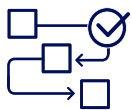


RIM

Regulatory Planning, Tracking and Registration Management

Regulatory Information Management (RIM) reduces data entry needs, improves data quality, and increases global oversight and control via robust capabilities to manage regulatory planning, tracking, and registration management.

Key Benefits



Standardized Processes

Save time and effort by ensuring more streamlined and consistent end-to-end workflows for regulatory planning and tracking across all markets.



Enhanced Collaboration

Make it simple for your teams to work together with various stakeholders on global and national levels.



Single Source of Truth

Eliminate siloes and harmonize your operations by establishing a single area for worldwide registration activities and records.



Product Specifics

Ensure compliance with support for investigational and marketed drug/medicinal products, as well as active substances, devices, site registrations and more.

Deliver real value
with RIM.

To learn more about RIM,
visit: lifesphere.arisglobal.com

Features

Regulatory Planning

Built-in support for planning your initial and maintenance submissions.

Enhanced Tracking

Track registrations, correspondence, commitments, and other key information.

Global Coverage

Centralize global submission planning and tracking.

Localized Support

Support specific regional requirements and regulatory knowledge.

IDMP Compliance

Support for the IDMP target operating model.

Interactive Dashboards

Gain actionable insights into regulatory activities and registrations.