



Sept '26

Insights from the Cutting Edge of Medtech

As Medtech evolves, staying ahead means keeping pace with industry changes. This month, explore Veranex expert insights on preclinical studies, FDA pre-submissions, EU medical device regulations, and cybersecurity. Plus, discover upcoming training and events to keep you informed and connected.

ANNOUNCEMENTS AND SPOTLIGHTS

Evaluate Your Preclinical Partner Through a First-in-Human Lens

A GLP study can satisfy a requirement and still leave clinical questions unanswered. Use this framework to pressure-test whether your preclinical plan is built to support regulatory readiness, investigator confidence, site preparedness, and downstream evidence.

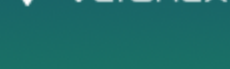
[Get the Framework](#)


The Impact of 24/7 ICU Monitoring on Research Success

In the realm of veterinary research, the value of continuous 24/7 ICU monitoring cannot be overstated. Jessica Amilcar, Director of Preclinical Veterinary Research at Veranex, sheds light on how this practice enhances study integrity and outcomes.

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E-LEARNINGS




Webinar

Clinical Evaluation under MDR Article 61(10)

Practical Application of the New Team-NB Position Paper

Dr. Ulrich Nitsche, M.D./Ph.D.
Global Director, Centre of Clinical Evaluation, TÜV SÜD



Julianne Bobela, Ph.D.
Scientific Affairs Director, Veranex



September 24, 2026

9:00-10:00 AM ET

15:00-16:00 CEST

[Register Now](#)

Advance your regulatory and quality expertise with upcoming Veranex trainings. This fall, explore expert-led courses with practical guidance you can apply to your work.

Upcoming Veranex Trainings Fall 2026

- MDSAP
- ISO 13485
- EU MDR
- Regulatory Affairs

[Explore Upcoming Trainings](#)


EVENTS: MEET US IN PERSON



Biocompatibility Insights 2026

Explore the Science Behind the Regulations

Copenhagen, Denmark

Exclusive Training Day: October 12

Full Conference: October 13 – 14

[Register Now](#)


LSI Europe

September 28 – October 2

Barcelona, Spain

Connect with our team in our private meeting room.



The Medtech Conference

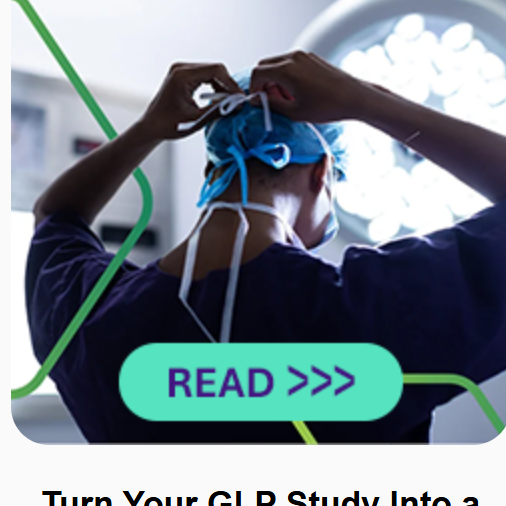
October 18 – 21

Boston, MA

Visit us at Booth #1106.

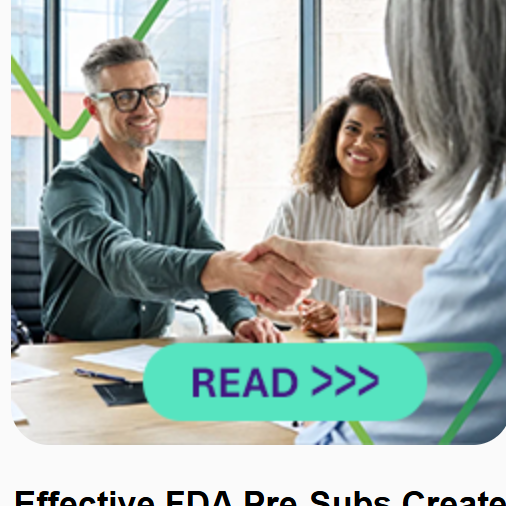
[Meet with Veranex](#)

INDUSTRY INSIGHTS FROM OUR EXPERTS


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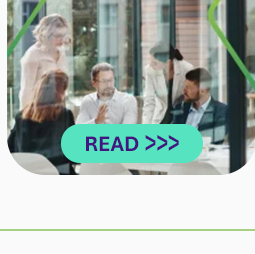
Turn Your GLP Study Into a Stronger Clinical Start

Preclinical strategy should do more than support a submission. Generate insights that strengthen clinical execution and market readiness.


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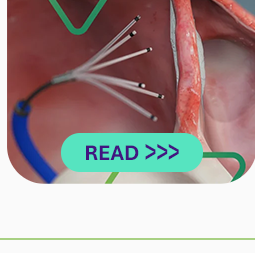
Effective FDA Pre-Subs Create Stronger Development Decisions

The most valuable FDA conversations start before the meeting. Explore three essentials for bringing questions that unlock useful, actionable feedback.


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Orphan and Breakthrough Devices Have a New EU Pathway

A closer look at emerging EU options that could help high-need device innovators build momentum with more tailored evidence expectations.


[READ >>>](#)

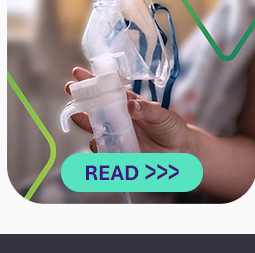
Preclinical Precision Cleared Recross Cardio for First-in-Human

See how carefully chosen models and clean GLP execution helped turn a novel PFO concept into an FDA-cleared early feasibility study.


[READ >>>](#)

Build FDA Cybersecurity Into Your Device From Day One

Discover how secure-by-design can turn cybersecurity from a late-stage requirement into a product advantage from the start.


[READ >>>](#)

Design for Adoption Around How Care Is Actually Delivered

The most revealing product insights may be waiting outside the lab. See how use-environment analysis brings real-world workflow into focus.

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