

EMA/156670/2025

**Programme of the 2025 annual meeting of
the European network of paediatric research
at EMA (Enpr-EMA)**

Version 1.0

**Thursday,
20 November 2025**

European Medicines Agency (room 1C) and online

Background

The European Network of Paediatric Research at the European Medicines Agency (Enpr-EMA) is a network of research networks, investigators and centres with recognised expertise in performing clinical trials in children. Enpr-EMA's main objective is to facilitate clinical trials in order to increase the availability of authorised medicines for children.

Enpr-EMA's annual meeting is a unique opportunity for networks as well as other stakeholders of paediatric clinical trials, such as health care professionals, parents, carers, patient representatives as well as medicine developers and regulators, to learn about the network's priority activities. The meeting aims to facilitate inter-network and stakeholder collaboration and to build competences at European level.

Objectives of the event

The 2025 Enpr-EMA annual meeting aims to share regulatory updates, showcase network activities, and explore innovations in paediatric clinical trials. Key topics include the modernisation of Good Clinical Practice (GCP), implementation of the EU Clinical Trial Regulation, and updates from Enpr-EMA activities and its working groups. The meeting will also highlight advances in trial design, translational research, and the use of real-world data and artificial intelligence.

Programme

Chairpersons: Pirkko Lepola and Gunter Egger

Time	Topic	Topic lead	Duration
09:00	<i>Arrival and registration</i> <i>(Virtual meeting room open for remote participants)</i>		
09:30	Welcome address	<i>Ralph Bax</i>	5'
09:35	Opening remarks by Enpr-EMA Co-chairs	<i>Pirkko Lepola</i> <i>Gunter Egger</i>	5'
	<u>SESSION I - Accelerating data use</u>		60 min
09:40	<i>European Health Data Space (EHDS)</i> <i>Introduction to EHDS</i>	<i>Daniel Morales</i>	15'
09:55	<i>Discussion – Q&A</i>		5'
10:00	<i>EMA Reflection Paper on the use of Artificial Intelligence (AI) in the medicinal product lifecycle</i> <i>Leveraging data and artificial intelligence (AI) in regulating human medicines in practice</i>	<i>Gabriel Westman</i>	15'
10:15	<i>Discussion – Q&A</i>		5'
10:20	<i>Artificial Intelligence in paediatric research</i> <i>Data protection and ethics and use of data from minors for further research</i>	<i>Julian Isla</i>	15'
10:35	<i>Discussion – Q&A</i>		5'

Time	Topic	Topic lead	Duration
10:40 – 11:10	Coffee break		30'
	<u>SESSION II – Enhancing trial design and conduct</u>		55 min
11:10	Translational research from lab to patients Latest innovations by EPTRI (European Paediatric Translational Research Infrastructure)	Donato Bonifazi	15'
11:25	Discussion – Q&A		5'
11:30	Innovative trial designs in practice • Two case studies	Dieter Haering Marius Thomas Gina Calarco Smith	30'
12:00	Discussion – Q&A		5'
	<u>SESSION III - Enpr-EMA activities 2024/25 and new initiatives</u>		50 min
12:05	Annual report from the coordinating group • Enpr-EMA activities, achievements and challenges	Pirkko Lepola Isabel Sanchez	15'
12:20	Update from selected Enpr-EMA working groups • Paediatric research nurses • Paediatric trial sites quality criteria	Pamela Dicks Pernille Skovby Ricardo Fernandes	15'
12:35	New initiative – introductions • European Rare Disease Research Alliance (ERDERA)	Daria Julkowska	15'
12:50	Discussion – Q&A		5'
12:55-13:55	Lunch		60'
	<u>SESSION IV – Impact of ethics principles and public involvement</u>		40 min
13:55	Declaration of Helsinki (DoH) 2024 in paediatrics	Dominique Sprumont	15'
14:10	Discussion – Q&A		5'
14:15	Patient & parent experience of advocacy and involvement	Sebastian Honoré	15'
14:30	Discussion – Q&A		5'
	<u>SESSION V – Regulatory updates 1 – GCP</u>		35 min
14:35	Good Clinical Practice (GCP): Implementation in clinical trials ICH E6 (R3) - Annex I - history, reasons, overview	Peter Twomey	15'
14:50	ACT EU and GCP modernisation	Peter Twomey	15'

