Infographic

IQVIA RIM Smart

Intelligent management of the complete regulatory lifecycle.

The situation

Managing the rising cost, volume and complexity of regulatory compliance is an increasing burden with no end in sight.

>2,000
new or modified regulations
released by the FDA since 1998

Source: PwC, as cited in "Disruptive Trends in Pharma Regulation" Pharmaboardroom.com, June 2018

53% of global life sciences CEOs
consider industry regulations a top disruptive business trend

Source: PwC, 21st Annual Global CEO Survey

The challenges

Key to your success is bringing together critical data, processes, technology and regulatory expertise in a comprehensive, sustainable approach to regulatory information management.

- Keeping up with new and changing regulations
- Disparate systems, complex processes
- Manual, high-touch, repetitive work
- High administrative burden of maintenance work
- Breaking down silos for timely access to information
- Regulatory demands in more countries

The solution

IQVIA RIM Smart.

Product and registration tracking
Universal, portfolio-wide multi-level tracking, reporting capabilities

Correspondence and commitments
Quickly, efficiently e-archive, track and respond to health authority communications

Submission planning
Preparation, workflow, management tools for dossier planning and resource, time management

IQVIA RIM Smart

Viewer / archive
Read, navigate eCTD, NeeS, other e-submissions in varying views

Publishing and validation
Create, edit and validate eCTD, NeeS and Paper submissions

Content management
Create compliant, submission-ready content (eCTD); synchronized with MDM; automated

Embedded RID
Up-to-date, country level reporting requirements

IQVIA RIM Smart delivers fully integrated, technology-led, intelligent management of the complete regulatory lifecycle so you can focus on what you do best – getting safe, effective products to market and keeping them there. It automates high-volume tasks, boosting speed, accuracy and efficiency, lowering costs, improving data quality and enabling global visibility across the portfolio. RIM Smart is the only solution with an embedded regulatory information database (RID).

Key benefits

Fast and accurate
- Automates high-volume, routine tasks
- Dramatically reduces cycle times
- Improves data quality and compliance performance
- Consumer-grade UI enables rapid adoption, higher ROI

Inter-connected
- Regulatory intelligence database embedded in the software
- Will seamlessly integrate with IQVIA's safety and quality systems
- Compatible with multiple CMSs and virtually every IT environment

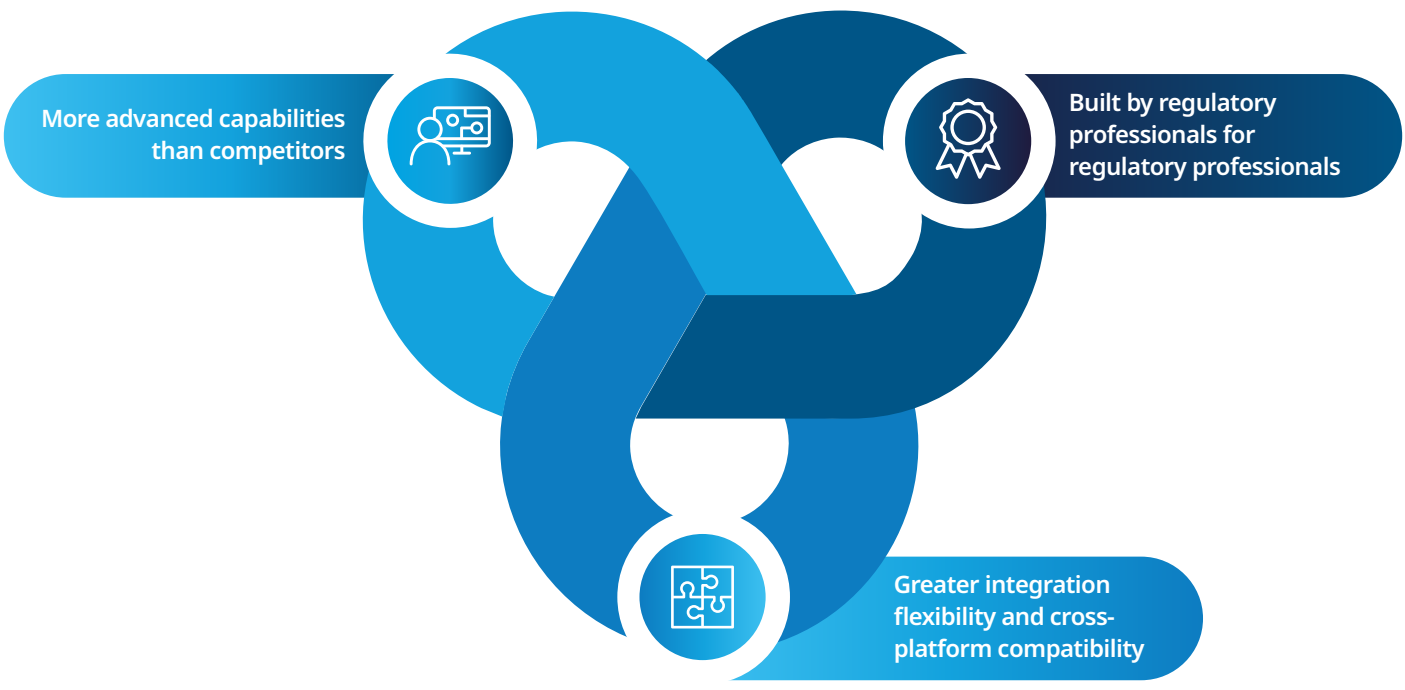
Cost-effective
- Reduces maintenance activity spend by >50%
- Fully integrated module array allows need-based configuration
- Cloud-based to deploy rapidly without capital expenditure or update disruptions

Intelligent
- AI and ML enhance proactive functionality, enabling better, more integrated management across entire regulatory process
- In future, will automate process of changing new regulations into workflows

The difference

IQVIA RIM Smart.

Based on industry needs, RIM Smart is a true end-to-end solution, with more functionality, integration, and planned enhancements than competitive solutions. We offer both pre-integration and open integration choices, allowing the flexibility to leverage existing investments. Over 1,900 strong across 100+ countries, our highly experienced in-house GRA services organization interacts with customers daily. Many have provided input into the development of IQVIA RIM Smart: we consider them Client 0.



Contact us to discuss your regulatory needs.

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