurosciences, Queensland Children's Hospital, Australia	Research	Wednesday
W14	Glutathione Synthetase Deficiency: a rare differential diagnosis of neonatal metabolic acidosis Dr Hebatallah Mansour^{1,2}, Dr Chern Tan^{1,2}, Dr Bernd Schwahn^{1,2,3} ¹Royal Manchester Children's Hospital, UK, ²Manchester Centre for Genomic Medicine, St Mary's Hospital, Manchester University NHS Foundation Trust, Health Innovation Manchester, UK, ³Division of Evolution & Genomic Sciences, School of Biological Sciences, Faculty of Biology, Medicine and Health, University of Manchester, UK	Clinical	Wednesday
W15	Determinants of Bone Density Change in Gaucher Disease: A Multisite Mixed-Effects Analysis Dr Carmen Minea¹, Dr Boliang Yu¹, Dr Patrick B. Deegan¹ ¹Cambridge University Hospitals, England	Research	Wednesday
W16	Future proofing The Metabolic Service: MCADD Clinic – a nurse led approach Claire Mizler¹, Nichola Cowie¹, Hannah Dixon¹, Rebecca Herbert¹ ¹Sheffield Children's Hospital Foundation Trust, UK	Clinical	Wednesday
W17	A homozygous intronic variant in NDUFAF8 causes optic atrophy and learning difficulties Dr Maria Moustakis¹, K Kavanagh², LS Taylor^{3,4}, D Brosnahan⁵, S Chamney^{5,6}, K McCreery⁶, M O'Regan⁷, D O'Rourke⁸, RW Taylor^{3,4}, CL Alston^{3,4}, C Howard¹ ¹National Centre for Inherited Metabolic Disorders, CHI at Temple Street, Dublin 1, Ireland, ²National Centre for Inherited Metabolic Disorders, Mater Misericordiae University Hospital, Dublin 7, Ireland, ³Newcastle Highly Specialised Mitochondrial Diagnostic Service, Newcastle upon Tyne Hospitals NHS Foundation Trust, UK, ⁴Mitochondrial Research Group, Translational and Clinical Research Institute, Faculty of Medical Sciences., Newcastle , UK, ⁵Department of Paediatric Ophthalmology, CHI at Crumlin, Dublin 12, Ireland, ⁶Department of Paediatric Ophthalmology, CHI at Temple Street, Dublin 1, Ireland, ⁷Department of Paediatric Neurology, CHI at Crumlin., Dublin 12, Ireland, ⁸Department of Paediatric Neurology, CHI at Temple Street., Dublin 1, Ireland	Clinical	Wednesday

W18	Enhancing Enzyme Replacement Therapy in Lysosomal Storage Disorders: Faster Infusion Rates Across the UK Antonio Ochoa-ferraro¹, Zoe Taylor², Faiza Adrees³, Daniel Bell⁴, Susan McMahon⁵, Evelyn Frank⁶, Chin Gan⁷, Siema Akram⁸, Niamh Finnegan⁹, Robert Dugdale¹⁰, Clara Monaghan¹¹, Thomas Wyllie¹² ¹University Hospitals Birmingham NHS Foundation Trust, Birmingham, UK, ²All Wales Adult Inherited Metabolic Disease Service (AWIMDS), University Hospital of Wales, Cardiff and Vale University Health Board, UK, ³Salford Royal NHS Foundation Trust, Northern Care Alliance NHS Foundation Trust (NCA), UK, ⁴Cambridge University Hospitals NHS Foundation Trust, UK, ⁵Inherited Metabolic Disorders Service, Royal Hospital for Children, UK, ⁶University College London Hospitals NHS Foundation Trust, UK, ⁷Great Ormond Street Hospital NHS Foundation Trust, UK, ⁸Birmingham Children's Hospital, Birmingham Women's and Children's NHS Foundation Trust, UK, ⁹Royal Free London NHS Foundation Trust, UK, ¹⁰Royal Manchester Children's Hospital, Manchester University NHS Foundation Trust, UK, ¹¹Royal Belfast Hospital for Sick Children, Belfast Health and Social Care Trust, UK, ¹²Noah's Ark Children's Hospital for Wales, Cardiff and Vale University Health Board, UK	Clinical	Wednesday
W19	Metabolic Pharmacist Roles Across the UK: Current Practice and Service Provision Antonio Ochoa-ferraro¹, Robert Dugdale², Daniel Bell³, Georgina Smith⁴, Siema Akram⁵, James Roddick⁶, Priya Dattani⁷, Clayton Wong⁸, Clara Monaghan⁹, Faiza Adrees¹⁰, Trinh Huynh¹¹, Chin Gan¹², Alice Ruston¹³, Thomas Wyllie¹⁴ ¹University Hospitals Birmingham NHS FT, UK, ²Royal Manchester Children's Hospital, Manchester University NHS FT, UK, ³Cambridge University Hospitals NHS FT, UK, ⁴Nottingham University Hospitals NHS Trust, UK, ⁵Birmingham Children's Hospital, Birmingham Women's and Children's NHS FT, UK, ⁶Inherited Metabolic Disorders in Scotland (IMD Scotland), NHS Greater Glasgow and Clyde, UK, ⁷University Hospitals of Leicester NHS Trust, UK, ⁸Royal Free London NHS Foundation Trust, UK, ⁹Royal Belfast Hospital for Sick Children, UK, ¹⁰Salford Royal NHS Foundation Trust, Northern Care Alliance NHS Foundation Trust (NCA), UK, ¹¹Evelina London Children's Hospital, Guy's and St Thomas' NHS FT, UK, ¹²Great Ormond Street Hospital For Children NHS FT, UK, ¹³Sheffield Teaching Hospitals NHS FT, UK, ¹⁴Noah's Ark Children's Hospital for Wales, Cardiff and Vale University Health Board, UK	Clinical	Wednesday
