

FROM THEORY TO PRACTICE:

Implementing the EU Health Technology Assessment Regulation



HYBRID MEETING – UTRECHT, THE NETHERLANDS, 30 JANUARY 2024

PROGRAMME

All times indicated in this programme are in CET.

09:00 – 09:15	<p>Welcome and introduction by Tamsin Rose, Moderator.</p> <p>Welcome by Sjaak Wijma, CEO of the National Health Care Institute, The Netherlands.</p> <p>Video message from Sandra Gallina, Director General of the Directorate General for Health and Food Safety, European Commission</p>
09:15 – 10:00	<p>Presentation and Q&A: “The EU regulation on health technology assessment: what’s in it and why it matters?”</p> <ul style="list-style-type: none"> • Roisin Adams, Chair of the Member State Coordination Group on HTA • Anne Willemsen, Co-chair of the subgroup for joint clinical assessments • Valentina Barbuto, Directorate-General for Health and Food Safety, European Commission
10:00 – 10:50	<p>Panel discussion: “The national perspective - Expectations, opportunities and the challenges ahead”</p> <ul style="list-style-type: none"> • Susanne Zöhrer, Ministry of Health, Austria • Marc Van De Castele, National Institute for Health and Disability Insurance, Belgium • Emer Fogarty, National Centre for Pharmacoeconomics, Ireland • Magali Boers, Ministry of Health, Luxembourg • Lonneke Timmers, National Health Care Institute, The Netherlands <p>Q&A and views from stakeholders</p>
10:50 – 11:10	<p>Coffee break</p>
11:10 – 11:55	<p>Panel discussion: “Ensuring engagement and cooperation in joint clinical assessments”</p> <ul style="list-style-type: none"> • Anna Nachtnebel, Austrian Social Insurance • Marijke de Vries, National Health Care Institute, The Netherlands • Robin Doeswijk, European Hematology Association • Derick Mitchell, IPPOSI - The Irish Platform for Patient Organisations, Science and Industry • Marjan Willaert, Pharma.be – Association of the Medicines Industry, Belgium <p>Q&A and views from stakeholders</p>

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11:55 – 12:40 Panel discussion: “Ensuring engagement and cooperation in joint scientific consultations”

- **Simon Roels**, National Institute for Health and Disability Insurance, Belgium
- **Dominique Hamerlijnck**, Dutch Lung Foundation, The Netherlands
- **Barbara Claus**, Ghent University Hospital, Belgium
- **Michael Berntgen**, European Medicines Agency

Q&A and views from stakeholders

12:40 – 12:45 Closing remarks by:

- **Tamsin Rose**, Moderator
 - **Roisin Adams**, Chair of the Member State Coordination Group on HTA
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