Westminster Health Forum policy conference:

Priorities for rare disease diagnosis, care and treatment in England

Timing: Morning, Tuesday, 16th July 2024

Taking place online

Draft agenda subject to change



8.30 Registrati	on
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9.00 Chair's opening remarks

Senior Parliamentarian

9.05 Assessing the current landscape for the treatment of rare diseases - progress of the England Rare Diseases Action Plan 2024

Kath Bainbridge, Head of Rare Diseases and Emerging Therapies, Department of Health and Social Care Questions and comments from the floor

9.30 Progress and next steps for newborn screening for early detection and diagnosis of rare diseases

Senior representative, screening

9.40 Improving the delivery of care for patients with rare diseases

developing effective care pathways | patient accessibility to care | specialised commissioning | transitions from paediatric to adult care | coordinated care services | implementing a holistic approach to care | Early Access to Medicines Scheme

Dr Larissa Kerecuk, Rare Disease Lead and Consultant Paediatric Nephrologist, Birmingham Children's Hospital **Adam Galloway**, Senior Children's Physiotherapist, Leeds Children's Hospital, Leeds Teaching Hospitals NHS Trust

Senior representative, NHS

Senior representative, patient group Senior representative, holistic care

10.05 Questions and comments from the floor

10.30 Approaches to workforce education for rare diseases and therapies - raising awareness among healthcare professionals to

improve diagnosis and treatment

Dr Lucy McKay, CEO, Medics4RareDiseases Questions and comments from the floor

10.50 Chair's closing remarks

Senior Parliamentarian

10.55 Break

11.05 Chair's opening remarks

Senior Parliamentarian

11.10 Priorities for regulation and licencing pathways for innovative treatments

Senior representative, guidelines

Questions and comments from the floor

11.35 Advancing research and innovation across the rare disease pathway

exploring the future potential of genomic medicine | clinical trials | next steps for children's treatments through the Rare Therapies

Launch Pad | improving research eco-system collaboration and the role of the Rare Disease Research UK Platform | coordinating data
and disease registries | licencing and regulation pathways

Catriona Crombie, Head of Rare Disease, LifeArc

Senior representative, patient group Senior representative, research Senior representative, pharma Senior representative, NHS

Questions and comments from the floor

12.30 Next steps for utilising innovation in genomics for care and treatment of rare diseases

Ana Lisa Tavares, Clinical Lead for Rare Disease Research, Genomics England

Questions and comments from the floor

12.55 Chair's and Westminster Health Forum closing remarks

Senior Parliamentarian

Jessica Lear, Westminster Health Forum