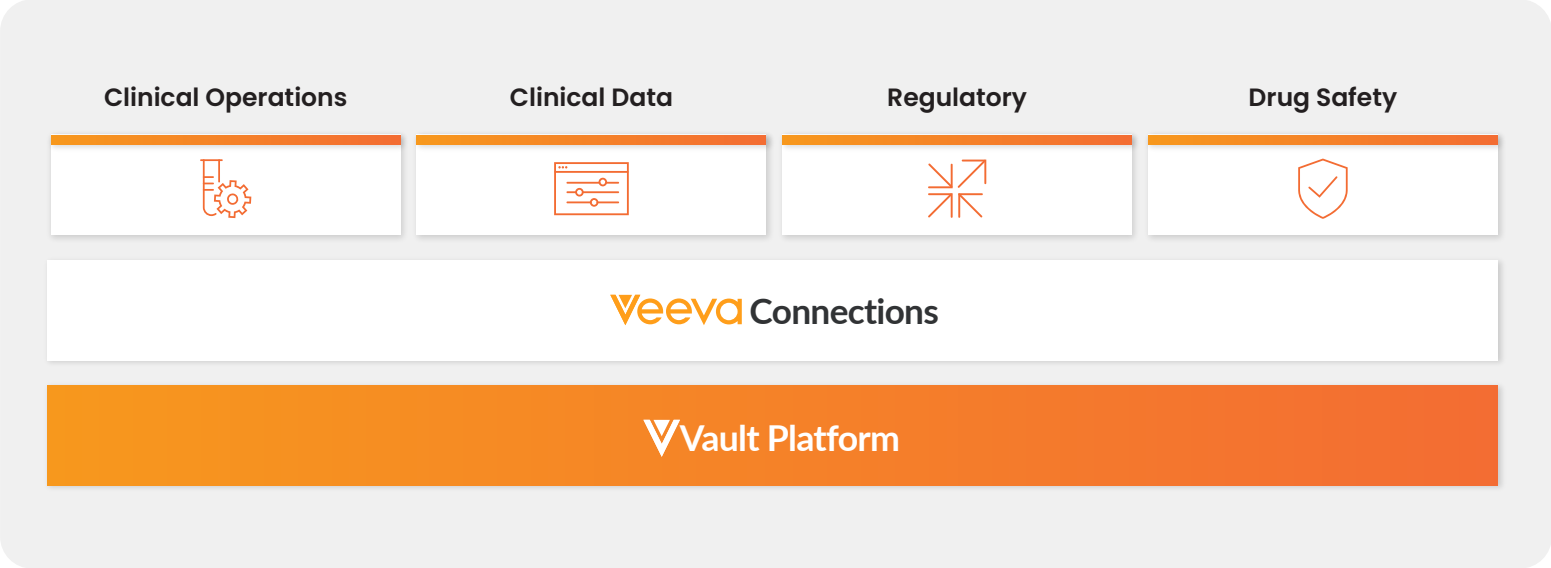
Veeva Development Cloud

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 across clinical, regulatory, and safety Vaults. Designed to streamline cross-functional business processes, the Connections break down silos and provide greater visibility, and automate manual tasks. View the Veeva Connections Resource Hub for a full list of available Veeva Connections.

Veeva Clinical Operations

Simplify and standardize clinical trial execution.

Veeva Clinical Operations unifies clinical systems and processes on a single cloud platform to enable end-to-end trial management.

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 Disclosures

Centralize clinical trial disclosures.

Veeva OpenData Clinical

Provide investigator and site data.

Veeva Clinical Data

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

Veeva Clinical Data 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 RIM

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

Unify regulatory systems and processes on a single cloud platform for end-to-end submission and registration management.

Veeva Registrations

Track global registrations.

Veeva Submissions

Plan and manage submissions.

Veeva Submissions Publishing

Publish to health authorities.

Veeva Submissions Archive

Access submissions history.

Veeva Safety

Safety suite of applications operate as a unified pharmacovigilance system on a single cloud platform to maximize operational efficiencies and improve patient safety.

Veeva Safety

Real-time management and oversight for adverse events.

Veeva Safety Workbench

Run advanced reports, queries, and analytics.

Veeva SafetyDocs

Centrally manage pharmacovigilance processes and content.

Veeva Safety Signal

Reliably detect, validate, and manage potential safety signals.