PRODUCT SHEET

Veeva Quality Cloud

Veeva Quality Cloud brings together QA, QC, and training on a single cloud platform to enable efficient quality management.

Quality applications exist on a single platform, allowing for the consolidation of quality processes that have been traditionally managed in siloed, disparate systems.

Veeva QualityDocs is the industry-leading GxP quality content management application.

Veeva Station Manager is a simple tablet-based application that ensures the right content is available 24/7 for operators on the manufacturing floor.

Veeva QMS manages and tracks quality processes such as deviations, complaints, CAPAs, change controls, audits, and risk management.

Veeva Batch Release aggregates data and content from QMS, LIMS, ERP, and regulatory systems to facilitate GMP release and market-ship decisions.

Veeva Validation Management is a paperless validation solution that optimizes the end-to-end validation lifecycle process.

Veeva Product Surveillance supports MedTech post-market surveillance processes and fully automated adverse event reporting to global health authorities.

Veeva Training is a GxP-optimized learning management system (LMS) that manages authoring, approval, assignment, and assessment of training materials in one place.

Veeva LearnGxP is an eLearning library of accredited courses and microlearning videos to help organizations meet regulatory requirements and drive personnel development.

Veeva LIMS is a laboratory information management system that facilitates quality control activities for GMP manufacturing.

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PRODUCT	ANNOUNCED	STATUS	CUSTOMERS
Veeva QualityDocs	2013	Very Mature	100+
Veeva Station Manager	2018	Mature	11–50
Veeva QMS	2016	Very Mature	100+
Veeva Batch Release	2023	Early	1–10
Veeva Validation Management	2021	Mature	11–50
Veeva Product Surveillance	2020	Early	1–10
Veeva Training	2018	Mature	100+
Veeva LearnGxP	2016	Mature	100+
Veeva LIMS	2021	Early	1–10

PRODUCT SHEET

Veeva QualityDocs

QualityDocs is a regulated quality content management solution. It is based on Veeva's proprietary GxP content reference model which enables best practices and standardization. The application manages content throughout its entire lifecycle, from creation to disposal. It allows internal and external parties to collaborate and share information, such as quality agreements and batch-related documentation, in a controlled manner directly within the system.

Announced	2013
Status	Very Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech
Customers	100+
Platform	Veeva Vault
Integrations	Lives with Training, QMS, LIMS, Validation Management, Station Manager, Product Surveillance

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PRODUCT SHEET

Veeva Station Manager

Station Manager is a tablet-based application that provides manufacturing operators access to content from QualityDocs. As a foundational element of business continuity planning, Station Manager syncs with QualityDocs when internet connectivity is available to ensure the latest versions of content are accessible even when offline. Station Manager surfaces only the relevant work instructions and procedures for a specific station in manufacturing to ensure operators readily have access to the content they need.

Announced	2018
Status	Mature
Customer type	Enterprise Pharma, Biotech, CDMO, MedTech
Customers	11–50
Platform	Veeva Vault
Integrations	Requires QualityDocs Lives with QualityDocs

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PRODUCT SHEET

Veeva QMS

QMS is a system designed to manage life sciences-specific quality processes. It provides best practices for handling complaints, deviations, audits, and change control. It also allows external partners to access the system to collaborate on investigations, audit findings, and corrective actions.

QMS is unified with other Quality Suite applications and connected to Registrations to enable coordination of product change control activities. Additionally, QMS, Batch Release and LIMS bring together quality assurance and quality control processes in one platform.

Announced	2016
Status	Very Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech
Customers	100+
Platform	Veeva Vault
Integrations	Lives with QualityDocs, Training, LIMS, Validation Management, Batch Release, Product Surveillance Connected with Registrations

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PRODUCT SHEET

Veeva Batch Release

Batch Release is an end-to-end solution that automates aggregation, reviews, and traceability of batch-related data and content to enable faster, more confident GMP release and market-ship decisions.

