



EUROPEAN FEDERATION
FOR EXPLORATORY
MEDICINES DEVELOPMENT

EUFEMED Workshop

*"Making
the Investigator's Brochure
truly fit for purpose
in early medicines
development"*

September 20, 2024, Warsaw (Poland)

www.oeslive.pl/eufemed

Preliminary Programme

Status: June 2024

Friday, 20 September 2024

- from 8:00 Registration
- 09:00 Welcome and Introduction
Jan de Hoon, UZ Leuven, Belgium
- 09:10 What does the PI of an early phase clinical trial need from an Investigator's Brochure?
Jeroen van Smeden, CHDR, Netherlands
- 09:40 What does a Regulator authorizing an early phase clinical trial need from an Investigator's Brochure?
Sandrine Tinton, AFMPS, Belgium
- 10:10 What are the challenges of pre-clinical and translational experts in interpreting, risk-assessing and explaining the so far existing results?
Daniela Arndt, PCS, Switzerland
- 10:40 – 11:00 Coffee Break
- 11:00 – 12.15 *Break-out sessions I*
1. Preclinical Aspects: e.g., what data need to be included, how to present this data, risk benefit for first-in-human.
Stephanie Plassmann, PCS, Switzerland
Thijs van Iersel, ICON, Netherlands
 2. Clinical / life cycle IBs: e.g., update of IB during drug cycle, how to keep the IB understandable and 'short', updates of risk-benefit, substantial amendments.
Henri Caplain, France
Nariné Baririan, Chiesi, Italy
 3. Regulatory Aspects: e.g., what is needed for correct reviews, how to present data clearly, when to make amendments, input from different countries.
Sandrine Tinton, AFMPS, Belgium
Ingrid Klingmann, Pharmaplex, Belgium
Claudia Riedel, BfArM, Germany
 4. Investigator / end-user Aspects: e.g., needs for interpreting safety of participants, difference early / late phase investigators
Yves Donazzolo, Eurofins Optimed, France
Jeroen van Smeden, CHDR, The Netherlands
Ronald Koning, Biokinetica, Poland

12.15 – 13:00 Lunch Break

13:00 *Break-out sessions 2*

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4. Investigator / end-user Aspects: e.g., needs for interpreting safety of participants, difference early / late phase investigators

Yves Donazzolo, Eurofins Optimed, France
Jeroen van Smeden, CHDR, The Netherlands
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14:00 Reports with discussion from the Break-out Sessions

Moderator: Ingrid Klingmann, Pharmaplex, Belgium

15:30 – 15:45 Coffee Break

15:45 Discussion and decision on concluding recommendations for the early phase IB guideline

Moderator: Jelle Klein, SGS, Belgium

16:30 End of the Workshop

Workshop Date and Time

Friday, September 20th, 2024

9:00 – 16:30

Venue

University of Warsaw
Faculty of Applied Linguistics

Dobra 55

00-312 Warsaw, Poland

Organizing society

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Webpage and online Registration

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