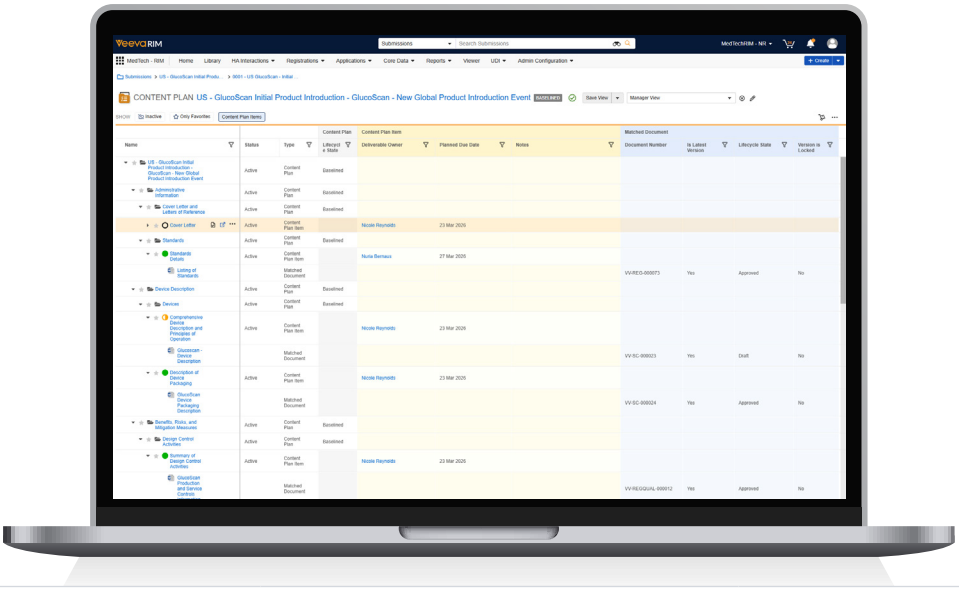
VRIM

Submission Content Plans

Simplify submission content planning, management, and compilation to save significant time and effort.

Veeva RIM Submission Content Plans (SCPs) provide a unique, structured, and collaborative approach to planning, tracking, and managing submission development. As the volume of submissions continues to grow, regulatory teams need a way to track submission development across multiple markets. Many companies are using multiple tools for this work, including Excel, Microsoft Project, emails, and file shares. With SCPs, teams can efficiently plan, track, and develop all content required for submissions in a single, collaborative space.



Benefits

Increase Submission Efficiency

Streamline communication and task delegation with a single source of truth, while reducing duplicative tasks and manual data entry with templates and automation.

Track Global Submission Statuses

Gain a holistic, real-time, view of individual document and submission progress across submission teams and markets.

Enhance Compliance

Improve global traceability of submitted documents, including what versions were submitted to which Health Authorities, and where to find the latest source documents when it's time to make an update.

Features

Content plan templates

Leverage our recommended constraints to standardize submission planning by submission type, product, or region.

Content plan viewer

Plan submissions, collaborate on content assembly, and track progress to completion using a holistic view that can be easily tailored to each user's preference.

Advanced document matching

Reduce manual tasks with Veeva's most recent improvements to automated and suggested matching for documents.

Drag-and-drop documents

Drag and drop Veeva RIM documents from your library, or external files from your desktop for automated assignment and document upload.

Real-time reports and dashboards

Automate status reporting on progress, tasks, and timelines with real-time dashboards and reports. Easily report on which documents have been submitted to which Health Authorities.

Foundation for continuous innovation

SCPs are a prerequisite for adopting more of Veeva RIM's high value features, including Active Dossier, Global Content Planning and Veeva Publishing.

Veeva RIM

SCPs are part of Veeva RIM, which streamlines global regulatory processes on a single, cloud-based platform. This enables medtech companies to:

- Ensure teams are developing reliable regulatory content with high data integrity
- Coordinate regulatory efforts across headquarters, affiliates, and partners
- Respond faster to changing regulations
- Increase end-to-end process efficiency from submission planning to publishing