Transform Your Research Operations

Prepare your cancer center for the future of clinical trials.

Overview

Veeva provides a suite of applications built specifically for research sites, cancer centers, and institutions with connectivity to sponsors and patients to reduce complexity and advance research.

Benefits

- Improve visibility into start-up and study activation
- Enhance quality and compliance
- Connect to 350+ industry sponsors and CROs
- Reduce patient burden and improve satisfaction

Products

Go paperless and enable remote monitoring today with Veeva SiteVault

Reduce administrative burden with a modern eRegulatory system. Manage documents in a system that is 21 CFR Part 11 and HIPAA compliant. Design your own workflows to support quality and speed study activation. Create your own reports and dashboards to see what's completed and what's missing. Enable remote monitoring for regulatory and source files. Veeva SiteVault simplifies regulatory compliance so you can get more done. Learn more >

Seamlessly exchange information with sponsors with Vault Site Connect

Vault Site Connect enables information to exchange seamlessly between sites and sponsors, so you can focus less on administrative tasks and more on treating patients. Streamline the delivery and receipt of IND safety reports, feasibility surveys, start up packages, and study documents. There's no additional cost, no integrations to manage, and no effort to get started. Learn More >

Make patient participation easier with Veeva eConsent

Transform the current paper-based consent process with Veeva eConsent, a free solution for sites and patients. Tailor sponsor-provided consents to the requirements of their ethics boards and deliver forms to your patient's mobile device for their consent. Learn more >