Datasheet

Veeva Vault

Regulated Content Management in the Cloud

Always Current, Never Stuck

For decades, life sciences companies have dealt with content management solutions that cobble together separate technologies, require months of custom development, and end up too complex to change.

Veeva Vault delivers a regulated content management platform and purpose-built applications for every major part of a life sciences company, from regulatory, to clinical, quality, and commercial. Built and delivered in the cloud, these solutions work out-of-the-box to manage specific documentation needs across the organization. There's no software to install, hardware to maintain, or costly updates to deploy. Also, Veeva's pace of innovation, which is only possible with a multitenant cloud architecture, means Veeva Vault applications will stay modern and fast. The results: happier users, dramatic process improvements, and reduced regulatory risk.

Access

Connect and collaborate globally. Gone are the days when you had to wait weeks (or even months) to provide document access to your trusted external partners. Vault connects internal and external collaborators quickly and securely without costly VPNs or company laptops. Simply add the user, set the access levels, and go. Now your partners can work with you where, when, and how you need them to.

Visibility

In any regulated environment, it is critical to know what content exists, what state it is in, and where it has been used. Vault's real-time reporting and dashboards help managers identify and remedy process bottlenecks. Users drill down through reports to answer questions about progress and readiness. Embedded action menus enable managers to resolve issues directly from the reports, so no time is wasted.

Control

Meeting regulatory requirements can be easier than it is today. Vault simplifies compliance by providing the functionality you need with unparalleled ease-of-use. Capabilities such as versioning, e-signatures, and change comparisons are native in the application. Business owners get control over even the most complex document processes. Vault automates manual steps and keeps collaborators on-task, to improve the efficiency of your team.

Cloud

Unlike traditional systems, Vault is easy to administer and configure. Need to add a new document attribute? No problem. Have to change a document type? Good as done. Want the latest capabilities? It's multitenant cloud; you're always up-to-date. We perform IQ/OQ validation on each release, and you control when to activate new functionality, so your applications stay current and compliant.

Learn more at veeva.com ›

Veeva Vault

Vault Applications

Vault eTMF

Streamline Document Collection, Management, and Analysis.

Vault eTMF brings sponsors, CROs, and investigator sites together in the regulated cloud. Sponsors get complete visibility with real-time dashboards and actionable reports. CROs get a smart eTMF that tracks outstanding content and automatically places documents into the eTMF. The result is complete visibility, faster study start-up, and an inspection ready eTMF.

Vault Investigator Portal

Portal Access for Clinical Trial Investigator Sites.

Vault Investigator Portal helps sponsors and CROs get off paper and maintain an audit-ready eTMF. The interface is simple—just documents and tasks. The backend is powerful—autofiling into the eTMF when relevant documents are exchanged. And since Veeva's portal provides a direct interface to Vault eTMF, companies get better reporting and visibility into study status.

Vault Submissions

Efficient Authoring and Assembly Drive Faster Submissions.

Vault Submissions unites contributors, partners, and affiliates in the cloud with a single destination for regulatory documents. Create global and regional submission dossiers to harmonize the planning process and provide real-time visibility into submission readiness.

Vault QualityDocs

Ubiquitous Access and Breakthrough Usability.

Vault QualityDocs provides a single global source for quality and manufacturing SOPs, reports, and validation documents. Vault makes managing quality documentation easy: there's real-time collaboration without downloading documents. Secure unfettered access for external partners. Ensure easy fulfillment of training requirements with "Read & Understood" tracking.

Vault MedComms

Single Source of Truth for All Medical Communications.

Vault MedComms provides an efficient way to manage global medical communication content. A common repository supports easy collaboration and version control, and an open API allows integration and accurate fulfillment across all communication channels and geographies.

Vault PromoMats

End-to-End Solution for Promotional Material Management.

Vault PromoMats ensures faster time-to-market and compliance from creation to distribution. It supports easy internal and external collaboration, a built-in digital asset library, MLR review, and single-click multichannel distribution and withdrawal with actionable insight to remove bottlenecks at every stage.

Regulated content managed easily and securely in the cloud.

Vault Platform

The Most Modern, Scalable, and Secure Platform for Regulated Content Management

Veeva Vault is the first cloud platform built from the ground up to meet the rigorous content management requirements of the life sciences industry. Employing the very latest in cloud software and hardware, the Vault Platform provides medical device, pharmaceutical, biologics, and in-vitro diagnostic companies with an enterprise-class platform to store and manage all of the company's important documents.

The Vault Platform brings an outstanding user experience and pace of innovation to the world of regulated content management, replacing the costly and painful content management solutions of the past. For the first time, life sciences organizations can manage their content in the cloud with confidence.

The Vault Platform supports all of the Vault applications with the flexibility, controls, and deep functionality required to make Vault the choice for regulated content management.