Batch Release brings together data and content from QMS, LIMS, ERP and Veeva RIM to simplify collaboration with external partners. It is tightly integrated with and requires QMS and QualityDocs. When used with LIMS and RIM, it offers faster time to value, but can be implemented with third party solutions and adjacent paper processes.

Announced	2023
Status	Early
Customer type	Enterprise Pharma, Biotech, CDMO
Customers	1–10
Platform	Veeva Vault
Integrations	Requires QMS and QualityDocs Lives with QMS, QualityDocs, LIMS Connected with Registrations

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PRODUCT SHEET

Veeva Validation Management

Validation Management streamlines commissioning, qualification, and validation activities across computerized systems, facilities, utilities, equipment, and processes. It facilitates the tracking of system inventory, the management of requirements, and oversight of project deliverables. Validation activities can be easily created and approved, test scripts executed digitally, and traceability and summary reports generated automatically.

Validation Management is unified with QualityDocs and QMS to connect quality events and key artifacts.

Announced	2021
Status	Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech
Customers	11–50
Platform	Veeva Vault
Integrations	Lives with QualityDocs, QMS, Training, LIMS, Product Surveillance

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PRODUCT SHEET

Veeva Product Surveillance

Product Surveillance helps with medical device post-market surveillance processes and adverse event reporting to global health authorities including the FDA, European Commission, and Health Canada. It standardizes and consolidates complaint reportability through a global decision tree and helps manage reporting timelines to ensure compliance across different health authorities. Quality and regulatory teams can allocate resources and prioritize submissions more effectively.

Product Surveillance also supports fully automated reporting with built-in XML payload generation and electronic data interchange (EDI) gateway. It is unified with core quality processes in QMS.

Announced	2020
Status	Early
Customer type	MedTech
Customers	1–10
Platform	Veeva Vault
Integrations	Requires QMS Lives with QualityDocs, Training, QMS, LIMS, Validation Management

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PRODUCT SHEET

Veeva Training

Training is a learning management system (LMS) designed for GxP compliance. It gives customers tools to manage learning content and curricula, and to deliver and track assignments. Training administrators can build curricula with many training types, including documents, videos, eLearning, classroom training, on-the-job training, assessments and more. Managers can track qualification and compliance status using reports and dashboards.

Training is unified with QualityDocs, ensuring access to source content and automating re-training based on changes. Document and training data are easily combined for reporting, and training assignments can be made in QMS workflows.

Announced	2018
Status	Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech
Customers	100+
Platform	Veeva Vault
Integrations	Requires QualityDocs Lives with QualityDocs, QMS, LIMS, Validation Management, Product Surveillance Connected with LearnGxP

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PRODUCT SHEET

Veeva LearnGxP

LearnGxP is an accredited training library that can be deployed with Veeva Training or other learning management system. The library contains interactive eLearning content and microlearning videos on topics such as the Fundamentals of Good Manufacturing Practices, Data Integrity, and Inspection Readiness. It is designed to help life sciences companies meet regulatory compliance requirements and provide professional development on industry-specific topics to their workforce.

Announced	2016 (acquired in 2021)
Status	Mature
Customer type	Enterprise Pharma, Biotech, CRO, CDMO, MedTech, Consumer
Customers	100+
Platform	N/A
Integrations	Requires Training or other learning management system

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PRODUCT SHEET

Veeva LIMS

Veeva LIMS enables Quality Control to optimize batch release testing, stability study management, and environmental monitoring. It drives digital method execution, specification adherence, and review by exception to accelerate the release of product.

LIMS promotes compliance by verifying user qualifications from Veeva Training, displaying effective test method procedures from QualityDocs, and initiating lab investigations directly in QMS from out-of-specification (OOS) results.

Announced	2021
Status	Early
Customer type	Enterprise Pharma, Biotech, CDMO, MedTech
Customers	1–10
Platform	Veeva Vault
Integrations	Lives with QualityDocs, QMS, Training, Validation Management, Batch Release, Product Surveillance

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