PRODUCT SHEET

Veeva Safety

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

Safety applications share a common data model which enables the end-to-end management of safety processes.

Veeva Safety is a global safety case intake, processing, and reporting system.

Veeva SafetyDocs manages safety-related content and processes including the pharmacovigilance master file, pharmacovigilance agreements, risk management plans and risk minimization measures, aggregate reports, literature articles, and safety signal investigations.

Veeva Safety Workbench is an advanced reporting tool designed for handling large volumes of data and complex analyses.

Veeva Safety Signal is for signal detection using statistical methods on data from Veeva Safety and standard industry sources. Signal detection activities can be tracked and automated using alerts and workflows.

PRODUCT	ANNOUNCED	STATUS	CUSTOMERS
Veeva Safety	2019	Mature	51–100
Veeva SafetyDocs	2019	Mature	11–50
Veeva Safety Workbench	2023	Early	1-10
Veeva Safety Signal	2023	Early	1-10

PRODUCT SHEET

Veeva Safety

Veeva Safety is a modern individual case safety report (ICSR) management system that manages the intake, processing, and submission of adverse events for clinical and post-marketed products.

Within one system, sponsors and CROs process and manage global and domestic adverse events for drug, biologic, vaccine, device, and combination products. Built-in gateway connections and reporting rules streamline submissions to health authorities and distributions to partners.

Central coding dictionary management automates semi-annual MedDRA, WHODrug, and EDQM updates.

Announced	2019
Status	Mature
Customer type	Enterprise Pharma, Biotech, CRO
Customers	51–100
Platform	Veeva Vault
Integrations	Lives with SafetyDocs Connected with Clinical Operations, EDC, RIM, and MedInquiry

PRODUCT SHEET

Veeva SafetyDocs

Veeva SafetyDocs manages pharmacovigilance-related content and processes.

Improve compliance, enable collaboration, and increase efficiency in the management of pharmacovigilance system master files (PSMF), pharmacovigilance agreements (PVAs), risk management plans (RMPs) and additional risk minimization measures (aRMMs), aggregate reports, literature articles, and safety signal investigations.

Announced	2019
Status	Mature
Customer type	Enterprise Pharma, Biotech, CRO
Customers	11–50
Platform	Veeva Vault
Integrations	Lives with Safety

PRODUCT SHEET

Veeva Safety Workbench

Veeva Safety Workbench is an advanced reporting and data analysis tool for high volumes of data from Veeva Safety. It is seamlessly integrated with Veeva Safety to ensure performant and reliable data transfers.

Design complex ad hoc reports with an intuitive user interface and explore the data using Dashboards and Case Series. Industry standard aggregate reports are included, with the ability to configure these to suit specific customer needs.

Announced	2023
Status	Early
Customer type	Enterprise Pharma
Customers	1-10
Platform	Veeva Vault
Integrations	Requires Veeva Safety

PRODUCT SHEET

Veeva Safety Signal

Veeva Safety Signal enables advanced signal detection using disproportionality and other methods on a variety of industry data sources such as FAERS, VAERS and EVDAS. Use Case Series to explore the underlying case data. It is seamlessly integrated with Veeva Safety to ensure performant and reliable data transfers.

Improve efficiency by automating signal detection activities using alerts and workflows and configure individual case reviews directly in Veeva Safety.

Announced	2023
Status	Early
Customer type	Enterprise Pharma
Customers	1-10
Platform	Veeva Vault
Integrations	Requires Veeva Safety