Fact Sheet

SmartSolve® RIM Submission Management

Intelligence. Automation. Integration.

As global regulations continue to advance and drive complexity, medical device, in-vitro diagnostic, and pharmaceutical companies require digitized solutions that are fit for purpose for their industry and drive transparency and acceleration of global product approvals and timely market registrations.

Situation

Rising cost pressures, evolving and divergent regulations, and elevated internal and external pressures drive organizations to leverage technology to accelerate market access activities whilst improving commercial performance. These innovations enable teams to author source content and prepare submission dossiers more efficiently while ensuring full compliance with the applicable requirements of regional health authorities.

Our solution

SmartSolve RIM Submission Management provides the industry's most advanced solution for **managing medical device, in-vitro diagnostic, and pharmaceutical regulatory product registration activities globally.**

SmartSolve RIM delivers a single, fully integrated, AI enabled solution developed on cloud architecture to ensure scalable, effective and compliant management of regulatory applications, positioning organizations to complete regulatory activities more efficiently and cost-effectively.

With SmartSolve RIM Submission Management, organizations can **efficiently assess, plan, forecast and coordinate the end-to-end processes** involved in assembling, publishing and dispatching regulatory dossiers to international health authorities within a single integrated platform to drive timely market access of global healthcare solutions.

SmartSolve® RIM Submission Management

Built with IQVIA's deep industry knowledge and experience, led by a large team of professionals with practical business and technology competence

Supports the entire regulatory submission management lifecycle, from early investigational stages to marketing and post-approval activities

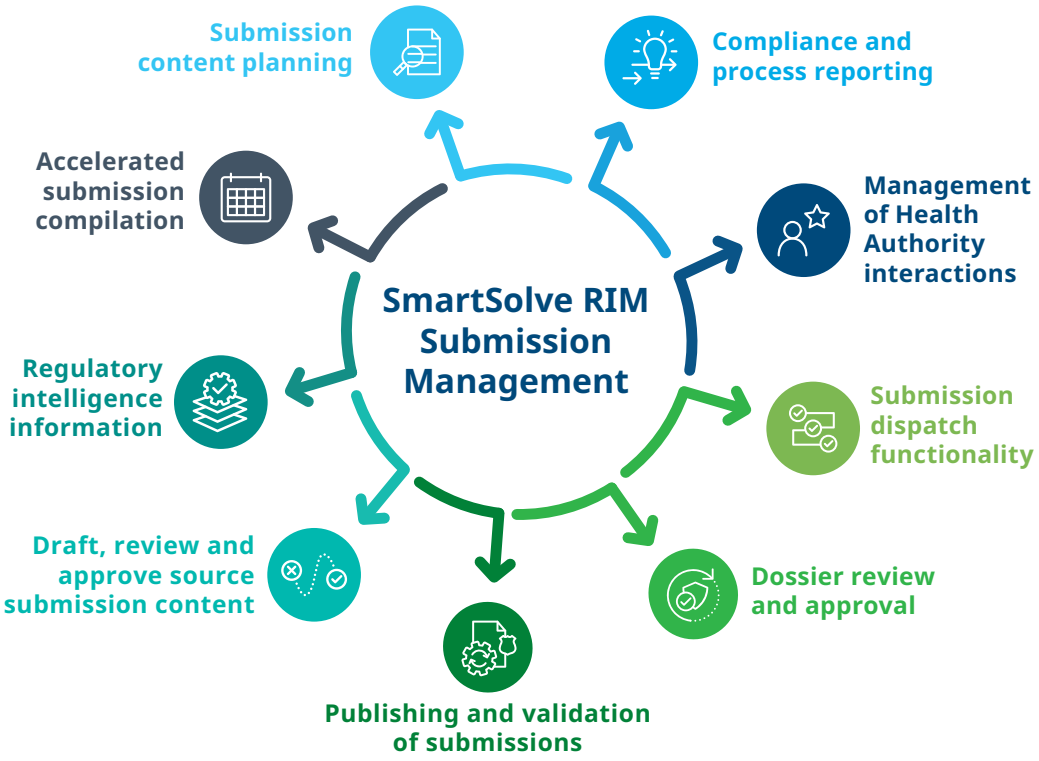
Enables full transparency of actions, data in use and project timelines across all involved regulatory groups

Supports all mandatory pharma (eCTD, NeeS) and MedTech (IMDRF, eSTAR and ASEAN CSDT formats) submission formats

Intuitive and easy to use, enabling efficient tasks completion with familiar tools and functions throughout the system

Drives operational efficiency enabling re-use of integrated PDF tools across workstreams, countries and regions

SmartSolve RIM Submission Management improves submission process efficiency, regulatory professional productivity and accelerates global market approvals.



SmartSolve RIM Submission Management streamlines global regulatory submissions with end-to-end planning, seamless system integration, and efficient dispatch. It supports Pharma and MedTech with ready-to-use templates, centralized oversight, and real-time visibility — helping teams accelerate approvals and stay compliant worldwide.

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.

IQVIA TECHNOLOGIES

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