

Drive innovation and performance in regulatory information management (RIM)

Revolutionize the product life cycle and accelerate time to market with AI-powered automation



Shift RIM into high gear with intelligent automation

Bringing innovative therapies and wellness-enhancing products to market can feel like a race to the finish line in the life sciences industry, especially when it comes to regulatory information management (RIM).

Don't let speed bumps like clinical data complexity, strict regulatory compliance requirements, and maintaining clean, current electronic master files (eTMF) stop product development in its tracks. Transform your approach to RIM with AI-powered automation and deliver lifesaving products to market in record time.

Rev up your R&D

Intelligent automation within RIM paves the way for more powerful cross-functional processes, including manufacturing, R&D, clinical trials, pharmacovigilance, and quality assurance.

- **Clinical data management:** Enhance research reliability and efficiency by automating data capture across different sources for consistent accuracy and accessibility.
- **Clinical reporting and audits:** Improve regulatory compliance and streamline the audit process using AI-driven software to generate reports and flag inconsistencies.
- **Clinical supply chain:** Optimize inventory management, track shipments, and forecast future supply needs with AI-powered automation and analytics.

Better than cruise control

Boost the entire product life cycle with intelligent automation benefits that go beyond the basics.

- **Enhanced operational efficiency:** Boost speed and accuracy and reduce errors by automating activities like data entry and compliance tracking.
- **Faster time to market:** Simplify processes and reduce the time required to prepare and submit regulatory documents for rapid turnaround.
- **Improved employee satisfaction:** Free employees from routine, repetitive tasks so they can focus on more meaningful, strategic goals.
- **Better patient outcomes:** Provide patients with the best possible treatments using intelligent automation to enhance RIM processes.

Automation by the numbers



3 million

documents automatically authored, archived, and migrated to a new RIM in one year

31,250

working hours per year saved by automating emailed invoices

150 hrs → 2 hrs

Total time for clinical trial data summarization and submission before and after automation

From manual to automatic

Deploying intelligent automation in your RIM system takes your processes from slow to seamless.



Automated audit trails

Audits are critical for patient safety, but collecting information from different platforms takes significant time. Automating audit trails and report generation within the RIM system leads to:

- **90% reduction** in processing time
- **\$570K** annual savings
- **Stronger regulatory compliance**



Binder creation

Binder creation is a key step in the submission application process, but it's a time-consuming effort. An automated RIM system library for binder creation results in:

- **90% reduction** in processing time
- **\$435K** annual savings
- **Shorter time to market**



Regulatory submission document creation

Creating the documents required for a drug submission is integral to gaining regulatory approval. Automating the submission document creation process provides:

- **90% reduction** in processing time
- **\$630K** annual savings
- **Faster drug submissions**



The right copilot for your RIM

AI-powered automation turbocharges your RIM system, but you need an experienced copilot to help steer you towards the finish line. The right technology partner supports you at every turn with advanced capabilities, life sciences industry expertise, and solutions tailored to your specific regulatory and operational needs.

UiPath and CGI offer a roadmap for intelligent automation across the product life cycle so you can:

- **Eliminate barriers to entry:** Full-featured automation platforms as a service allows you to only pay for what you need without substantial investments in infrastructure and hardware.
- **Drive faster ROI:** Shorter initial setup time and lower financial commitments means you'll experience the benefits of automation sooner – so you can get ahead of the competition.
- **Leverage scalability:** Become an industry leader by responding quickly to market demands, regulatory changes, and internal growth with an agile and scalable automation platform.

Put your RIM system on the fast track to success with UiPath and CGI.

Read [**Cut through the red tape: How AI-powered automation enhances regulatory information management to accelerate time to market**](#) to learn more.

