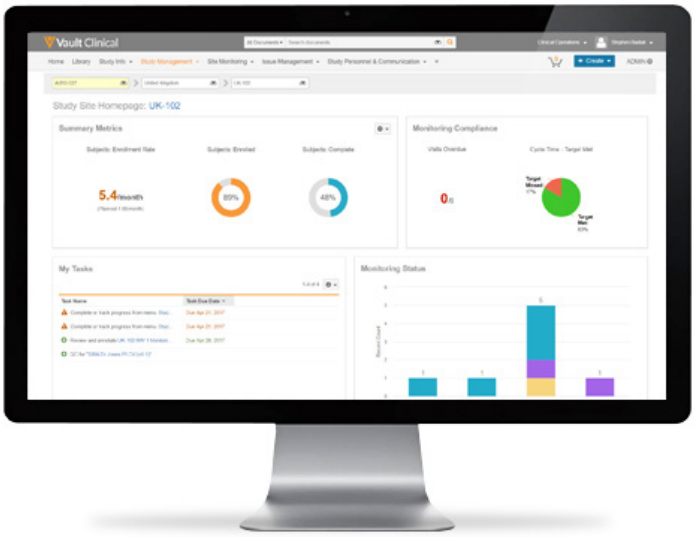
Veeva CTMS



Veeva CTMS is the only modern cloud application that makes it easy to unify clinical information and processes, streamline trial management, and gain complete visibility across the trial portfolio.

Study teams can manage the entire end-to-end clinical trial process and gain a global view into tasks in one unified and secure system. With simple navigation and a single source for clinical master data and study information, Veeva CTMS improves operational efficiency and enables faster, higher-quality trial execution. Streamline operations with a flexible, agile solution that easily adapts to your organization's unique clinical trial needs, study designs, and business processes.

Benefits

Enable Faster Trials. Easy-to-use dashboards generate real-time, actionable insights. Quickly identify sources of delay and take corrective action to restore momentum, improving overall trial efficiency and performance.

Improve Decision-Making. Strategic trial planning is easier with a full view of global operations in a single system. Get a complete and accurate view of clinical trial status to make informed decisions faster and enable proactive closed-loop issue management with real-time visibility across all studies.

Streamline Clinical Operations. By providing a seamless connection to the Veeva Clinical Operations Platform, organizations and their study partners have one source of truth for shared TMF, CTMS, and site payments data, eliminating complex integrations.

A Single Source of Truth

With the Veeva Clinical Operations Platform, study teams enjoy a consistent experience with single sign-on and avoid constantly switching between multiple systems. Submit trial information and documentation once and leverage it across different systems, sites, and countries. This single source of truth improves visibility and control and accelerates trial execution.

Study Planning and Management

Plan and track study milestones across trial activities to optimize resources and proactively plan for events such as aligning clinical supply arrival with the site initiation visit or assessing site performance across studies. Veeva CTMS enables seamless subject visit planning based on categories such as protocol, visit frequency, and procedures.

Subject Recruitment Planning

Plan the number of subjects that will be screened, enrolled, or randomized within a study and get a comprehensive view at the study, study country, and site levels. With metrics that update with actual recruitment data, you can track subject enrollment against goals to ensure studies are on time.

Site Monitoring

Manage all aspects of routine monitoring visits—pre-study, site initiation, interim monitoring, and closeout—in Veeva CTMS. CRAs can view key information such as enrollment metrics and violations at-a-glance on the CRA homepage, quickly author new monitoring visit reports, and track onsite monitoring activities, all in one application.

Issue Management

Easily capture, track, and manage protocol deviations, issues, and follow-up items across all studies. Quickly identify actions needed, assess clinical task statuses, and document quality issues to enable closed-loop issue management.

Investigator Relationship Management

Empower study teams with an accurate and complete view of interactions between sponsors, CROs, investigators, and site personnel. Track site communication logs, site monitoring visits, resources assigned to sites, and more to strengthen collaboration and improve study execution.

Site Payments

Unified with Veeva CTMS, Veeva Payments speeds payments to clinical research sites and provides financial visibility to all study partners. Designed to support complex trials, Veeva Payments enables sponsors and CROs to pay sites faster and more accurately.

Study Oversight

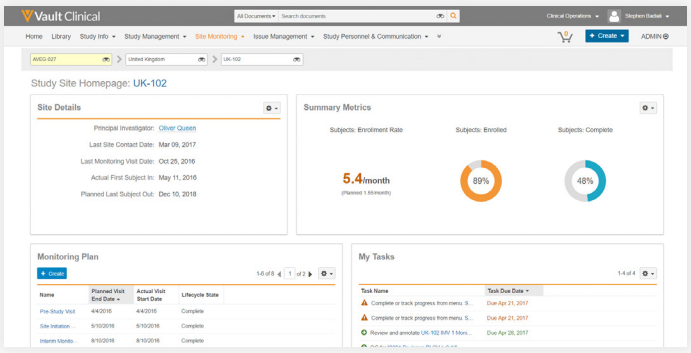
Monitor study performance, track CRO activity, and maintain communication logs to help ensure regulatory compliance with ICH/GCP guidelines. CRAs can also capture and track protocol deviations for effective issue management.

Interactive Dashboards and Reports

Create reports that show real-time operational metrics, documentation, and information by study, country, investigator, site activation status, and more. Study teams can take immediate action directly from dashboards, eliminating bottlenecks and increasing efficiency.

Connection to Veeva EDC

Eliminate duplicate data entry and transcription errors and provide real-time visibility into enrollment status at every site. Study managers and CRAs can move seamlessly from subjects or subject visits in Veeva CTMS to the applicable events and forms in Veeva EDC without a separate login.



Veeva Clinical Platform

The Veeva Clinical Platform is the first eClinical platform offering EDC, CDB, eCOA, eTMF, CTMS, Payments, Site Connect, Study Training, and RTSM on one enterprise-class cloud platform. For the first time, medtech companies can connect clinical operations and data management with a unified platform to create a single source of truth and streamline clinical trials from study startup to close.