DXC Regulatory Information Management
Simplify • Amplify • Comply

Accelerate innovation through a centralized, secure and automated management platform for regulatory affairs information and processes.

The sensitive nature of regulatory affairs (RA) content and processes demands a unified platform to streamline regulatory data and workflows. Additionally, it is essential that teams across these RA processes are able to efficiently collaborate. This requires continuous interaction in an intuitive and accessible collaboration environment.

The DXC Regulatory Information Management Platform acts as your central hub – a cloud-based and scalable open platform that delivers enterprise-level security and integrated data flow across RA ecosystems.

Business Value

Simplify key RA operational processes to bring products to market faster

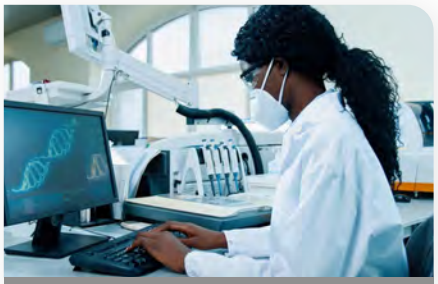
Amplify user experience through extensive collaboration tools

Comply with RA requirements and process while keeping efficiency and quality of data

Why choose the DXC Regulatory Information Management platform

Simplify

- Enhance productivity through effective automation features
- Reduce cost of ownership with an out-of-the-box solution that provides speed, scale and built-in security
- Monitor business performance through integrated data and real-time holistic reports



Amplify

- Streamline business processes and communications through embedded collabortive tools
- Increase output and improve data quality with advanced automation
- Increase performance, strengthen strategy and improve time to market with data insights

Comply

- Increase compliance by reducing manual errors with advanced automation.
- Utilize reliable rules engine for policy-driven control
- Comply with regulatory requirements across 13 regions, including electronic records management requirement 21-CFR-Part 11 and EudraLex Annex.

Boost your organization's efficiency in managing regulatory information through our end-to-end solutions and services.

DXC TRS Suite

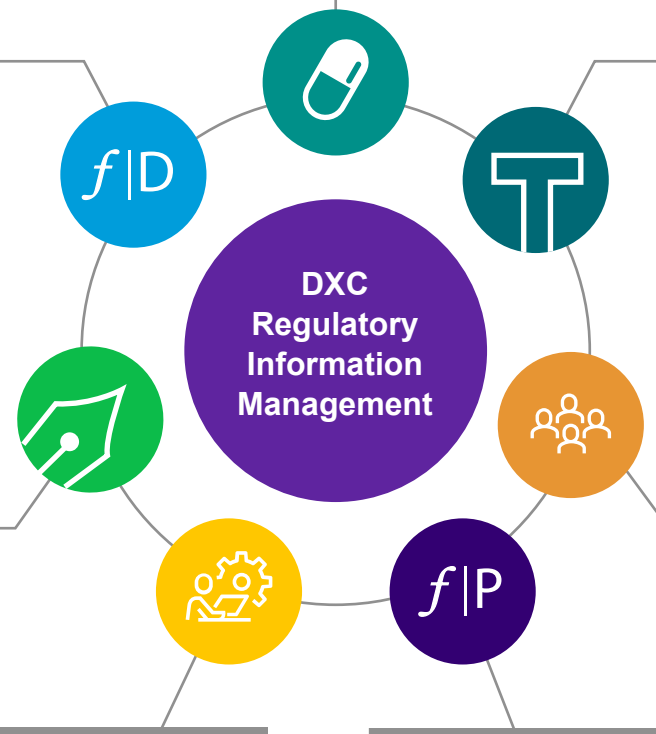
Efficiently manage drug applications through a suite of products and services that allows users to create, submit, validate, publish and manage regulatory content.

DXC FirstDoc & eTMF Tracker

Easily manage and track every product in your portfolio with best in practice eDMS for all your regulatory documents. Equipped with a trial master file planning functionality that is focused on study completeness, quality, and timeliness.

DXC ToolBox

Develop, modify, and validate documents, create bookmarks and hyperlinks, report metrics and perform other PDF transformation operations. DXC ToolBox has editions supporting legal, pharma, professional and standard operations.



DXC Writer

Accelerate the medical writing process through guidance-compliant template, enhanced collaborative tools and streamlined processes.

Professional Services

Maximize the value of Regulatory Information Management through DXC's highly capable and industry experienced team of professionals.

Regulatory Services

Ensure compliance with country or regional regulations in multiple markets covering document formatting, document publishing, dossier publishing and submission publishing.

DXC FirstPoint

DXC FirstPoint RIM.AI is an orchestration layer, designed to ingest and produce data from multiple different REST enabled & HTTP data sources with business defined automation, machine learning and artificial intelligence.

Get the insights that matter.
dxc.com/optin

To learn more about DXC Regulatory Information Management, please send an email to:

- Sales: RIM@dxc.com
- Licensing: ls_licensing@dxc.com
- Support: RIMSupport@dxc.com