

# RIMS

## Speed. Quality. Collaboration. Delivered in one simple, unified platform.

LifeSphere® RIMS delivers end-to-end regulatory information management in an all-new, easy-to-use cloud application that accelerates speed to market, reduces risk and streamlines collaboration across teams. With LifeSphere RIMS, regulatory affairs departments can seamlessly plan, track, and gain real-time visibility into all phases of the regulatory submissions process.

### Move fast without sacrificing data quality.

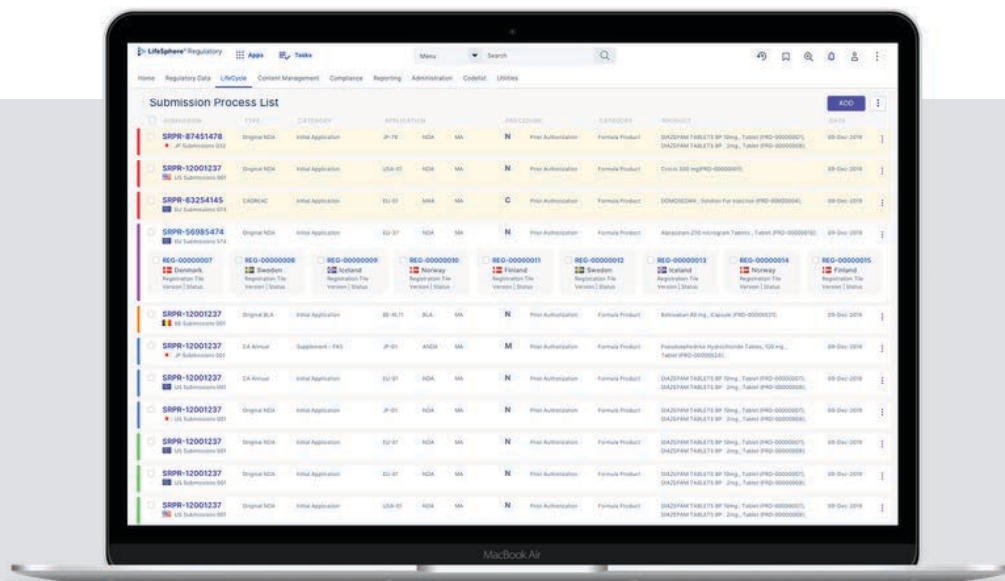
Accelerate speed to market by planning, executing, and tracking all regulatory activities in a single, unified RIM application, with seamless access to regulatory documents and full support for all major eCTD submission requirements. Built-in automation streamlines workflows to reduce administrative burden and provide greater confidence in product registration data quality.

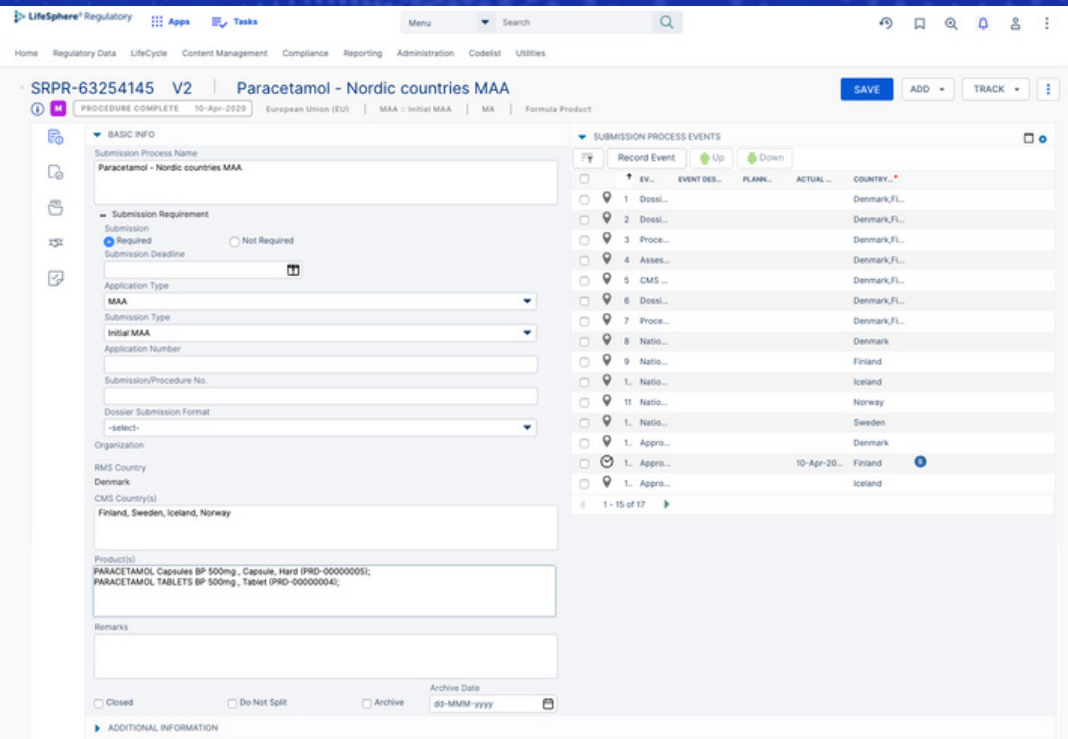
### Future-proof your compliance.

