PROVENTA INTERNATIONAL

IQVIA TECHNOLOGIES

VIRTUAL BOARDROOM

Preventing quality findings using an RBQM system

Thursday, 1st December 2022
10:00am EST

Discussion

- Current regulations and guidelines that support an RBQM approach to clinical trials
- Unpacking common audit findings in clinical trials
- Ensuring regulatory compliance and audit readiness using RBQM technology

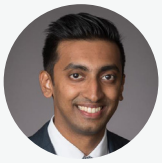
Speakers:

Gayle Hamilton — *Director, RBQM, Digital Trial Management Suite,* IQVIA Technologies

Gayle Hamilton is an experienced Risk-Based Monitoring project advisor and project lead with a strong background in clinical operations and project management. She has supported study trial teams and IT development in RBQM implementation across all phases and therapeutic areas, driving the development of processes, tools, and systems. Gayle is also experienced in assuring high-quality business performance of clinical operations within global projects.

Adrian Kizewski — *Associate Director, RBQM, Digital Trial Management Suite,* IQVIA Technologies

Adrian brings expertise spanning R&D and clinical life sciences, business analysis, process design & improvement, and product implementation. He is currently a lead for IQVIA's Digital Trial Management Suite focused on the Risk Based Quality Management SaaS solution. Adrian holds an MBA from the McDonough School of Business at Georgetown University in addition to an MSc in Pharmacology from The Johns Hopkins University School of Medicine and a BSc in Biochemistry from Temple University.



Sathya Ramnath — *Product Lead, RBQM, Digital Trial Management Suite,* IQVIA Technologies

Sathya is a Product Lead at IQVIA Technologies, where he drives business and product vision, strategy, and execution of Advanced Analytics for IQVIA's Risk Based Quality Management SaaS solution. He brings expertise spanning R&D, clinical operations, process design, artificial intelligence, and product implementation. Sathya holds a Bachelor's degree in Pharmaceutical Sciences from the Institute of Chemical Technology, India and an MBA from University of California San Diego.

Rachael Geedey — *Associate Director, RBQM, Digital Trial Management Suite ,* IQVIA Technologies

Rachael brings expertise in technology training, adoption and implementation across various clinical trial platforms. She is passionate about people, process and technology and has extensive operational experience supporting study teams in the use of RBQM technology. Rachael is currently an Associate Director supporting IQVIA's Risk Based Quality Management SaaS Solution.

About IQVIA Technologies:

IQVIA is a leading global provider of advanced analytics, technology solutions, and clinical research services to the life sciences industry. IQVIA creates intelligent connections to deliver powerful insights with speed and agility—enabling customers to accelerate the clinical development and commercialization of innovative medical treatments that improve healthcare outcomes for patients. With approximately 82,000 employees, IQVIA conducts operations in more than 100 countries. You can learn more at www.iqvia.com.

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