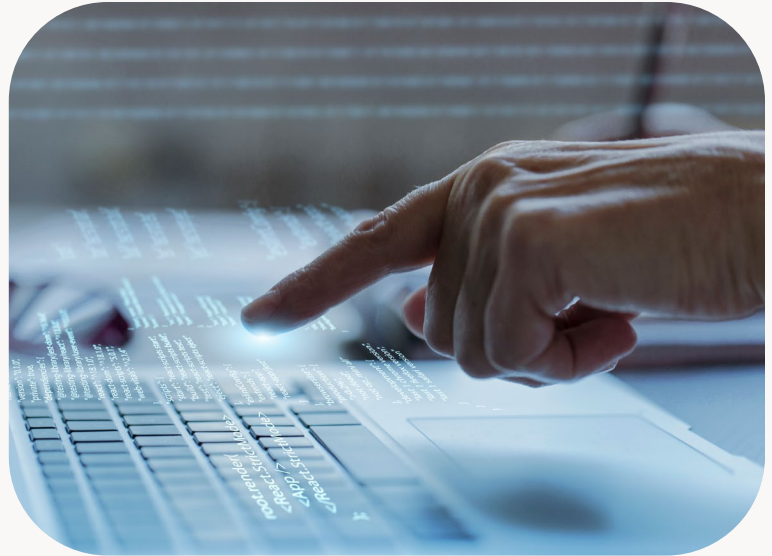


Global Consistency with Smarter Literature Review

This Top 3 Med Device company wanted to enhance regulatory operations with a centralized platform for literature monitoring. ReadCube helped reduce manual effort, accelerate insight generation, and support audit readiness. Regulatory teams now manage content more consistently across therapeutic areas and global regions.



The Opportunity

The Med Device manufacturer saw a clear opportunity to unify its approach to literature review across regulatory teams worldwide. With growing global requirements and expanding product lines, the company aimed to simplify review workflows, strengthen audit trails, and improve consistency across compliance submissions.

By streamlining how content was accessed, tagged, and shared, the company could free up expert capacity while accelerating insight generation and decision-making.

The Solution

The company adopted ReadCube to modernize regulatory literature workflows.

- ✓ Automated tagging improved search precision and review speed
- ✓ Shared libraries supported collaboration across global teams
- ✓ Alerts ensured teams stayed up to date with new publications
- ✓ Built-in version control enhanced audit readiness and traceability

The Results:

30%

improvement
in article data extraction
accuracy

5,250

hours saved
annually across users
worldwide

350

global users
aligned under one regulatory
content platform