Veeva SiteVault

High Quality Applications Built for Research Sites

Veeva SiteVault improves efficiency, quality, and collaboration for research sites with unified **CTMS, eISF and eConsent.** 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**

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.

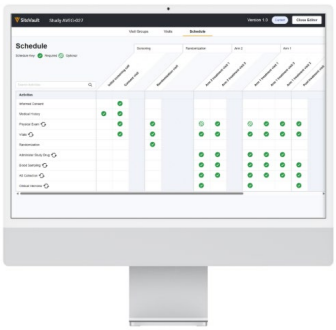
Faster Study Execution
Leverage actionable dashboards and metrics across studies and teams to identify bottlenecks and pinpoint priorities.

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

Veeva SiteVault

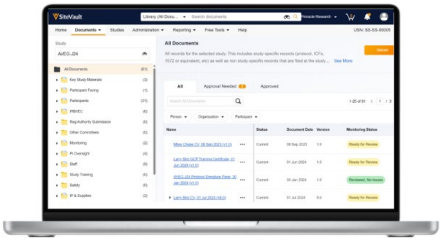
Veeva SiteVault CTMS

Coming Soon:
Available for Early Adopters in August 2025



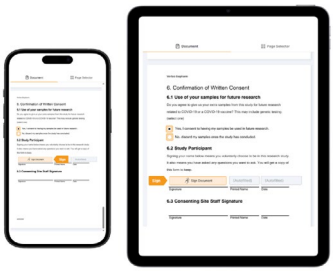
Veeva SiteVault CTMS allows study teams to capture and track patient activity throughout a trial, from scheduling study visits and procedures, to generating invoices and tracking payments. Since it's built on the same platform that powers the clinical operations of 450+ sponsors and CROs, and is seamlessly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



Veeva SiteVault eConsent makes patient participation in clinical trials easier, while reducing administrative burden for study teams and ensuring compliance for sites of all sizes. With flexible options to consent patients in-person or remotely, Veeva SiteVault eConsent is unified with Veeva SiteVault eISF to make the consent process efficient for study teams, and includes MyVeeva, an easy-to-use, patient-centric application to foster patient engagement.

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