A Unified Suite of Applications for Clinical Operations and Data Management

Decentralize clinical trials and streamline data management to bring agility and speed to clinical research.

Increasing demand for real-world evidence, data, and stringent global regulatory requirements is making clinical research more complex, resulting in longer product development cycles. The ability to streamline study processes and data management is crucial to driving innovation and getting medical devices and diagnostics to patients fast.

The Vault Clinical Suite is the industry's first platform for Medtech that combines CDMS, eTMF, CTMS, and payments to deliver the most comprehensive suite of clinical applications in the cloud. Veeva MedTech cloud software helps streamline clinical operations and data management with a single source of information and collaboration.

With dedicated applications for managing and tracking all clinical activities, including study documents, sponsors, site payments, and more, the suite of applications standardizes the entire clinical research process.

Key Business Benefits

- **Speed Trial Study Timelines:** Enable technology-driven process improvements throughout trials from database builds to monitoring.
- **Realtime Inspection Readiness:** Manage data in real-time for better decision-making and visibility into trial status and reporting.
- **Streamline Operations:** One source of truth for shared TMF, CTMS and site payments with connected solutions.
- **Unify Clinical Data & Operations:** Streamline processes and enable sponsors, CROs, and sites to collaborate, in the cloud, throughout the clinical trial process.

Savings Realized

40%
reduction in TMF reconciliation time with Vault eTMF

90%
reduction in monitoring visit preparation time

50%
faster study builds

Vault Clinical Suite Applications

- **Vault CDMS** – Accelerate study execution by combining EDC, coding, data cleaning, and reporting
- **Vault eTMF** – Real-time inspection readiness, full TMF status visibility into TMF status, access to all study partners
- **Vault CTMS** – Single source for clinical operations with global visibility and end-to-end trial process management
- **Vault Payments** – Speed payments to research sites and provide complete visibility to all study partners
- **Site Connect** – Connect sponsors and research sites by automating information flow

Solution Features

Fully Configured Studies

Veeva MedTech expert services team delivers fully configured studies to sponsors and CROs. Innovative features such as a rules engine for edit checks, drag-and-drop form design, and self-documenting specification creation enable Veeva to build studies quickly and efficiently with no downtime or migrations.

Study Planning and Visibility

Plan and track study milestones across trial-related activities to optimize trial resources and proactively plan for events such as aligning clinical supply arrival with site initiation visits or assessing site performance across studies.

Single Source for Clinical Master Data

Ensure high-quality data across clinical applications with one system of record for master study, study country, and study site information.

Easily Optimize for Your Trial

Streamline operations with a flexible, agile solution that easily adapts to your organization's unique clinical trial needs, study designs, therapeutic areas, and business processes. Easily apply protocol amendments and eliminate the need for manual study trackers.

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