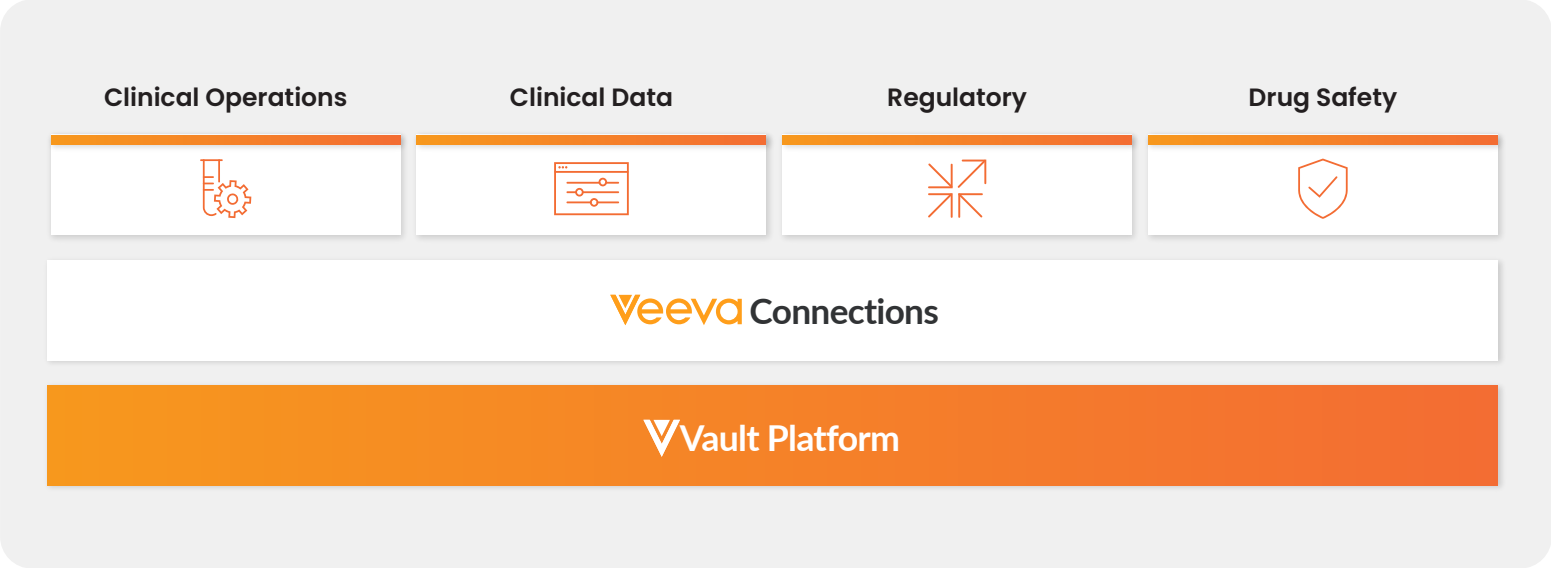
Veeva Development Cloud

SOLUTION BRIEF

Technology Foundation for Product Development

Veeva Development Cloud is the technology foundation for product development that brings together applications for clinical, regulatory, and safety to help organizations drive end-to-end business processes. Today, product development systems are not well integrated, which creates inefficiencies and slows down critical operations. Veeva is the first and only company to offer unified suites of applications that are connected on a single cloud platform. This enables organizations to centralize content and data across global departments for greater efficiency and compliance.



Veeva Connections

Veeva Connections are Veeva-delivered integrations that seamlessly transfer data and documents between clinical, regulatory, and safety Veevas. They are designed to streamline cross-functional business processes by breaking down silos, providing greater visibility, and automating manual tasks. View the Veeva Connections Resource Hub for a full list of available Veeva Connections.

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 eTMF

Enable active eTMF for real-time inspection readiness.

Veeva CTMS

Enable proactive trial management.

Veeva Payments

Pay clinical research sites faster.

Veeva Study Startup

Accelerate time to site activation.

Veeva RTSM

Randomize subjects and manage trial product supply.

Veeva Site Connect

Automate information sharing.

Veeva Study Training

Streamline and automate training.

Veeva Clinical Data

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

Veeva Clinical Data Management helps clinical teams collect, aggregate, clean, and manage trial data with agility and speed.

Veeva EDC

Collect, clean, and review study data.

Veeva CDB

Manage complete and concurrent study data.

Veeva eCOA

Capture responses directly from clinical trial participants.

Veeva eConsent

Simplify informed consent with an end-to-end process.

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.