Veeva MedTech

PRODUCT SHEET

Veeva Quality

Veeva Quality 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 nonconformances, complaints, CAPAs, change controls, audits, risk management & SQM.

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 digital solution that unifies data across the quality ecosystem to optimize the 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 compliance-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 facilitates quality control activities for GMP manufacturing by unifying QC data management and test execution processes within a single, cloud application.

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

Veeva MedTech

PRODUCT SHEET

Veeva QualityDocs

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

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

Veeva MedTech

PRODUCT SHEET

Veeva Station Manager

Station Manager is a tablet-based application that provides manufacturing operators access to content from QualityDocs. As a supportive element to 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	Medtech, Enterprise Pharma, Biotech, CDMO
Customers	11–50
Platform	Veeva Vault
Integrations	Requires QualityDocs Lives with QualityDocs

Veeva MedTech

PRODUCT SHEET

Veeva QMS

QMS is a cloud-based quality management system designed to manage life sciences-specific quality processes. It provides faster time to value with streamlined processes for handling complaints, nonconformances, audits, quality risk management, supplier quality management, and change control. It also allows external partners to access the system in real time to collaborate on investigations, audit findings, corrective actions, supplier change control and more.

QMS is unified with other Quality Cloud applications, and connects to RIM to coordinate product change control activities, Safety to manage complaints, and CTMS for study-related data used in QMS and protocol deviation management.

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

Veeva MedTech

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 QualityDocs, QMS, LIMS, ERP and 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 to centralize all batch release data.

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

Veeva MedTech

PRODUCT SHEET

Veeva Validation Management

Validation Management is a digital solution for faster, accurate validation. It 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 deliverables.

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

Veeva MedTech

PRODUCT SHEET

Veeva Product Surveillance

Product Surveillance supports 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 manages reporting timelines to ensure compliance.

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

Veeva MedTech

PRODUCT SHEET

Veeva Training

Training is a learning management system (LMS) designed to ensure training compliance, qualification, and job readiness. It gives customers tools to manage learning content and curricula, and to deliver and track assignments. Training administrators can build plans 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	Medtech, Enterprise Pharma, Biotech, CRO, CDMO
Customers	100+
Platform	Veeva Vault
Integrations	Requires QualityDocs Lives with QualityDocs, QMS, LIMS, Validation Management, Product Surveillance Connected with LearnGxP

Veeva MedTech

PRODUCT SHEET

Veeva LearnGxP

LearnGxP is an accredited training library that can be deployed with Training or other learning management systems. 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	Medtech, Enterprise Pharma, Biotech, CRO, CDMO, Consumer
Customers	100+
Platform	N/A
Integrations	Requires Training or other learning management system

Veeva MedTech

PRODUCT SHEET

Veeva LIMS

LIMS optimizes batch release testing, stability study management, and environmental monitoring for the quality control lab. It drives detailed sample management, digital test method execution, specification adherence, and review by exception to accelerate the release of product.

LIMS promotes compliance by verifying user qualifications from Training, displaying effective test method procedures from QualityDocs, and initiating quality events directly in QMS, ensuring proper resolution prior to batch disposition.

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