



High Quality Applications Built for Clinical Research

Veeva SiteVault improves efficiency, quality, and collaboration for research sites with unified **CTMS, eISF and eSource**. With Veeva SiteVault, sites seamlessly integrate with sponsors to streamline information sharing and speed trials.

"We need to let go of what we did before. We don't want technology to mirror the paper processes we previously had."

Meghan Blair, Abramson Cancer Center at Penn Medicine, Director of Regulatory Affairs

Connecting the Industry

9,300+ Sites

eISF + eConsent + CTMS
+ eSource (late 2026)

**2,000+
Integrations**

500+ Sponsors

EDC + eCOA + IRT + Safety
+ Feasibility + Training

80% of trials run on Veeva

Benefits

Save Time and Reduce Costs

Sites report spending 50% less time on document management, saving more than \$8k per study using Veeva SiteVault.

Work More Seamlessly with Sponsors

Share trial information across sponsors with no additional setup or complex integrations.

Relieve Overwhelmed Staff

Minimize duplicate data entry, inconsistent processes and multiple logins so study team members can focus on what matters most.

Faster Study Execution

Leverage actionable dashboards and metrics across studies and teams to identify bottlenecks and pinpoint priorities.

Simplify Patient Participation

Flexible eConsent allows sites to expand their reach and enhance their patient experience.

Purpose-Built Tech

A unified suite (eISF, CTMS, eSource) designed to handle all of your research needs in one place.

Guided by Sites, For Sites

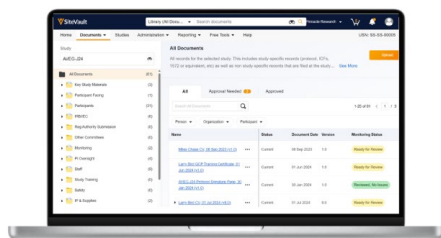
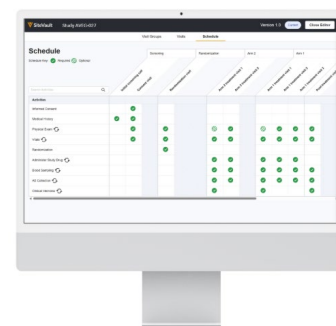
We don't build in a vacuum. Our Site Advisory Council—composed of leaders from large Academic Medical Centers to community-driven research clinics—directly guides our roadmap and design.

Our Rapid Innovation Cycle: We are continuously innovating to cover the full spectrum of your research needs. Delivering three product-rich releases every year, ensuring your site always has the latest tools to stay competitive.

Veeva's Suite of Research Site Applications:

Veeva SiteVault CTMS

Veeva SiteVault CTMS allows study teams to capture and track patient activity throughout a trial, from scheduling study visits and procedures, to generating invoices and tracking payments. Since it's built on the same platform that powers the clinical operations of 450+ sponsors and CROs, and is seamlessly connected to Veeva SiteVault eISF and Veeva SiteVault eConsent, Veeva SiteVault CTMS minimizes duplicate data entry to save study teams time and money.



Veeva SiteVault eISF

Veeva SiteVault eISF enables sites to stay organized and compliant without the effort, with an easy-to-use electronic investigator site file (eISF) that fosters consistent record-keeping processes, provides study teams and monitors with self-serve access to information, and simplifies document exchange with sponsors. Veeva SiteVault eISF supports compliance with global industry regulations and can be used across all of your studies.

Veeva eSource

Veeva eSource is a holistic data capture application that will provide research sites with everything they need to capture patient data digitally at the source, eliminating the need for paper. Veeva eSource will be built to integrate with Electronic Health Records (EHRs) and unified with the industry-leading Veeva Electronic Data Capture (EDC), which will transform the way that research sites collect and manage trial data.



Scan to learn more

Ready to transform your sites operations?

Connect with the Veeva site solutions team and learn how to get started.



Scan to get started