

OCTOBER '26

Insights from the Cutting Edge of Medtech



This month, we're taking you from the lab to the regulatory desk. Get a look inside our Paris cardiovascular lab, then hear from Veranex experts on SaMD, the RAPID pathway, reshoring, women's health, tariffs, and GLP study planning. Plus, discover upcoming training and events to keep you informed and connected.

CARDIOVASCULAR INNOVATION

STARTS HERE

Advancing Structural Heart Technologies in Paris

Success starts with the right model. For over 20 years, our veterinarian-led team in Paris has designed cardiovascular studies around how species differ from human anatomy, with bypass, imaging, and pathology in one environment. From proof of concept to GLP safety, one expert team supports every stage, advancing products that have treated 1 million+ patients.

Learn more about our cardiovascular expertise and meet our experts in Paris. In-person and remote sessions are available.

Our Cardiovascular Expertise



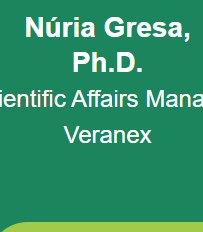
 TRAINING

Clinical Evaluation Under MDR

How to Define and Justify the Appropriate Level of Clinical Evidence

 October 26, 2026

 9:30 AM – 12 PM EST | 3:30 PM – 6 PM CET



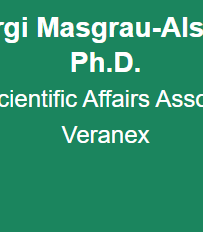
Núria Gresa, Ph.D.

Scientific Affairs Manager, Veranex



Laia Morató, Ph.D.

Sr. Scientific Affairs Manager, Veranex



Sergi Masgrau-Alsina, Ph.D.

Sr. Scientific Affairs Associate, Veranex

Register Now

 TRAINING

ISO 13485 Quality Specialist Training

Build foundational quality management knowledge for the medical device industry and prepare for TÜV SÜD Academy certification.

 October 26 – November 6, 2026

 8 AM – 12:30 PM EST | 2 PM – 6:30 PM UTC +1

Exam Date: November 17, 2026

Register Now

 THE MEDTECH CONFERENCE
OCT 18-21 | BOSTON, MA

Hosted by  AdvaMed
The Medtech Association

Join Thousands of Medtech Leaders at the Medtech Event of the Year

Booth #1106

 CRF®
TCT OCTOBER 31 – NOVEMBER 3, 2026
SAN DIEGO CONVENTION CENTER
SAN DIEGO, CA

Where Pioneers Unite to Shape the Future of Interventional Cardiology

 October 31 – November 3, 2026

 San Diego, CA

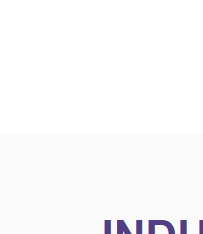
MedTech Strategist
INNOVATION SUMMIT
San Diego, CA

Pre-Conference Workshop:
Designing for Approval and Adoption:
Aligning Evidence, Regulation, and
Market Access Early

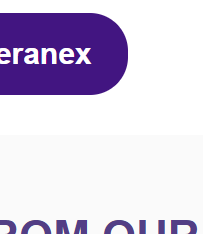
 November 3 – 5, 2026

 10:45 – 11:45 AM PST

 San Diego, CA



Joseph Ferrara
Chief Strategy Officer,
Veranex



Pablo Morales
Chief Medical Officer,
Veranex

Meet with Veranex

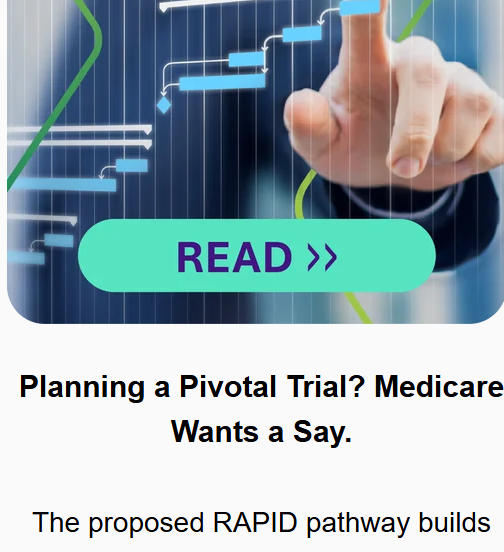
INDUSTRY INSIGHTS FROM OUR EXPERTS



READ >>

Five Questions to Help You Determine If Your Software Is a Medical Device

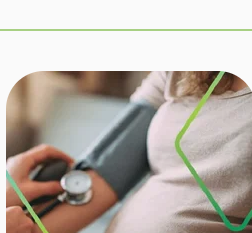
Understanding where your software falls on the spectrum is typically the first regulatory decision you'll make for the product.



READ >>

Planning a Pivotal Trial? Medicare Wants a Say.

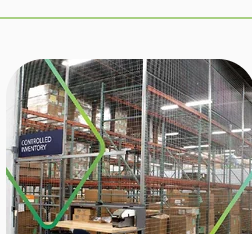
The proposed RAPID pathway builds CMS coverage outcomes into IDE studies for Breakthrough Devices. Here's what it means for your trial design.



READ >>

Reshoring a Medical Device? Make the Move Count Twice.

If your medical device is showing its age or if complaints, cost pressure, supplier obsolescence, or a competitor closing in are forcing the issue, the reshoring window is the right time to address Gen 2.



READ >>

It's Not What You Think: Evaluating Your Right to Play in Women's Health

Decades of male-centric data hide real risk in women's health. Learn the red flags and what market readiness looks like.



READ >>

Tariff-Proof Your Device, Starting With Materials

Early material and component choices decide whether tariffs and supplier changes become setbacks or non-events.



READ >>

Two Hours on Surgery. Fifteen Minutes on Pathology.

Imbalances in study planning are behind three costly GLP failures, and each one is preventable.

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