

PRODUCT SHEET

Veeva Quality Basics

Veeva Quality Basics brings together quality processes, content, and training management into a single platform. Where you are managing your documents, you can also assign and track training, as well as document quality events such as Change Controls, Deviations, Audits, and Supplier Quality Management (SQM).

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

Veeva QMS manages and tracks core quality processes such as audits, deviations, change controls, standalone CAPA, and supplier quality management.

Veeva Training is an industry-specific LMS ensuring that your team is ready to perform their jobs while enabling GxP compliance and audit readiness.

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

Veeva LIMS Basics

Veeva LIMS Basics empowers virtual life science companies to digitize and scale all QC and Stability functions. It centralizes specifications and external results for consistent data, compliance, and accelerated batch release.

Veeva RIM Basics

Veeva RIM Basics provides an end to end solution for managing regulatory submissions. It helps organizations streamline the complex process of preparing, managing, submitting, and maintaining regulatory documentation required by global health authorities.

Veeva Submissions is used to plan, collaboratively author, review, and approve regulatory documents. It includes dashboards and reports to allow submission managers to track the status of documents in real time.

Veeva Submissions Archive is a secure repository of submission-published output and includes a viewer for easy access to historical applications submitted to health authorities.

Veeva Submissions Publishing streamlines the assembly, formatting, and publishing of regulatory submissions, ensuring compliance with health authority requirements.

Veeva Clinical Basics

Veeva Clinical Basics unifies eTMF and CTMS in a single system, providing seamless collaboration across clinical trial operations and document management. This ensures that study teams can manage trial activities, monitor subject recruitment, and maintain a complete, inspection-ready TMF—enhancing efficiency and compliance throughout the clinical trial lifecycle.

Veeva eTMF is the leading trial master file application used to ensure the quality, timeliness, and completeness of a TMF. Veeva eTMF Basics supports both active management, archival, and the transfer of a TMF from a Veeva CRO.

Veeva CTMS is a trial management system that provides end to end study management and study capabilities.

Veeva Commercial Content Basics

Veeva PromoMats provides an end-to-end MLR review and digital asset management solution for managing compliant promotional content.

Veeva Medical Basics

Veeva MedComms is a unified source of truth for medical content. It streamlines the creation, approval, and distribution of scientific materials, ensuring Medical Affairs teams deliver consistent, compliant responses globally.

Veeva CRM Basics

Veeva CRM for Medical is a purpose-built engagement platform designed specifically for Medical Science Liaisons (MSLs). It empowers medical teams to foster deep, scientific relationships with Key Opinion Leaders (KOLs) by capturing rich interaction data and providing the insights needed to drive meaningful scientific exchange.