

# IQVIA RIM Smart

*Streamline your regulatory processes with our end-to-end regulatory information management system, IQVIA RIM Smart.*



Free your team from labor-intensive maintenance to focus on optimizing the availability of global healthcare solutions. RIM Smart can help your organization streamline the global regulatory approval processes, optimize resource utilization, and guarantee the accuracy of information that underpins regulatory compliance activities.

## The challenges of regulation

Regulatory organizations across the globe are dealing with a significant burden: maintaining and managing a vast amount of information across many diverse sources.

From strategy to submission data, organizations have to face the challenge of how to efficiently gather, analyze, and disseminate regulatory information – information that underpins crucial regulatory and business strategies.

At the same time, disparate systems and complex processes increase the risk for error. What's more, constant changes in global health authority requirements and resource constraints make the task of putting regulatory data to work even harder.



# How to take the hard work out of compliance

RIM Smart enables organizations to remain focused on what matters most: getting safe, effective products to market and keeping them there.

RIM Smart puts seven key components to work, addressing the most common and high-volume regulatory activities, including product and registration tracking, submission planning and publishing, health authority correspondence logging, and administering regulatory information.

These feature-rich components can be implemented independently and users can add additional modules as they grow. However, when used together, they deliver a seamlessly integrated system that's built for automating and reusing regulatory data in the most efficient way possible.

Intuitive workflows provide visibility and accountability across the organization while cloud services ensure information is globally accessible and secure.

Managing the labeling process, such as handling artwork and translation, has never been easier when paired with RIM Smart Labeling.

RIM Smart has been designed by regulatory professionals to make the regulatory process as simple and as fast as possible, while helping organizations achieve the highest levels of compliance.

RIM Smart is the modern solution for life sciences organizations that want to effectively manage and maintain their regulatory information.

## Product and registration tracking

Detailed planning and tracking of full product registration lifecycle management capabilities including xEVMPD. IDMP ready.

## Publishing and validation of submissions

All formats for all regions available.

## Integrated dossier review and approval functionality

Including peer comments and communications.



## Key capabilities



## Embedded regulatory information database

Information for drugs, biologics, medical devices and IVDs across over 110 countries, regions, and international organizations.

## Submission content planning

Including task assignments and scheduling.

## Health authority interactions

Complete management of health authority interactions, including correspondence and commitment tracking.

## Labeling and content management

A complete labeling solution with dedicated content management.

# A single solution, optimized for both pharmaceutical and MedTech

One product designed for life sciences is RIM Smart, which was built to deliver for all pharmaceutical and MedTech organizations. For companies that operate in one or both markets, RIM Smart offers an integrated, automated and intelligent way of managing the complete regulatory lifecycle of pharmaceutical, medical device and combination products.

## Effective content and certificate management

RIM Smart smooths change control and resubmission processes by recording submission content, and can also make MDSAP and ISO 13485 audits and inspections easier to manage with approval certificate storage and translation using a built-in tool

### Streamlining query processing

To ensure ease of activity monitoring and management, enquiries from healthcare agencies, notified bodies, dealers and other third parties can be linked into registration submissions.

### Integrated document handling

Global submission builds can be handled quickly and simply with source documents that can be dragged and dropped from repositories into country-specific templates, with the option to integrate directly into IQVIA's SmartSolve eQMS.

