DXC Regulatory Information Management (RIM) Platform

DXC RIM Platform manages regulatory affairs content, data and workflows in one central location, helping teams accelerate innovation and reduce time to market.

Speed, scale and flexibility in one central platform

The sensitive nature of regulatory affairs (RA) content demands a unified platform to handle regulatory data and workflows. Teams need simple, always-ready access to all relevant data. They also require an intuitive, customizable user interface that is easy to navigate. Collaborative tools and configurable reporting are crucial for success, but legacy technology can frustrate teams and slow processes.

Business challenges. Achieving compliance is a struggle. Collecting data and documenting processes is time consuming. Problems are compounded when life sciences organizations run fragmented legacy systems using disparate technologies. In some cases, they need to resort to spreadsheets to manage key operational processes. Bringing products to market and managing RA requires a centralized, enterprise-grade solution. DXC RIM Platform provides that single, connected ecosystem.

DXC solutions. Think of DXC RIM Platform as your central hub. This cloud-based, open platform scales with you, growing as your business grows. DXC RIM Platform delivers enterprise-level security while integrating and connecting data flow, without friction, across RA ecosystems. You gain simplified management of RA content, data and workflows. Along with RIM systems, you can also integrate electronic document management (EDM) systems and publishing solutions like electronic common technical documents (eCTD).

DXC will advise your team on how to deploy the RIM platform. You can expect a dynamically scalable SaaS service, managed by DXC (multitenant with no customer customization, or multi-instance with customer customization). If needed, we can deliver on-premises, customer-managed solutions.
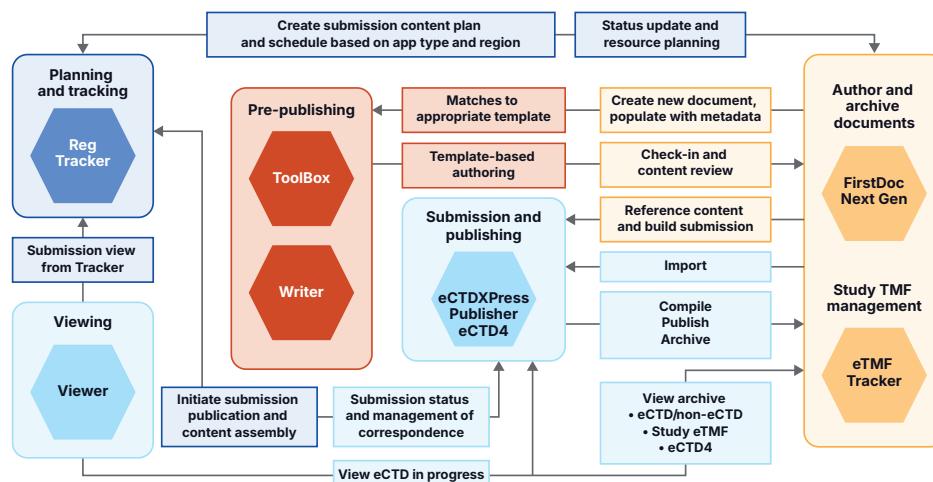
Key benefits

- Bring products to market faster by managing key RA operational processes
- Generate insights and business value for compliance
- Reduce costs and increase operational efficiency with automation
- Deploy flexible, scalable and low-cost SaaS options
- Simplify operations and amplify user experience

DXC RIM Platform: End-to-end enterprise management

All DXC solutions come mobile-ready for user bases of any size.

DXC RIM Platform in action

DXC RIM Platform provides a clear framework for RA life-cycle management in four key areas:

- Discovery
- Clinical
- Regulatory
- Safety

Across the enterprise, DXC RIM Platform transforms, standardizes and streamlines content, data, workflows and end-to-end processes. You can easily digitize content to prove compliance more efficiently. Our modern, web-enabled platform supports extensible, common core services that:

- Track, store and control documents
- Connect people and processes
- Drive data insights

DXC RIM Platform business value

Once deployed, DXC's RIM Platform globalizes the management of key RA processes, tasks and systems into one single, integrated platform. This enables companies to streamline compliance across the regulatory ecosystem.

DXC RIM Platform delivers:

- A reliable rules engine for policy-driven control
- Compliance with the electronic records management requirement 21-CFR-Part 11 of FDA regulations for electronic documentation and electronic signatures (ERES)
- Low ownership cost with an out-of-the-box solution that provides speed, scale and built-in security
- Advanced automation and AI to increase output and improve data quality
- Data integration from different systems, providing a real-time, holistic view of business performance
- Embedded collaborative tools to improve efficiencies and streamline business processes and communications
- Data insights to increase performance, strengthen strategy and improve time to market

Learn more about DXC's RA solutions

Capabilities

Controlled data management. Digitize regulatory content to improve data quality, increase content visibility, automate workflows and reduce risk.

Enhanced automation and productivity. Increase compliance and data accuracy with advanced automation and AI capabilities.

Cost reduction. Maximize ROI with low upfront investments and predictable as-a-service fees.

Why DXC?

DXC helps pharmaceutical, biotech, and medical device organizations modernize regulatory operations by connecting data, systems and workflows through AI, automation and digital twin technologies. We equip businesses with a single, connected platform to transform, standardize and enhance compliance and operational efficiency. This support empowers you to accelerate innovation and reduce time to market for new therapies, devices and medicines.

DXC