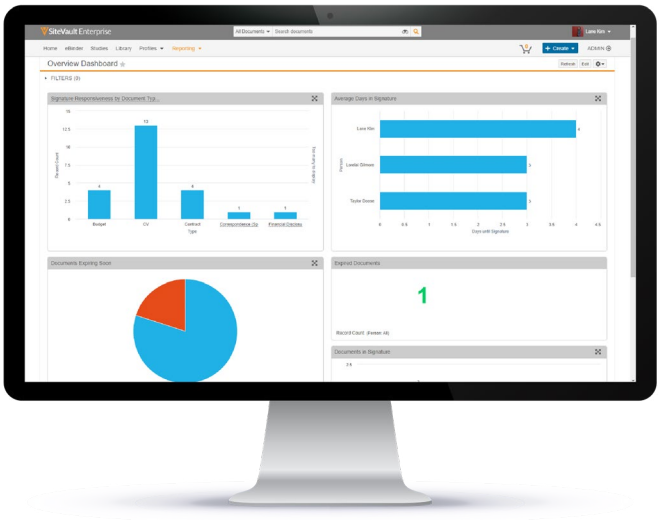
Veeva SiteVault

Transform Your Research Operations

Your workspace, digitally connected to patients and sponsors.



Overview

Veeva SiteVault reduces the administrative burden in clinical trials by simplifying the management of the regulatory and source documents in one system so that sites can spend time where it matters most – with patients.

Manage regulatory and source documents in a system that supports 21 CFR Part 11 and HIPAA requirements. Save time with built-in tools to remotely consent patients, capture electronic signatures, enable remote monitoring, certify copies, and view reports.

Benefits

- Save time and work remotely
- Provide a better experience for your patients and sponsors
- Always be inspection ready

Join Over 1,000 Successful Research Sites Across 60+ Countries

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ALLIANCE OHIO SLEEP MEDICINE INSTITUTE CENTER OF SLEEP MEDICINE EXCELLENCE™ JBR CLINICAL RESEARCH

Choose the plan that is right for you

SiteVault Free

100% Free eRegulatory and eConsent solution.

- Full eRegulatory System
- eConsent
- Remote Monitoring
- Electronic Signatures
- Auto-Filing and Auto-Naming
- Version Compare
- Real-Time Collaborative Authoring
- Upload Source Documents in Bulk
- Exchange documents with sponsors and CROs
- Built-in Workflows
- Reports and Dashboards
- Secure Cloud Platform
- Unlimited Studies and Users
- 25-Year Document Retention Period

SiteVault Enterprise

Includes everything in **SiteVault Free**, as well as:

- Configurable Workflows
- Configurable Reports and Dashboards
- Configurable User Groups
- Scan Documents on the Go With Veeva Snap
- Open API
- Enterprise Single Sign-on
- Unlimited Document Retention Period