W20	Service improvement: Clinical Nurse Specialist (CNS) led clerking of Neuronal Ceroid Lipofuscinosis (CLN2) Batten disease patients Katie Raven¹, Anja Keating¹, Dr Laura Guilder¹ ¹UHBW, Bristol, UK	Clinical	Wednesday
W21	Understanding the experiences and support needs of families living with Ketotic Hypoglycaemia to inform service delivery Dr Katie Redwood¹, Camille Newby, Naimo Abdullah ¹UHBW NHS, Bristol, UK	Clinical	Tuesday
W22	Galactosaemia and cheese: cheese analysis for lactose and galactose content in the UK, 2022–2026 Louise Robertson^{1,2}, Sue Bevington², Karen Ford², Simon Tapley³, Prof Anita MacDonald⁴ ¹University Hospitals Birmingham NHS Foundation Trust, UK, ²Galactosaemia Support Group, UK, ³University Hospitals Bristol and Weston NHS Foundation Trust, UK, ⁴Birmingham Women's and Children's Foundation Trust, UK	Research	Wednesday

W23	An Exercise and Dietary Induced Fat-Free Mass Increase in Two Patients with Phenylketonuria (PKU) Sarah Howe ¹ , Louise Robertson ² , Clare Dale ³ ¹ University Hospitals Birmingham, Birmingham, UK, ² University Hospitals Birmingham, Birmingham, UK, ³ University Hospitals Birmingham, Birmingham, UK	Clinical	Wednesday
W24	Characterization of Maple Syrup Urine Disease in a Chilean Cohort: Genotypic Variations and Metabolic Control Maria Florencia Salazar ¹ , Verónica Cornejo ¹ , Felipe Peñaloza ¹ , Eugenia Rojas-Agurto ¹ , Catalina Arancibia ¹ , Carolina Arias ¹ , Juan Francisco Cabello ^{1,2} , Pilar Peredo ¹ , Vicente Quiroz ¹ , María Jesús Leal-Witt ¹ ¹ Laboratory Of Metabolic Diseases, Inta, University of Chile, Chile, ² Department of Neurology, Boston Children's Hospital, USA	Research	Wednesday
W25	Estimating prevalence of long chain fatty acid oxidation disorders in the UK Saikat Santra ¹ , Erru (Christy) Yang ² , Lay Ean Tan ² , Nin Dass ² ¹ Birmingham Women's and Children's NHS Foundation Trust, UK, ² Ultragenyx Pharmaceutical Inc., UK	Research	Wednesday
W26	Multiple acyl CoA dehydrogenase deficiency (MADD): single-centre review of paediatric clinical, dietetic and biochemical disease severity spectrum Rachel Skeath ¹ , Erin Emmett ¹ , Alistair Horman ¹ , Ralph Wigley ¹ , Spyros Batzios ¹ , Laura Byers ¹ , Anupam Chakrapani ¹ , Sarah Cawtherley ¹ , Marjorie Dixon ¹ , Megan Dorman ¹ , Jennifer Dunne ¹ , Emma Footitt ¹ , Ide Herlihy ¹ , Mel McSweeney ¹ , Chloe Millington ¹ , Sophie Taylor ¹ , Karin Tuschl ¹ , James Davison ¹ ¹ Great Ormond Street Hospital for Children NHS Foundation Trust, UK	Clinical	Wednesday
W27	Cognitive and Behavioural Profiles of Children and Young People with Inherited Metabolic Disorders: A Retrospective Clinical Audit Molly Smith ¹ , Magdalini Hadjikal ¹ , Dr Clare Williams ¹ , Dr Emma Matthews ¹ ¹ Great Ormond Street Hospital NHS Foundation Trust, UK	Audit	Wednesday
W28	Paediatric Aldolase A deficiency: Case report, treatment & biomarkers of an ultra-rare inherited metabolic disorder Dr Dexter Tarr ¹ , Victoria Whiteley ¹ , Prof Judith JM Jans, James Cooper ¹ , Oliver Parkes ¹ , Dr Bernd Schwahn ¹ ¹ Manchester Foundation Trust, UK, ² UMC Utrecht, Netherlands	Clinical	Wednesday
W29	Lived Experience: Insights into Familial Chylomicronaemia Syndrome Dr Elise Wilby ¹ , Jill Prawer ² , Laura Smith van Carroll ¹ ¹ Metabolic Support, UK, ² Action FCS, UK	Research	Wednesday
W30	Single-centre experience of parenteral nutrition when acutely managing methylmalonic acidaemia and propionic acidaemia Georgina Wood ¹ , Dr Dexter Tarr ¹ , Dr Arunabha Ghosh ¹ ¹ Willink Biochemical Genetics Unit, St Mary's Hospital, Manchester University NHS Foundation Trust, UK	Clinical	Wednesday