

Wednesday, May 21, 2025

## **EUFEMED Pre-conference workshop: The most suitable participant for phase 1 clinical trials**

13:00 - 13:15 Welcome and Introduction

13:15 - 15:15 The most suitable participant for phase 1 clinical trials - Plenary Session

How did the healthy volunteer evolve over the last 10 years?

Which populations should be included in phase 1 clinical trials?  
Sponsor, regulator and ethics committee perspectives.

What is the place of patients in First-In-Human trials?

15:15 - 15:45 Coffee break

15:45 - 16:45 Parallel breakout sessions on the above topics

16:45 - 17:00 Coffee break

17:00 - 18:00 Feedback from break-out sessions

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# Mastering increasing complexity in developing innovative therapies

21-23 May 2025



Together in Lamot Conference Centre  
Mechelen, Belgium



22 May 2025

Thursday, May 22, 2025

## EUFEMED-HEALIXIA Joint Conference day

9:00 - 9:15	Welcome and introductions – Jan de Hoon, EUFEMED & Erik Present, Healixia
9:15 - 9:45	Keynote Presentation - Genetic medicines: setting the scene for treatments of the future – Olivier Harari, Regeneron Genetic Medicines, USA
9:45 - 10:45	<p>Medicines Development for innovative therapies: challenges in early and late clinical development</p> <ul style="list-style-type: none"> <li>• Challenges faced by the sponsor: lessons learned and hurdles to overcome (Speaker invited)</li> <li>• Challenges faced by the investigator: focus on the treatment of ALS as a rare disease – Philip Van Damme, KU Leuven, Belgium</li> <li>• The patient's perspective as participant in a First-in-Human clinical trial (Speaker invited)</li> </ul>
10:45 - 11:15	Coffee break
11:15 - 12:45	<p>Medicines Development for innovative therapies - challenges in regulation and reimbursement</p> <ul style="list-style-type: none"> <li>• Joint Health Technology Assessment (HTA) in Europe: today and tomorrow? – Marc Van de Casteele, RIZIV-INAMI, Belgium</li> <li>• Perspective of industry (Speaker invited)</li> <li>• Perspective of regulators (Speaker invited)</li> </ul>
12:45 – 13:45	Lunch

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13:45 - 15:15 Oxford Debate - Diversity YES, but in Phase I? – With Keith Berelowitz, PatientRx Ltd, UK and Henri Caplain, Consultant, France

15:15 – 15:45 Coffee break

15:45 - 17:15 Parallel session A: The new ethical framework

- Impact of the new versions of the Declaration of Helsinki, ICH-GCP and WHO guidelines for our future performance of clinical trials – Barbara Baroutsou, IFAPP, Greece
- The VolREthics Initiative – What will it change? – François Bompart, INSERM Ethics Committee, France
- The eConsent Initiative – What It Is, What It Isn't, and Tools to Implement eConsent – Hilde Vanaken, TCS, EFGCP, Belgium

Parallel session B: Real World Data in Belgium: will Phase IV studies become obsolete?

- Secondary use of data as a source of Real World Data/Evidence – Annelies Verbiest, UZA, Belgium
- The Federated Health Innovation Network (FHIN): transform healthcare with data (Speaker invited)
- The We Are Platform: a sustainable civil-scientific ecosystem for personal health data – Elfi Goesaert, VITO, Belgium

17:30 - 18:00 Closing Keynote Presentation: Evolving a new framework to optimize drug development – Richard Hargreaves, Bristol Myers Squibb, USA

19:00 - 22:00 Social Event with dinner at Salons Van Dijck, Mechelen

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Friday, May 23, 2025

## EUFEMED Conference day 2

9:00 - 10:30	<p>Update on safety biomarkers in early clinical development</p> <ul style="list-style-type: none"> <li>• Interest and use of emerging safety biomarkers in the drug development – Philippe Detilleux, Sanofi, France</li> <li>• Kidney safety biomarkers in human and approach to interpret emerging exploratory biomarkers – Emmanuel Krupka &amp; Olivier Roux, Sanofi, France</li> <li>• CNS safety biomarkers in human and approach to interpret emerging exploratory biomarkers: the preclinical perspective – Greet Teuns, Johnson &amp; Johnson, Belgium</li> <li>• CNS safety biomarkers in human and approach to interpret emerging exploratory biomarkers: the clinical perspective – Geert Jan Groeneveld, CHDR, The Netherlands</li> </ul>
10:30 – 11:00	Coffee break
11:00 - 11:45	Research in the spotlight: Poster pitches of selected abstracts and Audience Voting for best presentation award
11:45 - 13:15	Artificial Intelligence in Clinical Development: Buzzword, vision, or reality? (Speakers invited)
13:15 - 13:30	Summary & End of conference – Ingrid Klingmann, EUFEMED
13:30 - 14:00	Farewell lunch

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