



Aug '26

Insights from the Cutting Edge of Medtech

From human factors and evidence planning to regulatory readiness and clinical strategy, the right expertise can change how quickly teams see risk and act on it. Explore the latest insights, trainings, and events from across Veranex.

ANNOUNCEMENTS AND SPOTLIGHTS

With Director of Clinical Affairs

KAREN MINOR

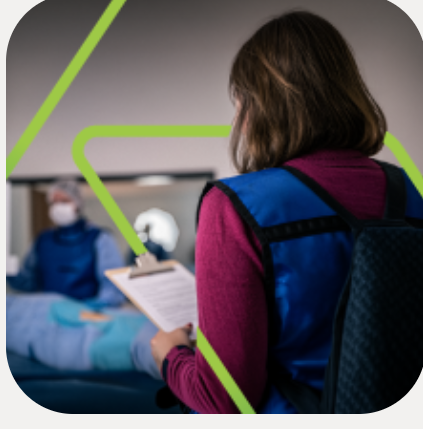
Episode 3 | IVD Study Insights

Insights on the unique aspects of IVD clinical studies, the importance of specimen strategy, and how evolving technology like AI impacts study design.



Connect Human Factors to Every Stage of Development

From early user research and formative testing to labeling, validation, and production readiness, human factors help keep the user at the center of device development. Our integrated PDE teams bring that perspective into the decisions that shape product success.

[Explore More](#)


Use the Evidence You're Already Generating More Strategically

Integrated evidence planning helps align studies, endpoints, and decisions already in motion so evidence can support regulatory, payer, clinical, and commercial needs.

[Learn More](#)


E - LEARNINGS

The Next Era of FDA Inspections: Digital, Data-Driven, and QMSR-Ready



Instructor

Paige Sutton-Smith

Senior Quality & Regulatory Affairs Manager, Veranex

Date: August 18

Training of Internal Auditors for MDSAP Country-Specific Requirements



Instructor

Cedric Razaname

Director, Quality and Regulatory Affairs, Veranex

Date: October 5

[Register Now](#)

EVENTS: MEET US IN PERSON



August 26 - 27

Boston, MA



September 28 - October 1

Barcelona, Spain

[Learn More](#)

INDUSTRY INSIGHTS FROM OUR EXPERTS



Veranex Client Atraverse Makes Successful Strategic Exit

We surfaced a workflow gap nobody had spec'd for. Three years later, J&J acquired the system.



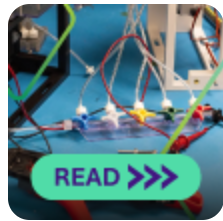
The Medical Device Workshop Readiness Checklist

A failed workshop doesn't just waste a day. It derails entire programs. Run this checklist first.



Closing the Imaging Gap in Preclinical Neurosurgery

MRI goes dark once the craniotomy starts. Intraoperative ultrasound catches brain shift while surgeons can still correct it.



Legacy Medical Devices: Refresh Insights for Your Next Generation

What got your Gen 1 device to market won't get you through Gen 2. Three regulatory shifts catching teams off guard, and where the budget should move instead.



What the Proposed EU MDR Revision Means for You

The EU's proposed MDR/IVDR overhaul touches certification timelines, clinical evidence, and innovation pathways. Catch up if you missed it.



How to Avoid FDA Warning Letters

Warning letters are public, permanent, and preventable. Catch up on the prevention and response guide if you missed it.

[Product Design & Engineering](#)

[Preclinical & Pathology](#) | [Clinical Research \(CRO\)](#)

[Regulatory Quality](#) | [Commercialization](#)



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