CRITICAL PATH INSTITUTE

View Now | C-Path in Europe: Moving Global Regulatory Science Forward

In support of C-Path's mission to catalyze innovation that accelerates the path to a healthier world, we're excited to invite you to this webinar to learn more about how C-Path's new headquarters in Amsterdam will continue to improve public health, share expertise, data, risks and costs to move global regulatory science forward by facilitating public-private partnerships with members from the biopharmaceutical industry, government regulatory agencies, academic institutions, and patient groups in Europe.

Speakers include C-Path scientific leadership who will present on-going activities on key topics, including:

- Digital Technologies
- Real-World Evidence
- Model Informed Drug Development
- Complex Clinical Trials

Following presentations, audience questions will be answered by a panel of representatives from EFPIA, academia, EMA and patient groups.

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Topic	**Speaker**
Welcome and Introduction	Kristen Swingle Tomas Salmonson
C-Path global vision and mission	Klaus Romero
EMA Regulatory Science Strategy (RSS) & How C-Path Can Contribute	Cecile Ollivier
Drug Development Tools	Klaus Romero

Complex Clinical Trials and Digital Technologies	Terina Martinez Martijn Muller
Patient-Centric Drug Development and Real-world Evidence	Scottie Kern Jeff Barrett
Repurposed Drugs	Marco Schito
Collaborations and Training in Regulatory Science	Jeff Barrett Huong Huynh
Q&A for C-Path Speakers	C-Path Speakers
Panel Discussion (industry, academia, regulators, patient groups)	Lada Leyens Ralf Herold Dimitrios Athanasiou Franz Koenig
Closing remarks	Cecile Ollivier

C-Path Speakers

- Kristen Swingle: Interim President & Chief Operating Officer (US)
- Tomas Salmonson: Member, Board of Directors (US) & Member, Board of Directors (Netherlands)
- Klaus Romero: Chief Science Officer & Executive Director of Clinical Pharmacology (US)
- Cecile Ollivier: EU Regulatory Science Advisor (US) & Member, Board of Directors (Netherlands)
- Terina Martinez: Executive Director Duchenne Regulatory Science Consortium & Critical Path to Therapeutics for the Ataxias (US)
- Martijn Muller: Senior Scientific Director, Critical Path for Parkinson's Consortium (US)
- Scottie Kern: Executive Director, Electronic Clinical Outcome Assessment Consortium (US)
- Jeff Barrett: Senior Vice President; Rare Diseases Cures Accelerator Data and Analytics Platform (RDCA-DAP) Lead (US)
- Huong Huynh: Director, Regulatory Science (US)

Panelists

- Lada Leyens: Global Regulatory Lead, Clinical Trial Innovation & Digital Health at Roche
- Ralf Herold: Senior Scientific Officer, Taskforce Regulatory Science & Innovation at EMA
- Dimitrios Athanasiou: PDCO Member at EMA – EURORDIS Board, Member in World Duchenne Organization, European Patients Forum Greek Patient Association
- Franz Koenig: Associate Professor, Medical University of Vienna

To learn more about C-Path in Europe, visit https://c-path.org/c-path-in-europe/ and read C-Path European Nonprofit Established in Amsterdam.

FDA acknowledgement