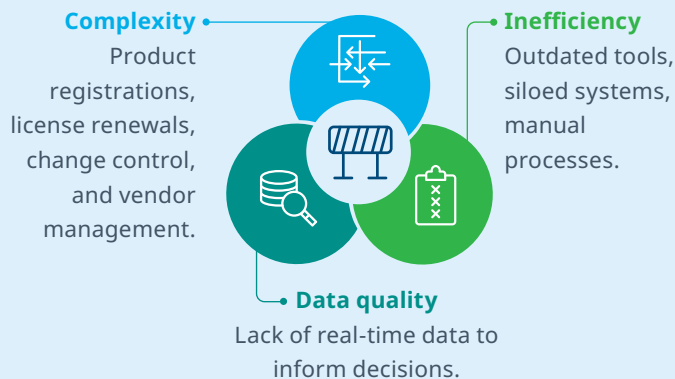


Streamline MedTech Regulatory Operations with SmartSolve® RIM

Your unified, intelligent solution for regulatory compliance activities





Challenges in MedTech regulatory operations



Solution

SmartSolve RIM consolidates product registration details, tracks license renewals, manages vendors, and supports change impact analysis.

Key features — SmartSolve RIM

-  **Centralized interface:** Access product applications, approvals, Health Authority correspondence, task tracking and submission status in one screen.
-  **Automation:** Create new applications, authorizations and submission content in seconds!
-  **Notifications and alerts:** Expiring licenses, health authority updates, drill-down reports.
-  **Vendor oversight:** Verification of third-party vendors and related SKU data.

Benefits of SmartSolve RIM

Comprehensive compliance
Reduce regulatory risk and operational complexity with traceable, centralized actions that simplify product registrations, license renewals and change control management.



Efficient change management
Streamline and automate the evaluation of regulatory changes across multiple applications and jurisdictions.

Seamless collaboration
Improve decision-making and eliminating data silos with cloud-based access and accurate, up-to-date information.

About SmartSolve®: SmartSolve is an AI-enabled, Microsoft Azure-based platform that helps Life Sciences organizations streamline and automate global quality management and regulatory compliance. [SmartSolve® eQMS](#) centralizes enterprise-wide quality processes, from design and manufacturing to post-market surveillance, while [SmartSolve® RIM](#) manages regulatory submissions, product registrations, and health authority interactions. Built on industry best practices, SmartSolve connects teams, data, and workflows in a single platform to drive an optimized focus on patient safety, product quality and commercial performance.

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