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## PROGRAMME

## EU-X-CT PUBLIC STAKEHOLDERS' FORUM MAKING CROSS-BORDER ACCESS TO CLINICAL TRIALS A REALITY

12 April 2024

10:00 – 17:00 CEST

Crowne Plaza Brussels Airport Da Vincilaan 4 1831 Diegem

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WEBSITE

EU-X-CT C/O EFGCP EU-X-CT Page

# INTRODUCTION

The pan-European public-private EU-X-CT Initiative aims at facilitating cross-border access to clinical trials in concrete terms. We are collecting information on the regulatory, ethical, social security and organizational conditions for the involved stakeholders, experiences and best practices in all European countries. This Public Stakeholder Forum will be the opportunity to jointly review the national conditions for patients, clinical investigators, academia/public funders and industry sponsors in need for cross-border participation of patients in clinical trials. The EU-X-CT gap analysis will be the basis for our panel and open forum discussions to come to conclusions about most efficient European and national strategies and activities for improvements in the near and mid-term future.

	PROGRAMME COMMITTEE	
Bettina Ryll	Stockholm School of Economics Institute for Research	Sweden
Emilie Prazakova	Roche	Czech Republic
Ingrid Klingmann	EFGCP	Belgium
Lisbeth Oxholm Snede	Patient advocate	Denmark
Maja Pizevska	Berlin Institute of Health	Germany
Marén Koban	Merck Healthcare KGaA	Germany
Sabine Kläger	ECRIN	Germany
Solange Corriol-Rohou	AstraZeneca	France
Teodora Lalova-Spinks	KU Leuven	Belgium

## ACKNOWLEDGEMENT

#### THE EU-X-CT THANKS ITS CONTRIBUTORS FOR THEIR CONTINUED SUPPORT

**INITIATIVE LEADERSHIP** 





#### 12 APRIL, 2024 - CEST SCHEDULE

09:00	Registrations & Welcome Networking Coffee	
	SESSION 1: WHERE ARE WE TODAY?	
10:00	Introduction to EU-X-CT	
	Welcome from the industry members	
	Welcome from the academia members Welcome from the patient members	
10.20	·	
10:20	Round table: Patient experience Moderator(s):	
	Lisbeth Oxholm Snede, Founder of Patients Unite	
10:50	Keynote: Overcoming borders will make Europe a more attractive place for clinical trials	
	EU Commission DG R&I	
	Discussion	
11:20	The issues of cross-border access to clinical trials in Europe: The EU-X-CT gap analysis	
	Moderator(s):	
	Susan Bhatti, Merck Group	
12:20	Panel and Open Forum Discussion:	
	<ul><li>Is this picture accurate and comprehensive?</li><li>Where are the assessment gaps?</li></ul>	
	• How to close them?	
	<ul> <li>Which other EU partnerships suffer from the lack of cross-border access to clinical trials?</li> <li>Moderator(s):</li> </ul>	
	Solange Corriol-Rohou, AstraZeneca	
13:00	Lunch	
	SESSION 2: WHERE DO WE WANT TO GO?	
14:00	European needs in overcoming borders in clinical research	
	Cross-border access in precision medicine trials: PCM4EU and PRIME-ROSE	
	Moderator(s): Sabine Kläger, ECRIN	
14.20	The way forward: How do we make progress in cross-border access to Clinical Trials?	
14:30	Moderator(s):	
	Bettina Ryll, Stockholm School of Economics Institute for Research	
	Panel and Open Forum Discussion	
15:30	Break	
15:50	Open Forum Discussion: How can EU-X-CT achieve the most urgent goals and how to	
	make them sustainable?	
	Moderator(s):	
	Ingrid Klingmann, EFGCP	
16:50	Conclusions and next steps	
	Ingrid Klingmann, EFGCP	
17:00	End of meeting	