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.

>2000
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**

**High administrative
burden of
maintenance work**

**Disparate systems,
complex processes**
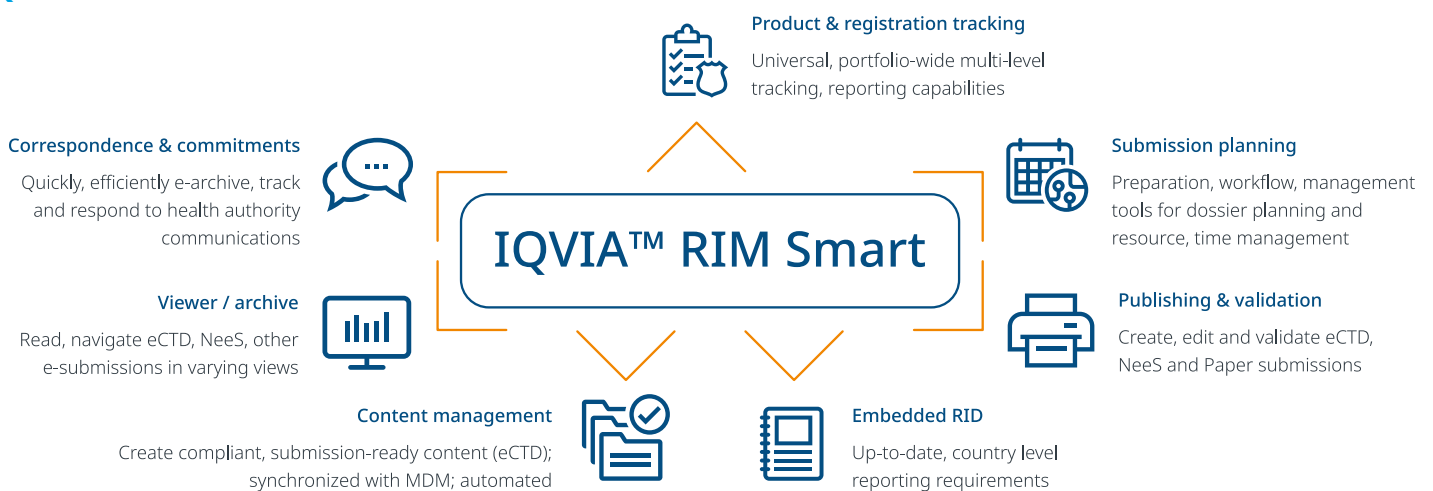
**Breaking down
silos for timely
access to information**

**Manual, high-touch,
repetitive work**

**Regulatory
demands in
more countries**

THE SOLUTION
IQVIA RIM Smart

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

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

IQVIA™ RIM Smart

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

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

Publishing & 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

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

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

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

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.

**More advanced capabilities
than competitors**

**Built by regulatory professionals
for regulatory professionals**

**Greater integration flexibility and
cross-platform compatibility**

CONTACT US TO DISCUSS YOUR REGULATORY NEEDS.
www.IQVIA.com