

Scalable Compliance and Literature Monitoring Worldwide

A Top 3 Med Device company implemented ReadCube to modernize regulatory workflows and unify literature monitoring across product lines and regions. By replacing siloed systems with a centralized platform, the company reduced review cycle times, improved audit readiness, and ensured global teams could access the content they needed faster and with greater consistency.



The Opportunity

With increasing complexity across divisions and therapeutic areas, this leading Med Device organization saw a strategic opportunity to align its literature workflows under a single, scalable system. The goal was to simplify regulatory reporting, reduce duplication, and create traceability across reviews.

By consolidating tools and enabling real-time collaboration, the company could respond to regulatory requirements more efficiently while improving compliance confidence across regions.

The Solution

ReadCube provided an intelligent infrastructure for global regulatory operations.

- ✓ Centralized libraries ensured consistent access and eliminated tool silos
- ✓ AI-powered tagging and alerts accelerated discovery and reduced oversight risk
- ✓ Built-in version control and stakeholder mapping improved audit readiness
- ✓ Automation reduced manual workload and enabled cross-functional efficiency

The Results:

100s

of libraries
unified across global
regulatory and review teams

1

scalable platform
supports insight generation
across business units

**Significant
efficiency gains**

by reducing manual review
and reconciliation effort