Infographic

IQVIA RIM Smart for MedTech

An integrated, automated registration management system

The regulatory landscape is undergoing significant transformation

Medical device and in vitro diagnostics (IVD) companies face rising global regulatory complexity and associated costs, directly impacting market access and regulatory compliance activities.

150%
Increase in regulatory mandates in the last 5 years[1]

Every **13 minutes**, a new or changed healthcare regulation or reference document is published somewhere in the world[2]

Over **8,000** new regulations and reference documents added per year[3]

1 IQVIA Regulatory Intelligence May 2022
2 IQVIA Regulatory Intelligence data — October 2022
3 IQVIA Regulatory Intelligence data — October 2022

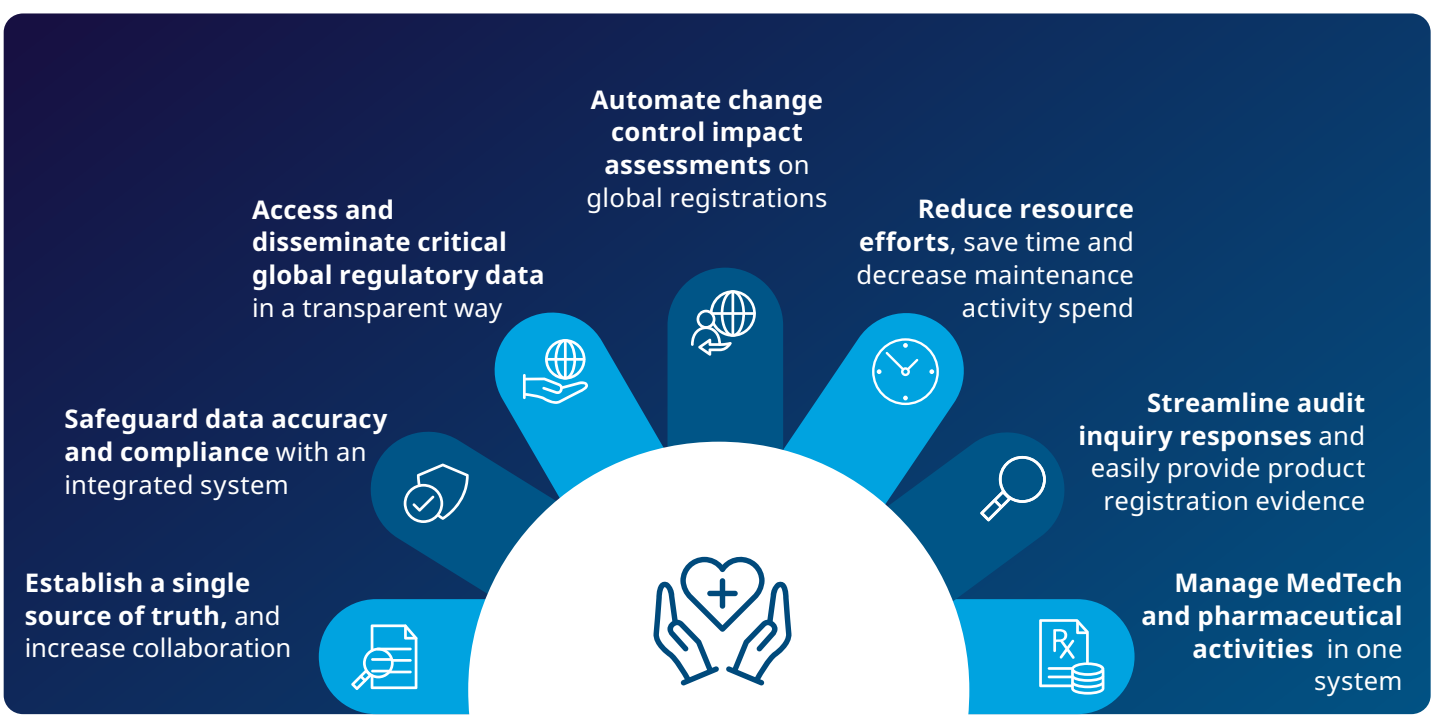
Companies face numerous challenges within the rapidly evolving and innovative medical device and IVD industry.

- Knowledge bottlenecks created by siloed information, decentralized systems, inefficient technology and resistance to change
- Complex and varied regulatory requirements that often differ by country and region
- Finding and retaining skilled regulatory professionals
- Significant risk exposure due to using shared drives and spread sheets to manage activities
- Manual and labor-intensive regulatory intelligence tasks

By adopting strategies and technology to address these challenges, you can better manage regulatory risks and ensure product safety and effectiveness.

The solution: IQVIA RIM Smart for MedTech

RIM Smart delivers a complete submission and registration solution, inclusive of variation activities, and brings transparency to global regulatory market access activities that drive commercial growth.



RIM Smart consolidates global submission and registration activities and data management into a single application, and offers an integrated, automated and intelligent way of managing the complete regulatory lifecycle.

Simplify content and certification management
Ease change control and resubmission processes, and simplify access to evidence needed in MDSAP and ISO 13485 inspections

Streamline query processing
Link inquiries from regulatory agencies, notified bodies and other third parties into registration submissions and post submission updates

Integrate document handling
Quickly address global submission builds by dragging and dropping source documents from repositories into country-specific templates

Global registration visibility and tracking
View item country-level registration across the globe, and track registration status activity at country, manufacturing site and local levels

Intelligent workflow
Easily manage tasks and commitments associated with global submission and registration activity

Complete expirations oversight
Maintain business continuity with a complete view of global registration and third-party license

RIM Smart's **seamless integration with IQVIA Regulatory Intelligence database** enables you to efficiently map regulatory tasks and dependencies across countries, simplifies impact assessment and helps ensure timely execution of remediation activities. When **combined with IQVIA SmartSolve eQMS, the result is an end-to-end change management solution** that allows you to evaluate potential outcomes of global registration changes and determine the impact before finalizing plans.

Drive commercial growth, increase productivity and improve registration data and content accuracy with IQVIA. **Learn more**

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