



June '26

Insights from the Cutting Edge of MedTech

This month, we're exploring what it takes to navigate complexity in today's medtech landscape, from the critical role of ethnographic research in shaping better solutions to the evolving regulatory and operational demands facing teams across the product lifecycle. Inside, you'll find insights on strengthening your preclinical approach, closing compliance gaps, and making smarter decisions, along with the latest updates and opportunities to connect with experts in the field.

This Month's Service Spotlights

Preclinical Execution Built Around the Result You Need

"Our job is not just to run the study. It is to partner with clients to get the right outputs, with the quality, collaboration, and problem-solving needed to move their program forward."

Michael Sweet

Director, Preclinical Study Management

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Design Instructions Users Actually Follow

Great devices fail when instructions are unclear in real workflows. We craft packaging artwork, iconography, and instructional content, including IFUs and QRGs, to strengthen comprehension and usability for submission and launch.

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Learning Opportunities



Valentina Lintas
Quality & Regulatory Affairs Managers

Corrective & Preventive Actions (CAPA) Training
June 10th, 2026

[Register Now](#)



Bruno Strappa
Quality & Regulatory Affairs Manager



Cédric Razaname
Director, Quality & Regulatory Affairs, Head of QMS

IEC 62304 Software Lifecycle Training
June 18th, 2026

[Register Now](#)

Webinar:

Prepare for FDA's Evolving Inspection Approach

July 30th, 2026

[Register Now](#)



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Veranex's Biocompatibility Insights 2026

Explore the Science Behind the Regulations: Emerging Trends and Clinical Realities

Copenhagen, Denmark
Exclusive Training Day: October 12
Full Conference: October 13-14

[Register Now](#)



CRF®
NEW YORK VALVES

The Structural Heart Conference

June 24, 2026 - June 26, 2026
New York, NY



Industry Insights From Our Experts



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What MedTech Investors Will Demand in 2026 and Beyond

If you can't walk someone through your integrated evidence plan in 90 seconds, the meeting ends there.



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Europe Is Rewriting the Rules on Medical Devices

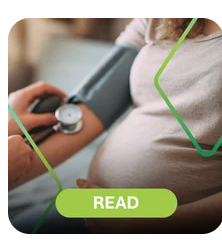
For startups, mid-size manufacturers or "strategics" these changes will reshape your compliance roadmap.



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Missing Cybersecurity Artifacts Can Kill Your Submission

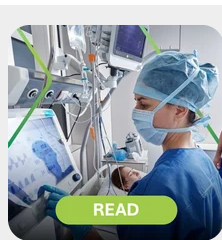
What software & engineering teams need to plan, document, and prove before you get to the finish line. Start by keeping a "Zero Trust" mindset.



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Market Access: Evaluating Your Right to Play in Women's Health

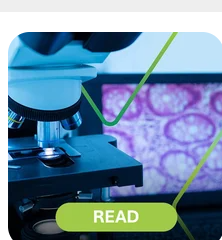
Why the blind spot between personal experience and population reality is where well-intentioned solutions go to die.



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FDA Opened a New Door on Real-World Evidence. Don't Walk Through It Unprepared

Three regulatory pathways are now more accessible. IF your evidence strategy is designed to take advantage of them.



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Six Signs You're Not Ready for GLP: The Histology Perspective

Histopathology requirements frequently get treated as presumptive afterthoughts, until tissue processing reveals that your study wasn't ready to begin.

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