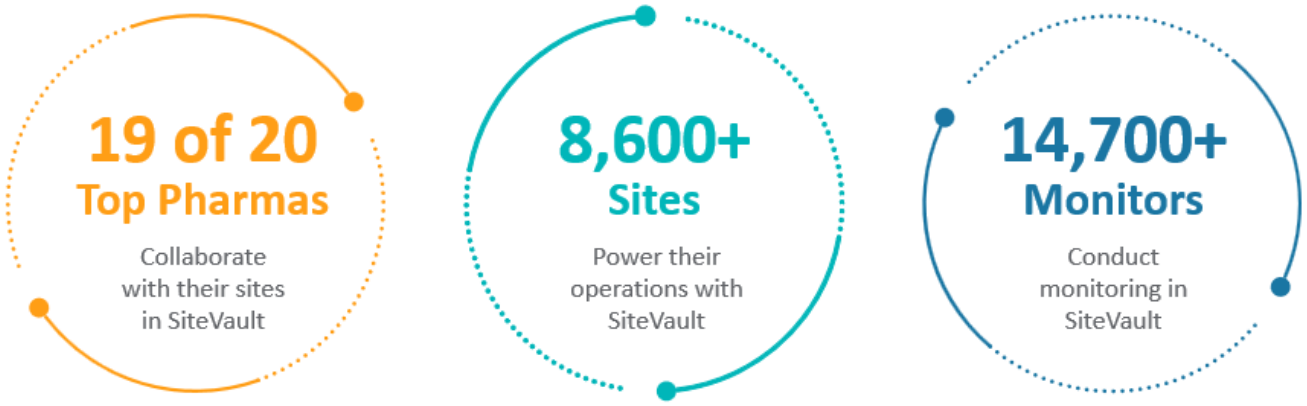
Veeva SiteVault

Unified Solutions Built for Research Sites

Veeva SiteVault improves efficiency, quality, and collaboration for research sites, unifying **CTMS, eISF and eConsent** within a single site platform. SiteVault seamlessly integrates with sponsor systems 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 SiteVault.
- **Work more seamlessly with your 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.
- **Speed 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.

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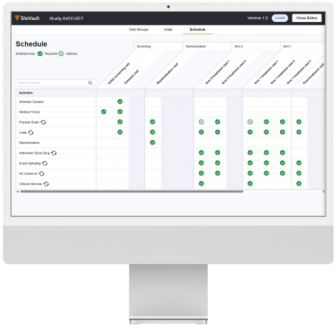
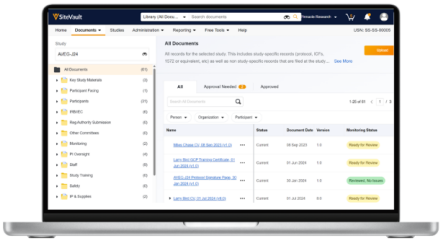
SiteVault CTMS

SiteVault CTMS streamlines clinical trial management by giving study teams a centralized system to manage protocols, track participant activity, and oversee study financials from start to finish. From pre-award budgeting and visit scheduling to post-award invoicing and payment reconciliation, SiteVault CTMS supports every phase of a study. Built on the same platform trusted by 450+ sponsors and CROs, it reduces duplicate data entry and keeps operations aligned, saving sites time, effort, and unnecessary costs.



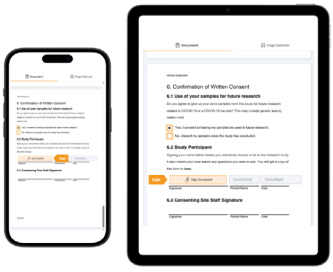


SiteVault eISF

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. SiteVault eISF supports compliance with global industry regulations and can be used across all studies.

SiteVault eConsent

Veeva 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 eConsent makes the consent process efficient for study teams, and includes MyVeeva, an easy-to-use, patient-centric application to foster patient engagement.



Scan to register your site for Veeva SiteVault