Datasheet

Veeva SiteVault

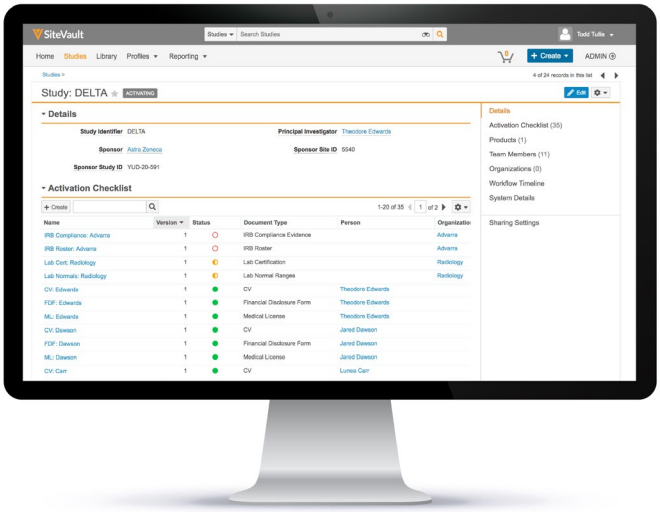
Less Red Tape, More Research

Simplify research by managing regulatory documents and processes in a modern eRegulatory system

Overview

Veeva SiteVault reduces the administrative burden in clinical trials by simplifying the management of regulatory documents and processes across all studies to speed study execution.

SiteVault can be used for all trials regardless of what technology sponsors are using, as well as the site file for investigator initiated trials.



Benefits

Reduce administrative burden: Replace manual and paperbased processes by managing regulatory documentation and trial information in a modern, Part 11 compliant system.

Speed study activation: Eliminate repetitive tasks and accelerate turnaround timelines to activate studies sooner

Increase visibility: Gain real-time visibility into document status to improve compliance and oversight

Top Companies Use Veeva

Veeva is the leading provider of clinical operations technology to more than 200 sponsors and leading research organizations.

iact HEALTH · UNIVERSITY OF LOUISVILLE · VERTEX · BAYER · gsk · astellas · parexel · MERCK

Veeva SiteVault

Choose the plan that is right for you

Standard Features	Free	Enterprise
Full eRegulatory system Provide investigators and staff with easy access to study documents through an intuitive electronic regulatory binder that supports compliance with 21 CFR Part 11 and HIPAA requirements.	✓	✓
Electronic signatures Simplify approvals and replace printing, faxing, and scanning with fully electronic signature workflows that investigators and staff will love.	✓	✓
Remote monitoring Provide monitors with secure, direct access to study binders from any location.	✓	✓
Auto-filing and auto-naming Improve compliance by automatically naming and filing study documents into eBinders. Quickly update CVs, medical licenses, and staff information across multiple studies with a single action.	✓	✓
Standard workflows Save time and centralize tasks with built-in workflows for eSignatures and certified copies.	✓	✓
Standard reports Improve visibility with powerful reports that provide visibility into open tasks, upcoming expiration dates, and signature turn-around timelines.	✓	✓
Secure cloud platform Veeva SiteVault is designed to meet the rigorous content and data management requirements of the life sciences industry.	✓	✓
Unlimited studies and users Both versions support an unlimited number of users and studies and comes with full customer support from Veeva.	✓	✓
Enterprise Features		
Configurable workflows and user groups Modify standard workflows to fit your SOPs. Provide visibility into study documents to different groups across your organization.		✓
Self-service dashboards and reports Design reports and dashboards based on metrics you care about most to identify bottlenecks and take immediate action.		✓
Real-time collaborative authoring Seamless integration between Veeva Vault and Microsoft Office Online provides real-time collaborative authoring.		✓
Open API Reduce duplicate data entry and streamline processes by integrating SiteVault with your other with research systems.		✓
Enterprise single sign-on Quickly sign and access documents using the same set of credentials as your other applications.		✓
Unlimited document retention period Preserve and archive your regulatory documents for as long as you use Veeva SiteVault.		✓