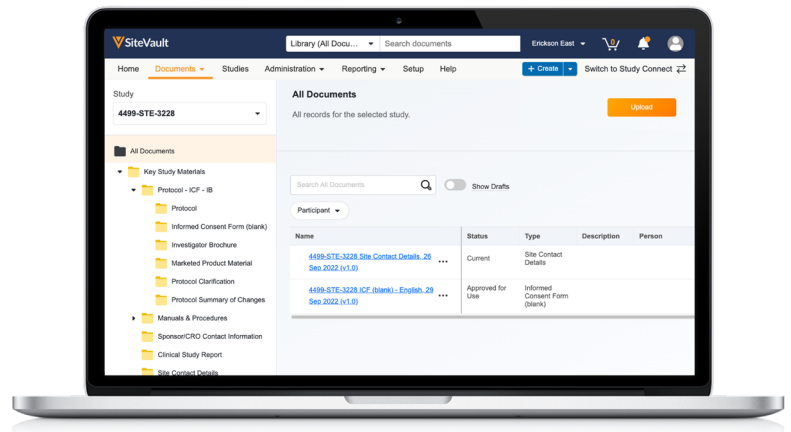


# Veeva SiteVault eISF

## 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 fees up staff time. SiteVault supports compliance with global industry regulations and can be used across all of your studies.



## Benefits

### Save Staff Time and Reduce Cost

File, search and access protocol and regulatory documents more easily to maintain compliance with less burden, spending 50% less time on document management, saving >\$10k per study.

### Work More Seamlessly with Sponsors

Simplified document exchange and self-service monitoring makes working with sponsors easier and less transactional.

### Stay Inspection-Ready

Maintain version control, audit trails, and complete document histories so you're always prepared for monitoring visits or regulatory inspections.

### Collaborate Across Teams

Give coordinators, investigators, and regulatory staff real-time access to the documents they need, in a standardized format — no emails, printing, or chasing signatures required.

“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

## Features



### Electronic Investigator Site File

Stay organized and efficiently manage regulatory documents across all your studies and sponsors with the industry standard eISF reference model.



### Advanced Reports & Analytics

Prioritize your work and make informed decisions with visibility into document expiration, eSignature turnaround timelines, workload and more.



### Monitoring

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



### Real-time Document Collaboration

Allow multiple users to edit documents in SiteVault at the same time by connecting your Microsoft Office 365.



### Digital Delegation

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



### Vault Mobile App

Complete document training, reviews, and eSignatures with biometric verification on the go with a purpose-built and secure app



### eSignatures, Approvals, & Training

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



### 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.



### Central Documents

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



### Study Archiving

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



Scan to register your site  
for Veeva SiteVault