Veeva Vault

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.



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.



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 decisionmaking to help identify and remedy process bottlenecks, track progress, and gain visibility into readiness.



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.

Vault Clinical Data Management

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

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

Vault EDC

Collect, clean, and review study data.

Vault Coder

Rapid coding for clinical terms.

Vault 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.

Vault Study Startup

Accelerate time to site activation.

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

Veeva eConsent

Improve patient experience.

Vault Quality

Modernize quality management while driving compliance and operational innovation.

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

Vault QMS Easily manage all quality processes.

Vault QualityDocs Document control for all GxP documents.

Vault Training Ensure compliance and role-based qualification.

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

Vault Validation Management Execute paperless validation.

Manage product registrations globally.

Vault Submissions Publishing

Vault Product Surveillance Simplify postmarket surveillance for medical devices.

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

Veeva Learn GxP Accredited GxP eLearning courses.

Enable proactive trial management.

Pay clinical research sites faster.

Automate information sharing.

Vault Safety

Vault Registrations

Vault RIM

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

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

Vault Safety

Vault SafetyDocs Centrally manage pharmacovigilance content.

Speed submission development.

Archive published dossiers securely in the cloud.

Vault Submissions Archive

Vault Submissions

Vault Signal

Manage signals from detection through risk evaluation and mitigation.

Real time management and oversight for adverse events.

Automate publishing during submission development.

Copyright © 2024 Veeva Systems Inc. All rights reserved. Veeva, V, Vault and Crossix are registered trademarks of Veeva Systems Inc. Veeva Systems owns other registered and unregistered trademarks. Other names used herein may be trademarks of their respective owners.

Veeva CDB

Vault CTMS

Vault Payments

Veeva Site Connect

Manage complete and concurrent study data.