Reduce risk and stay compliant with the latest regulatory requirements thanks to scalable cloud architecture that provides continuous innovation via free and seamless upgrades. LifeSphere RIMS has been built from the ground up with a deep understanding of the regulatory lifecycle, offering a best-in-class standard configuration with full support for xEVMPD and IDMP data standards.

### Streamline collaboration.

LifeSphere® RIMS is unified regulatory application designed to streamline cross-functional collaboration across the entire organization. It seamlessly connects with LifeSphere® EasyDocs, a modern cloud document management platform, to provide a single source of truth for regulatory data and documents across R&D business processes, and LifeSphere® Publishing, a submissions management system with full support for all major global eCTD requirements. The result is real-time collaboration and visibility across all core regulatory processes.





The screenshot displays the LifeSphere RIMS interface for a regulatory submission. The main header shows the submission ID 'SRPR-63254145 V2' and the product name 'Paracetamol - Nordic countries MAA'. Below this, there are sections for 'BASIC INFO' and 'SUBMISSION PROCESS EVENTS'. The 'BASIC INFO' section includes fields for Submission Process Name, Submission Requirement (Required/Not Required), Application Type (MAA), Submission Type (Initial MAA), Application Number, Submission/Procedure No., Dossier Submission Format, Organization (Denmark), RMS Country (Denmark), and CMS Country(s) (Finland, Sweden, Iceland, Norway). The 'SUBMISSION PROCESS EVENTS' section shows a table of events with columns for Event ID, Event Description, Planned Date, Actual Date, and Country. The table lists 17 events, with the most recent one dated 10-Apr-2020.

DELIVERING RESULTS TO 250+  
GLOBAL CLIENTS

30  
YEARS AS A LEADER IN LIFE  
SCIENCES TECHNOLOGY

80%  
OF THE TOP 50 GLOBAL  
PHARMA AS CUSTOMERS

100%  
COMPLIANCE WITH  
GLOBAL REGULATIONS

## Features

### Simplify the Regulatory Lifecycle

Manage the entire regulatory lifecycle in a single platform. Streamline planning and tracking of interactions, commitments and obligations, as well as document and dossier management, publishing, reporting, and data standards compliance.

### Superior Automation Capabilities

Automation-assisted workflows – including intelligent task assignment, submission planning and performance tracking – and pre-configured templates help expedite regulatory submission and dossier planning.

### Easy Access to a Central Source of Documents and Data

Access up-to-date documents and data easily with seamless connectivity to LifeSphere® EasyDocs, and third-party document management systems.

### Easy Integration

Connects with existing systems via open architecture, leveraging API connectors and web services.

### Support for Medical Devices

Supports medical device registrations, submissions tracking, and compliance to FDA GUDID and EU EUDAMED UDI regulations.

### Full xEVMPD Compliance and IDMP-Readiness

Full support for xEVMPD, with UDI and IDMP-readiness to ensure future compliance.

### eCTD Submission Lifecycle Support

Manage the submission lifecycle, with full support for all major global eCTD requirements, including FDA applications and submissions.

### No Extensive Training Needed

Designed to be easy and intuitive to use, with an interface designed for users of all skill levels.

### Worry-Free Implementation

Fast, knowledgeable teams adhere to industry best practices to get your teams up and running quickly.

## About ArisGlobal

ArisGlobal transforms the way today's most successful life sciences companies develop breakthroughs and bring new products to market. Our end-to-end drug development technology platform, LifeSphere®, integrates our proprietary Nava® cognitive computing engine to automate all core functions of the drug development lifecycle. Designed with deep expertise and a long-term perspective that spans more than 30 years, LifeSphere® is a unified platform that boosts efficiency, ensures compliance, delivers actionable insights, and lowers total cost of ownership through multi-tenant SaaS architecture.

Headquartered in the United States, ArisGlobal has regional offices in Europe, India, Japan and China.

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