Veeva Vault Medical Device & Diagnostics

Strategic Regulatory Platform for Medical Device & Diagnostics

Can you Access, Search and Manage your Regulatory Data in a Single, Connected System?

Medical device and diagnostic regulatory teams face ever-increasing volumes of information that needs to be managed—submissions, global registrations, post-market data, and promotional materials. But their systems are siloed, disconnected, and antiquated, stifling efficiency and increasing compliance risk.

Most regulatory affairs professionals lack modern tools to address the needs of today's global regulatory environment and end up with challenges like these.

Separated Content and Data

Prevents traceability & audit readiness

Disconnected Technology

Inhibits collaboration across teams

Lack of Visibility & Control

Increases compliance risk and inefficiency

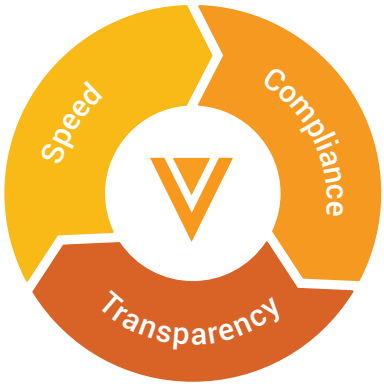
Good News. It can be.

Regulatory teams across the organization should be connected throughout the product lifecycle by maintaining a single source of truth that feeds critical commercial processes.

Using the **Veeva Vault RIM Suite and Vault PromoMats applications**, regulatory and commercial teams can work together to **ensure compliance** with approved labeling, **shorten regulatory review times**, and bring a **consistent, compliant message** to the market.

Vault RIM Suite

Authoritative source for regulatory information management



Vault PromoMats

Regulated marketing management to ensure end-to-end compliance

Veeva Vault Medical Device & Diagnostics

Better Together: Vault RIM Suite + Vault PromoMats

Purpose-built for medical device and diagnostics companies, the Vault RIM Suite and Vault PromoMats bring regulatory and marketing processes together with a single source of truth for labeling, intended use, and approved claims.

Vault RIM Suite + Vault PromoMats =

Insights with Real-time Data
Immediately see submission status, product registration, approved content and more

Centralized Repository
Mitigate compliance risk and shorten review times by maintaining a Single Source of Truth across the organization

Simplified Search
Reduce time & effort by searching content and data with a single query across unified systems

US & EU Ready
Handle 510(k), PMA, EU MDR, and IVDR requirements with confidence

Digital Publishing
Publish and withdraw approved assets to any digital channel from single source of truth

Cross-Linked Systems
Utilize key regulatory documents as references and claims substantiation

Built-In AI
Shorten timelines with auto claims-linking and continuous innovative improvements

Reduced Maintenance
Lessen IT burden by utilizing common platform

To learn more, visit veeva.com/MedDevice