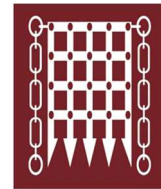


Westminster Health Forum policy conference

The future for UK regulation of medicines, medical devices and clinical trials – transparency and public trust, safety and enforcement, and support for innovation

Timing: Morning, Thursday, 16th July 2020

Taking Place Online



WESTMINSTER
HEALTH FORUM

Draft agenda subject to change

- 8.30 Registration
- 9.00 **Chair's opening remarks**
Daniel Zeichner MP
- 9.05 **Priorities for regulation and the role of the MHRA**
Jonathan Mogford, Policy Director, MHRA
Questions and comments from the floor
- 9.35 Break
- 9.40 **Meeting the challenge of regulating and developing medicines during the Covid-19 Pandemic**
Dr Aldo Faisal, Reader in Neurotechnology, Department of Bioengineering and Department of Computing, Imperial College London and Director, UKRI Centre in AI for Healthcare, Imperial College London
- 9.50 **'Transforming care: industry view on seizing opportunities to support access to innovation'**
Hilary Hutton-Squire, General Manager, UK and Ireland, Gilead Sciences
- 10.00 Questions and comments from the floor
- Stakeholder priorities for medicine regulation - safety, research integrity and innovation**
- 10.15 **Developing a regulatory ecosystem that supports innovation, provides regulatory certainty, and is transparent**
Erin Brooks, Head of Regulatory, Roche
- 10.25 **'Research transparency, public involvement and proportionality'**
Juliet Tizzard, Director of Policy, Health Research Authority
- 10.35 **Key issues for patient care - research, communication and access to treatment**
Neil Bennett, Director of Research, Action Duchenne
- 10.45 **Developing methods and technologies to ensure safe and effective medicines**
Dr Jan Turner, Director, Safer Medicines Trust
- 10.55 **Prioritising support for the uptake and adoption of innovations**
Patrick Stephenson, Director of Innovation and Healthcare, Fujitsu
- 11.05 Questions and comments from the floor
- 11.30 **Chairs closing remarks**
Daniel Zeichner MP
- 11.35 Break
- 11.40 **Chair's opening remarks**
Anne Marie Morris MP
- 11.55 **The future for collaboration with the EU, and the UK's relationship with the EMA**
Sarah Faircliffe, Legal Director, Bird & Bird
- 12.05 **'Using technology to tackle irresponsible medical ads - partnership with online platforms and statutory enforcement partners'**
Jane Eldridge, Head of Casework, Advertising Standards Authority
- 12.15 **Key issues for ensuring quality in clinical trials and the medicine approval process - research integrity, public trust and workforce development**
Eugene Pozniak, Managing Director, Siyemi Learning and Programme Director, European CME Forum
Phil Brown, Director of Technical and Regulatory, Association of British HealthTech Industries
Professor Pamela Kearns, Director, Institute of Cancer and Genomic Sciences and Director, Cancer Research UK Clinical Trials Unit, University of Birmingham
Questions and comments from the floor with **Sarah Faircliffe**, Legal Director, Blue & Bird and **Jane Eldridge**, Head of Casework, Advertising Standards Authority
- 12.55 **Chair's and Westminster Health Forum closing remarks**
Anne Marie Morris MP
Michael Ryan, Deputy Editor, Westminster Health Forum