Time	Topic	Topic lead	Duration
	<i>ICH E6 (R3) - Annex II – pragmatic trials, decentralised trials, use of real-world data sources</i>		
15:05	<i>Discussion – Q&A</i>		5'
15:10 – 15:40	Coffee break		30'
	<u>SESSION VI – Regulatory updates 2 – EU Regulatory Network</u>		65 min
15:40	<i>Clinical Trial Assessment (CTA) update</i> <i>Paediatric clinical trial assessments under the new EU Clinical Trial Regulation – from PIP to CTA</i> <i>Feedback from ACT EU workshop on assessment on paediatric clinical trials/interpretation of CTR article 32</i>	<i>Laura Fankhauser</i> <i>Anette Solli Karlsen</i>	30'
16:10	<i>Implementation of ICH E11A guideline on paediatric extrapolation in Paediatric Investigation Plans (PIPs)</i>	<i>Dominik Karres</i>	15'
16:25	<i>Discussion – Q&A</i>		5'
16:30	AOB	All	5'
16:35	Conclusions	<i>Pirkko Lepola</i> <i>Gunter Egger</i>	10'
16:45	End of the meeting		

List of speakers / chairpersons:

Surname	Name	Affiliation
<i>Bax</i>	<i>Ralph</i>	<i>European Medicines Agency</i>
<i>Bonifazi</i>	<i>Donato</i>	<i>European Paediatric Translational Research Infrastructure (EPTRI)</i>
<i>Calarco Smith</i>	<i>Gina</i>	<i>ACRO - Association of Clinical Research Organizations, IQVIA</i>
<i>Dicks</i>	<i>Pamela</i>	<i>Scottish Children's Research Network</i>
<i>Egger</i>	<i>Gunter</i>	<i>European Medicines Agency, Co-chair of Enpr-EMA</i>
<i>Fankhauser</i>	<i>Laura</i>	<i>Hopp Children's Cancer Center Heidelberg (KiTZ), Germany</i>
<i>Fernandes</i>	<i>Ricardo</i>	<i>STAND4Kids - Supporting Paediatric Trials in Portugal</i>
<i>Haering</i>	<i>Dieter</i>	<i>Novartis</i>
<i>Honoré</i>	<i>Sebastian</i>	<i>Cure Lowe Foundation</i>
<i>Isla</i>	<i>Julian</i>	<i>Foundation 29</i>
<i>Julkowska</i>	<i>Daria</i>	<i>European Joint Programme on Rare Diseases</i>
<i>Karres</i>	<i>Dominik</i>	<i>European Medicines Agency</i>

Surname	Name	Affiliation
<i>Lepola</i>	<i>Pirkko</i>	<i>Finnish paediatric research network (FINPEDMED), Chair of Enpr-EMA</i>
<i>Morales</i>	<i>Daniel</i>	<i>European Medicines Agency</i>
<i>Sanchez</i>	<i>Isabel</i>	<i>European Medicines Agency</i>
<i>Skovby</i>	<i>Pernille</i>	<i>Danish paediatric medicine research network (DanPedMed), Paediatric Committee (PDCO)</i>
<i>Solli Karlsen</i>	<i>Anette</i>	<i>Federal Institute for Drugs and Medical Devices, Germany (BfArM), Paediatric Committee (PDCO)</i>
<i>Sprumont</i>	<i>Dominique</i>	<i>Research Ethics Committee of Vaud (Switzerland), European Network of Research Ethics Committees (EUREC)</i>
<i>Thomas</i>	<i>Marius</i>	<i>Novartis</i>
<i>Twomey</i>	<i>Peter</i>	<i>European Medicines Agency</i>
<i>Westman</i>	<i>Gabriel</i>	<i>Swedish Medical Products Agency (MPA)</i>