RBQM LIVE24

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CluePoints

SEPTEMBER 24–25

What to Expect at RBQMLIVE 2024

For clinical trial professionals, it's vital to expedite the process of bringing effective medicines, therapies, and devices to market. Achieving this demands collaboration among the industry's brightest minds to uncover new insights and advance innovative ideas.
So, we invite you to attend RBQMLive 2024!

WHAT IS RBQMLIVE?

As the industry-leading event of the year, RBQMLive brings industry experts, clinical trial specialists, and curious listeners together to share insights and discuss the acceleration of RBQM. Join virtually to hear the transformative shifts in RBQM technology, global regulations, AI, and more.

HOW TO ACCESS THE CONTENT

RBQMLive is a **free, virtual event.** Once you register, you'll receive a link to log in from your computer and access all the sessions—whether you're in the conference room or relaxing on your couch.

After the event, we'll send you a new link to **access the content on demand.** This way, you can catch up on any sessions you missed or re-watch your favorites at your convenience.

HIGHLIGHTS FROM LAST YEAR

1. Explored evidence-based insights on the evolution of RBQM maturity
2. Discussed opportunities and challenges with evolving guidance updates
3. Analyzed real-world applications of artificial intelligence (AI) and machine learning (ML) in RBQM
4. Saw how RBDM identifies real-time "outliers" and enables prompt corrective action
5. Examined the role of audit trail review in detecting risks and improving data quality
6. Gained insights into effective collaboration strategies for study oversight

Check out the full 2024 agenda on the next page for a comprehensive list of the exciting sessions and content we're offering during this year's sessions.

RBQMLIVE 2024 AGENDA

SEPTEMBER 24th | DAY 1 SESSIONS

Keynote: Unlocking the Path to RBQM Maturity | 8:30 – 9:30 am*
Speaker: Ken Getz, Executive Director & Professor, Tufts Center for the Study of Drug Development

Panel: Taking RBQM into New Dimensions – Understanding the Risk-Based Data Management (RBDM) Revolution | 9:40 – 10:50 am*

Speakers: Miguel Valenzuela, Associate Director of Clinical Operations, RBQM, Alnylam

Paul MacDonald, Senior Director Vault CDMS Strategy at VeevaPanel

Patrick Nadonly, Global Head, Clinical Data Management, Sanofi

Panel: The Emergence of AI & ML in RBQM: What's here now and what is to come? | 11:00 – 12:00 pm*

Speakers: Marcin Makowski, Head of Centralized Monitoring & Data Analytics, GSK

Bob Zambon, Vice President, Technology Strategy & Strategic Partnerships, Syneos Health

SEPTEMBER 25th | DAY 2 SESSIONS

Panel: RBQM Industry Guidance Evolution – Adapting to Modernization | 8:30 – 9:30 am*

Speakers: Olivia Feiro, Director, Risk Management & Analytics, CSL Behring

Lotte Smets, Director Data Management, Julius Clinical

Daniel DiJohnson, Risk-Based Quality Management SME, Senior Manager, Parexel

Enhancing Data Quality through Advanced Audit Trail Review | 9:40 – 10:50 am*

Speakers: Paola Peshkepija, Senior Associate Central Monitor, Pfizer

Jennifer Nielsen, Senior Specialist, Clinical Data Management, Lundbeck

Panel: Integrated Data Review: Optimizing Study Oversight | 11:00 – 12:00 pm*

Speakers: Brian Barnes, Director of Risk Management Strategy – Process Excellence & Oversight, BioNTech

Jess Thompson, Founder, Clinical Research Project Management Association

Alaine Heffernan, Director Centralized Monitoring, Allucent

*All times are in Eastern Standard Time (EST)

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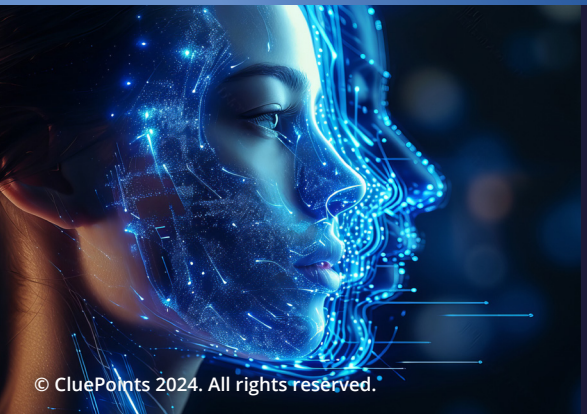
Syneos Health · BIONTECH · Julius Clinical · CS · Alnylam · Allucent · CRPM



THE FUTURE OF RBQM AWAITS

WILL YOU BE THERE?

From clinical trial specialists to thought leaders and curious listeners, we welcome everyone to join our industry-leading RBQM event and connect with CluePoints, the premier event sponsor.

Register today to reserve your spot.