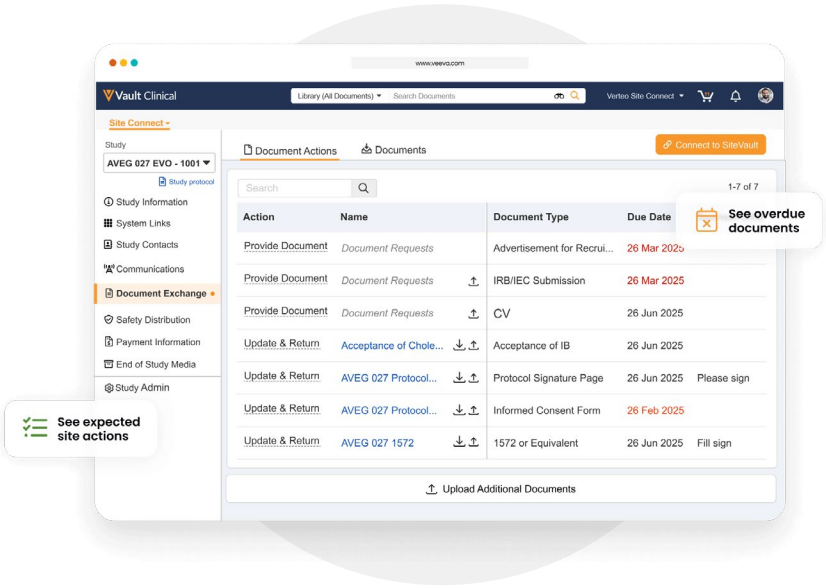
FEATURES BRIEF

Veeva Site Connect



Veeva Site Connect allows sponsors and research sites to collaborate on a trial by automating the flow of information to and from sites during startup, execution, and closeout.

Information flow includes protocols, essential document packages, safety reports, and payment letters. Required media is sent on closeout, including completed CRFs. Information sent and received is automatically filed in eTMF. Research sites manage tasks, documents, and data in Site Connect. Optionally, sites can connect their SiteVault for enhanced functionality.

Benefits

One system for sites and sponsors to collaborate

Sponsors can execute collaboration tasks across the trial lifecycle for all their sites in one application. Sites work in an optimized user interface that's the same across sponsors - giving them a standard way to work across all trials.

Open for use by any site, anywhere

Site Connect is accessible for all sites everywhere. Sites that decide to use Veeva SiteVault as their eISF get the added benefit of connecting their study for seamless bidirectional document exchange.

Faster, higher quality trials

By simplifying and standardizing site-sponsor collaboration, sponsors and sites reduce administrative burden so they can conduct higher quality trials at a lower cost.

Features

Document Exchange

Easily manage and distribute site documents automatically during study initiation to speed study start-up.

Safety Letter Distribution

Guarantee delivery and tracking of important safety letter information sent to sites so that principal investigators stay informed.

Study Information

Access study information such as name, description, site number, and a link to the protocol in one convenient location.

Study Announcements

Share important announcements across all sites, such as deadlines, key dates and notifications.

Study Contacts

Provide sites key contact information that's relevant for their study—such as role, name, phone, and email.

End-of-Study Media

Import, distribute, and track final subject data—such as completed CRF output—from EDC to clinical research sites, including auto-filing in eTMF.

Payment Information

Deliver reimbursement data and payment letters to clinical research sites and receive site invoices directly within Vault Clinical.

eTMF and eISF Connectivity

Sites may optionally use SiteVault so that clinical documents are automatically exchanged with Vault Clinical. This reduces manual steps associated with sponsor TMF and site eISF reconciliation activities.

About the Veeva Clinical Platform

The Veeva Clinical Platform improves clinical research by providing the most complete and highest-quality solution built for the unique needs of patients, research sites, and trial sponsors. With seamless connection and data flow across all stakeholders, Veeva Clinical Platform enables faster, more efficient trials that achieve higher data accuracy, and deliver a better experience for sites and patients.