Veeva Vault

FEATURES BRIEF

Regulated Content and Data Management in the Cloud

Always Current, Constantly Innovating

Veeva Vault is a cloud-based content management platform and suite of applications that provides life sciences companies a single source of truth to reduce complexity and increase business agility. Veeva Vault not only manages regulated documents, but also tracks critical information from product development to commercialization. Managing this unique combination of documents and data, organizations improve efficiency and gain deeper insight.

Access

Connecting all parties globally promotes and streamlines collaboration. Quickly provide secure access to internal and external users—incorporating all stakeholders into key processes and enabling greater visibility, and control.

Visibility

In any regulated environment, it is critical to know what content exists, what state it is in, and where it is used. Vault's real-time reporting and dashboards empower managers with information for faster decision-making to help identify and remedy process bottlenecks, track progress, and gain visibility into readiness.

Control

Veeva Vault simplifies compliance by providing the functionality you need with unparalleled ease-of-use. Easily gain control over document and data processes with capabilities such as versioning, e-signatures, and controlled workflows. Vault also automates manual steps and keeps collaborators on-task to improve efficiency.

Vault Platform

Veeva Vault is the first cloud platform built from the ground up to meet the rigorous content management requirements of the life sciences industry. With a modern user experience and uniquely designed for both content and data on a single platform, organizations can seamlessly manage end-to-end processes.

The Vault Platform leverages the latest in cloud technology and is delivered and accessed through the web for greater ease-of-use. Hosted at SOC I Type II and ISO 27001 certified global data centers, every release is IQ and OQ qualified reducing the validation efforts.

Veeva Clinical Data Management

Accelerate study timelines with modern, innovative applications for clinical data.

Veeva Clinical Data Management Suite (Veeva CDMS) is redefining data management to help clinical teams manage today's trials with agility and speed.

Veeva EDC
Collect, clean, and review study data.

Veeva CDB
Manage complete and concurrent study data.

Veeva Coder
Rapid coding for clinical terms.

Veeva Clinical Operations

The only suite of unified clinical operations applications on a single cloud platform.

The industry's first and only suite of unified clinical operations applications—including study start-up, eTMF, CTMS, payments, and site connect on a single cloud platform—to accelerate trial execution and deliver real-time visibility.

Veeva Study Startup
Accelerate time to site activation.

Veeva CTMS
Enable proactive trial management.

Veeva eTMF
Enable active eTMF for real-time inspection readiness.

Veeva Payments
Pay clinical research sites faster.

Veeva eConsent
Improve patient experience.

Veeva Site Connect
Automate information sharing.

Veeva Quality

Modernize quality management while driving compliance and operational innovation.

Seamlessly manage your quality processes and content with the Veeva Quality suite of applications. All parties have access to a single authoritative source, enabling greater visibility and control.

Veeva QMS
Easily manage all quality processes.

Veeva Product Surveillance
Simplify postmarket surveillance for medical devices.

Veeva QualityDocs
Document control for all GxP documents.

Veeva Station Manager
Deliver the right content to the shop floor.

Veeva Training
Ensure compliance and role-based qualification.

Veeva Learn GxP
Accredited GxP eLearning courses.

Veeva Validation Management
Execute paperless validation.

Veeva RIM

Bring speed and agility to your regulatory team with unified RIM.

Manage the planning, execution, and oversight of all regulatory activities within a single, unified RIM platform.

Veeva Registrations
Manage product registrations globally.

Veeva Submissions
Speed submission development.

Veeva Submissions Publishing
Automate publishing during submission development.

Veeva Submissions Archive
Archive published dossiers securely in the cloud.

Veeva Safety

Veeva Safety is the only modern application for the collection, management, and real-time oversight of adverse events.

Veeva Safety
Real time management and oversight for adverse events.

Veeva SafetyDocs
Centrally manage pharmacovigilance content.

Veeva Signal
Manage signals from detection through risk evaluation and mitigation.