Veeva SiteVault

Owned by sites. Loved by sponsors.

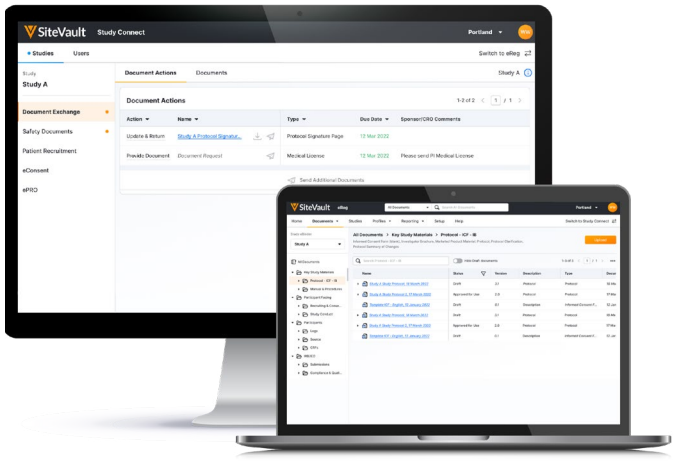
> *"Veeva SiteVault helps us dedicate more time to patient recruitment and care."*
>
> **– Luis Ruda,**
> *Director, Alliance Clinical Research*

Overview

Veeva SiteVault is a free, easy-to-use application that connects clinical research sites to sponsors and CROs to reduce the number of systems and logins used to run clinical trials.

Manage your studies and exchange information across sponsors and CROs in a single, site-owned system.

Veeva SiteVault brings better, more efficient collaboration between sites and sponsors, and more time to focus on your patients.



Benefits

- **One place to work with sponsors** - Easily exchange study information through a single application that's connected to more than 400 sponsors and CROs that use Veeva Clinical applications.
- **Stay organized and compliant, without the effort** - Centralize regulatory documents and simplify processes with an eReg system that's fully validated and easy to set up.
- **No cost** - Getting started is easy and there's no risk, or cost, to you. You get comprehensive support and a team of former clinical research professionals to ensure your success.

5,000+
Sites across 80+ countries use Veeva SiteVault

400+
Sponsors use Veeva Clinical applications

65%
Of all global industry trials run on Veeva

Work With Your Sponsors

Use **Study Connect** to work with sponsors and CROs using Veeva Clinical applications. Setup is easy and sites retain full visibility of historical data in a secure, site-owned system. Sites will be invited to use Study Connect by the trial sponsor or CRO.

- **Seamless Document Exchange**
  Receive and exchange start-up packages, regulatory documents, invoices, payment info, and end-of-study media to reduce manual steps and redundant requests.
- **Auto-file Documents into SiteVault eRegulatory**
  Provide investigators and staff with quick access to information with a system that automatically files study documents into SiteVault eRegulatory.
- **Receive and Acknowledge Safety Letters**
  Simplify the receipt and acknowledgement of safety letters and quickly file them into SiteVault eRegulatory to save time and ensure investigators stay informed.
- **Manage eConsent**
  Make patient participation in clinical trials easier by providing convenient access to consent forms and key study information.

Manage Your Studies

Use **eReg** as your 21 CFR Part 11 compliant investigator site file (ISF). Spend less time searching for the right document and chasing signatures. There is no limit to the number of studies, documents, or users. Sites can sign up for free at any time.

- **Remote Monitoring**
  Provide monitors with secure access to review regulatory and source documents.
- **Digital Delegation**
  Get more done with an intelligent system that auto-populates the delegation log while you work.
- **Workflows**
  Complete document tasks right in the system with eSignature, certified copy, and training workflows.
- **Reports and Dashboards**
  Quickly see open tasks, expiring documents, and more with configurable reports and dashboards.
- **eConsent**
  Deliver a better patient experience while creating efficiencies for your site. Veeva eConsent may be provided by your sponsor or enabled as a paid add-on to use across all studies.
- **Enterprise Add-on**
  Includes Single Sign-on, API support for integrations, field customization, and dedicated enterprise-level customer support.

> *“With Veeva SiteVault, communication is much improved and we have a better relationship with our sponsors. In one study, we went from sending about 140-150 emails to the sponsor down to 30-35 during study start-up.”*
>
> ***– Jim Sanders,***
> *President, ClinOhio Research Services, LLC.*