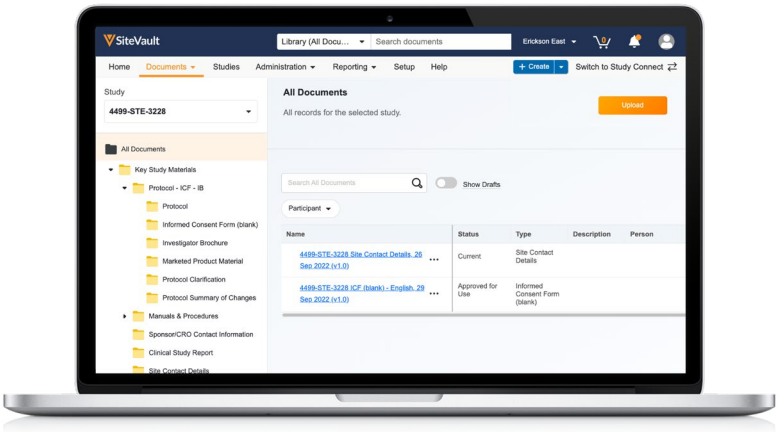
Veeva SiteVault eISF

Overview

Veeva SiteVault eISF is an easy-to-use electronic investigator site file (eISF) that reduces the burden of managing paper binders so research teams can focus on what matters most.

Sites save time by establishing consistent record-keeping processes and providing study teams and monitors with self-serve access to information, which reduces costs and frees up staff time. Veeva SiteVault eISF supports compliance with global industry regulations and can be used across all of your studies.



Benefits

Save Time and Reduce Costs
Sites report spending 50% less time on document management, saving more than $10k per study using Veeva SiteVault eISF

Relieve Overwhelmed Staff
File, search and access regulatory documents more easily to maintain compliance with less burden.

Work More Seamlessly With Sponsors
Simplified document exchange and self-service monitoring makes working with sponsors easier and less transactional.

"We are reducing study activation timelines by 40%, spending half the time completing regulatory tasks, and saving tens of thousands of dollars per study with Veeva SiteVault eISF. Veeva has helped us dramatically improve the speed and efficiency of study execution for more than 50 studies across 18 different sponsors."

Justin Deck, Chief Clinical Officer, Tilda Research

Veeva SiteVault

Veeva SiteVault eISF Features

Electronic Investigator Site File
Stay organized and efficiently manage regulatory and source documents across all your studies and sponsors with the industry-standard eISF reference model.

Monitoring
Provide secure, direct monitor access to source and regulatory documents to save time on visit preparation and collaborate with monitors in real time.

Easy Information Exchange with Sponsors
450+ Veeva eTMF customers can connect to your SiteVault for seamless exchange of documents and trial information.

Unified with CTMS & eConsent
Minimize duplicate data entry to improve study efficiency, quality and collaboration with seamlessly connected CTMS and eConsent, built on the SiteVault platform.

eSignatures, Approvals & Training
Finalize documents faster, streamline training completion, and eliminate manual trackers by completing tasks right in the system.

Advanced Reports & Analytics
Prioritize your work and make informed decisions with visibility into document expiration, eSignature turnaround times, workloads, monitoring, and more.

Digital Delegation
Simplify Delegation of Authority Log management and gain clarity into active assignments with a fully digital workflow to ensure DOA compliance.

Central Documents
Store central documents (e.g. licenses, certifications, CVs) in a single location and automatically update them across relevant studies.

Real-Time Document Collaboration
Allow multiple users to edit documents in SiteVault at the same time by connecting to your Microsoft Office 365.

Vault Mobile App
Complete document training, reviews and eSignatures with biometric verification on the go, with a purpose-built and secure mobile app.

MyVeeva for Patients
Make trial participation more accessible and convenient, with a simple, intuitive app for patients to learn about their trial, view and sign documents, and more.

Study Archiving
Automate archiving readiness checks and compliantly archive studies in minutes. Veeva maintains documents and data for a 25-year retention period.

API Integration
Connect SiteVault to your other systems with the public Vault API, already used to connect to over 100 unique applications.

Compliant & Validated
Gain peace of mind with a solution that supports 21 CFR 11, Annex 11, HIPAA and GDPR. Veeva maintains all validation and security documents for you.

Dedicated Support
Save time and get the most out of SiteVault with a dedicated expert invested in your team's success and a 98.5% customer satisfaction rating.

Scan to register your site for Veeva SiteVault