### Global registration visibility and tracking

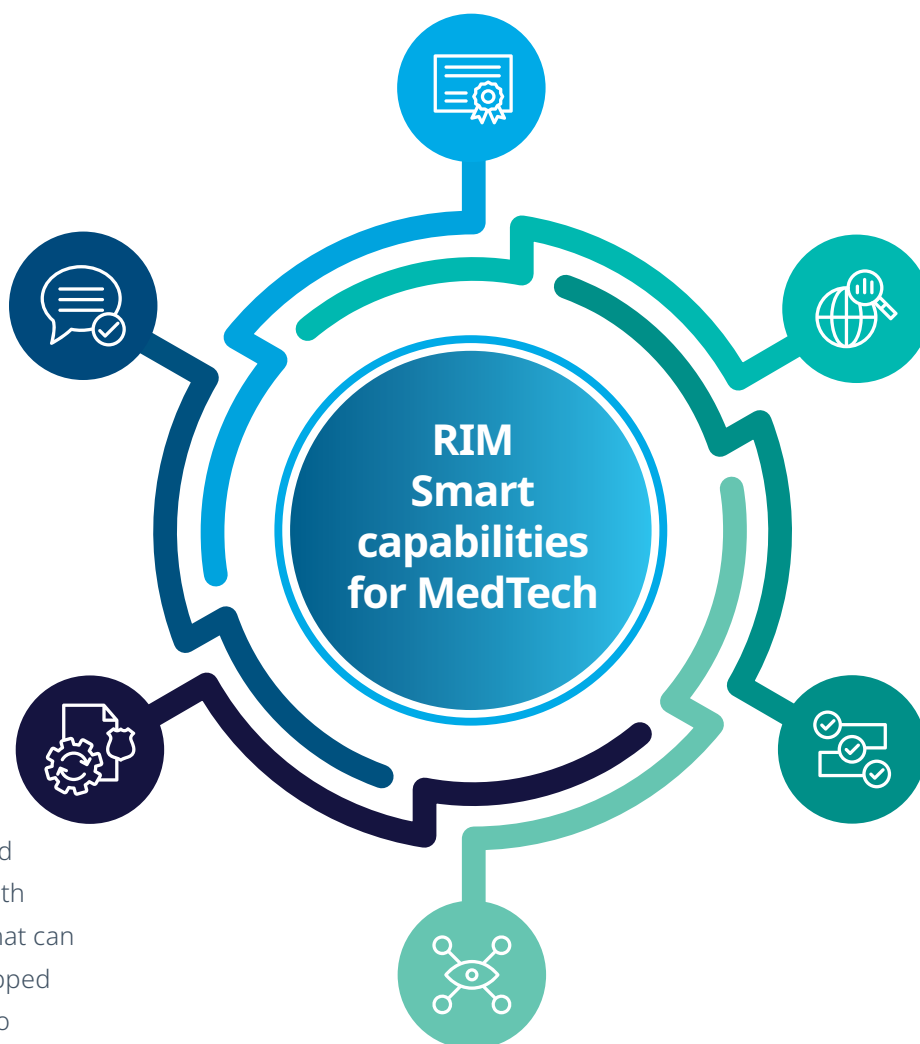
RIM Smart provides item country-level registration visibility across the globe, with registration status activity tracking at a country-submission level.

### Intelligent workflow

Easily manage tasks and commitments associated with global registration activity within RIM Smart, using a list or calendar format.



### Complete expirations oversight

RIM Smart helps maintain business continuity with a complete view of global registration and third-party license expirations.



# The benefits of RIM Smart

## RIM SMART ENABLES YOU TO:

-  Safeguard data accuracy and compliance with a single integrated system for structuring, managing, and reusing regulatory information across the product lifecycle.
-  Access and disseminate critical global regulatory data in a way that creates better upstream and downstream visibility across the organization.
-  Save time by automating the analysis and submission process for any requirement changes.
-  Increase focus on your core competencies: RIM Smart can help reduce resource efforts and cut maintenance activity spend.

# Our vision

IQVIA's vision is to empower your organization to focus its resources on what's important: bringing valuable products to market and keeping them there.

## Our vision is built on three core foundations:

<b>CONNECTED INTELLIGENCE</b>	Immersing users within an environment to provide complete awareness with actionable insights that are never more than one click away.
<b>ROBOTIC AUTOMATION</b>	With its intuitive automation, simplification and integration, RIM Smart streamlines processes, improves data accuracy, and delivers exponential increases in productivity.
<b>REGULATORY INSIGHT</b>	Uncovering strategic, regulatory, and compliance updates in real time, actively monitoring and alerting users to changes or emerging trends.

IQVIA's vision goes beyond just optimizing diverse quality, regulatory, and safety systems: it pioneers the transformation of compliance from a burden to a competitive advantage.



# The RIM Smart difference

## RIM SMART OFFERS:

-  Integration to IQVIA Regulatory Intelligence, providing global regulatory requirements and operational insights across the product lifecycle.
-  A complete end-to-end regulatory solution that supports product growth for organizations of all sizes.
-  A single source of truth for all your regulatory data and activities.
-  Intuitive submission management and publishing that enables rapid adoption across teams.
-  Smooth integration with quality systems including IQVIA SmartSolve® eQMS.

## CONTACT US

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