

EMA/202091/2026
Version 1.0

**Programme of the 2026 annual meeting
of the European network of paediatric research
at EMA (EnprEMA)**

Thursday, 5 November 2026

European Medicines Agency (2C) and online

Background

The European Network of Paediatric Research at the European Medicines Agency (EnprEMA) brings together research networks, investigators and centres with recognised expertise in conducting clinical trials in children. Its main objective is to facilitate high-quality paediatric clinical research and thereby support the development and availability of authorised medicines for children.

The EnprEMA annual meeting provides an opportunity for member networks and other stakeholders involved in paediatric clinical trials, including healthcare professionals, parents, carers, patient representatives, medicine developers and regulators, to exchange views on the network's priority activities. The meeting is intended to strengthen collaboration between networks and stakeholders and to support competence building at European level

Objectives of the event

The 2026 EnprEMA annual meeting aims to support exchange on priority areas for strengthening paediatric clinical research in Europe. The meeting will focus on advancing platform trial readiness, improving trial infrastructure and methodological approaches, strengthening patient engagement and recruitment facilitation, and discussing regulatory science developments relevant to paediatric drug development.

Programme

Chairpersons: Ricardo Fernandes 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		5 min
09:35	Opening remarks by EnprEMA Co-chairs	Ricardo Fernandes Gunter Egger	15 min
	<u>SESSION I – Supporting paediatric drug development through models, biomarkers</u>		
09:50	Disease and progression models to support paediatric drug development	Cecile Ollivier	20 min
10:10	Framework for model evaluation and qualification of biomarkers to support paediatric drug development	Efthymios Manolis	20 min
10:30	Discussion – Q&A	Sara Vennberg	30 min
11:00	Coffee break		20 min
	<u>SESSION II – Platform trials: building awareness, capacity and evidence</u>		
11:20	Optimising paediatric academic-industry platform trials: successes, lessons learned and needs	Pamela Kearns	20 min

Time	Topic	Topic lead	Duration
11:40	Platform trials to inform paediatric drug development: an industry perspective	Angeliki Siapkara	20 min
12:00	Discussion – Q&A	Dominik Karres	30 min
	RECENT DEVELOPMENTS		
12:30	Update on the assessor's guidance on best practices for paediatric clinical trials	Monique Al Anette Solli Karlsen	20 min
12:50	Lunch		60 min
	<u>SESSION III – Advancing patient partnership and patient experience in paediatric medicines development: from engagement to evidence</u>		
13:50	Current capability: insights from recent PPI surveys	Segolene Gaillard Maria Mavris	20 min
14:10	Developments regarding Patient Panels	Laura Pioppo	20 min
14:30	Discussion – Q&A	Tomasz Grybek	25 min
14:55	Coffee break		20 min
	RECENT DEVELOPMENTS		
15:15	Reporting guidelines and paediatric drug development: SPIRIT-C & CONSORT-C	Martin Offringa	20 min
15:35	Preparedness for public health emergencies (PHE)	Laura Fregonese	20 min
15:55	First-in-child/first-in-human workshop – reporting back	Gilles Vassal	20 min
16:15	AOB and conclusions	Ricardo Fernandes Gunter Egger	15 min
16:30	End of the meeting		

List of speakers / chairpersons:

Surname	Name	Affiliation
Al	Monique	CCMO - Central Committee on Research Involving Human Subjects, The Netherlands, CTCG – Clinical Trials Coordination Group, MedEthicsEU

Surname	Name	Affiliation
<i>Egger</i>	<i>Gunter</i>	<i>European Medicines Agency, Co-chair of EnprEMA</i>
<i>Fernandes</i>	<i>Ricardo</i>	<i>STAND4Kids - Supporting Paediatric Trials in Portugal, Chair of EnprEMA</i>
<i>Fregonese</i>	<i>Laura</i>	<i>European Medicines Agency</i>
<i>Gaillard</i>	<i>Segolene</i>	<i>French paediatric research network (Kids France)</i>
<i>Grybek</i>	<i>Tomasz</i>	<i>Paediatric Committee (PDCO), EURORDIS – European Organisation for Rare Diseases,</i>
<i>Karres</i>	<i>Dominik</i>	<i>European Medicines Agency</i>
<i>Kearns</i>	<i>Pamela</i>	<i>ITCC - Innovative Therapies for Children with Cancer</i>
<i>Manolis</i>	<i>Efthymios</i>	<i>European Medicines Agency</i>
<i>Mavris</i>	<i>Maria</i>	<i>European Medicines Agency</i>
<i>Offringa</i>	<i>Martin</i>	<i>The Hospital for Sick Children, Canada</i>
<i>Ollivier</i>	<i>Cecile</i>	<i>Critical Path Institute (C-Path)</i>
<i>Pioppo</i>	<i>Laura</i>	<i>European Medicines Agency</i>
<i>Siapkara</i>	<i>Angeliki</i>	<i>EFPIA - European Federation of pharmaceutical Industries and Associations, AstraZeneca</i>
<i>Solli Karlsen</i>	<i>Anette</i>	<i>Paediatric Committee (PDCO), Norwegian Medical Products Agency</i>
<i>Vassal</i>	<i>Gilles</i>	<i>ITCC - Innovative Therapies for Children with Cancer</i>
<i>Vennberg</i>	<i>Sara</i>	<i>Paediatric Committee (PDCO), Swedish Medical Products Agency</i>