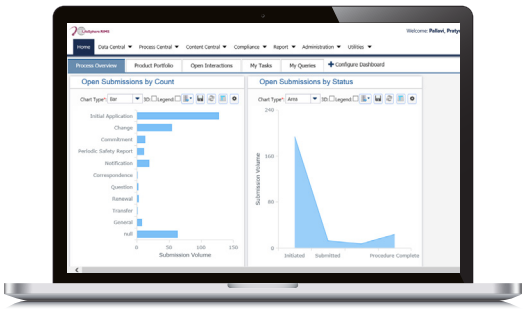
LifeSphere® RIMS

Enabling Intuitive, Mobile-enabled Regulatory Affairs Processes by all Stakeholders

Amid ever-changing regulatory requirements and the move toward data standards-based electronic submissions, today's legacy regulatory information management systems (RIMS) are no longer adequate to support the volume and complexity of regulatory documentation and information required for global compliance and maintenance of global product lifecycles worldwide.

LifeSphere RIMS is a cloud-based regulatory solution that provides intuitive handling of all regulatory affairs processes and makes it easy for all stakeholders – headquarters, regulatory operations, regional managers and local affiliates – to adapt to changing global regulations.



A Proven RIMS Designed for Ease of Use

Regulatory data tracking processes are highly complex, requiring insight into all aspects of the regulatory submission to meet timelines and address health authority questions. Regulatory and other departments must not only meet current and future regulatory requirements but also be able to collaborate with affiliates and other stakeholders.

LifeSphere RIMS is a powerful RIMS solution, built with a thorough understanding of the complexity of regulatory data tracking processes. The solution also ensures full support for xEVMPD and IDMP data extraction, review, validation and electronic submission needs.

Intuitive Interface and Visibility

LifeSphere RIMS boosts end-user acceptance and product adoption with minimal product training. Improved collaboration with affiliates or infrequent users is made possible with a more intuitive process-based user interface, role-specific workflow reminders, tasks lists and wizard-based approach to data entry.

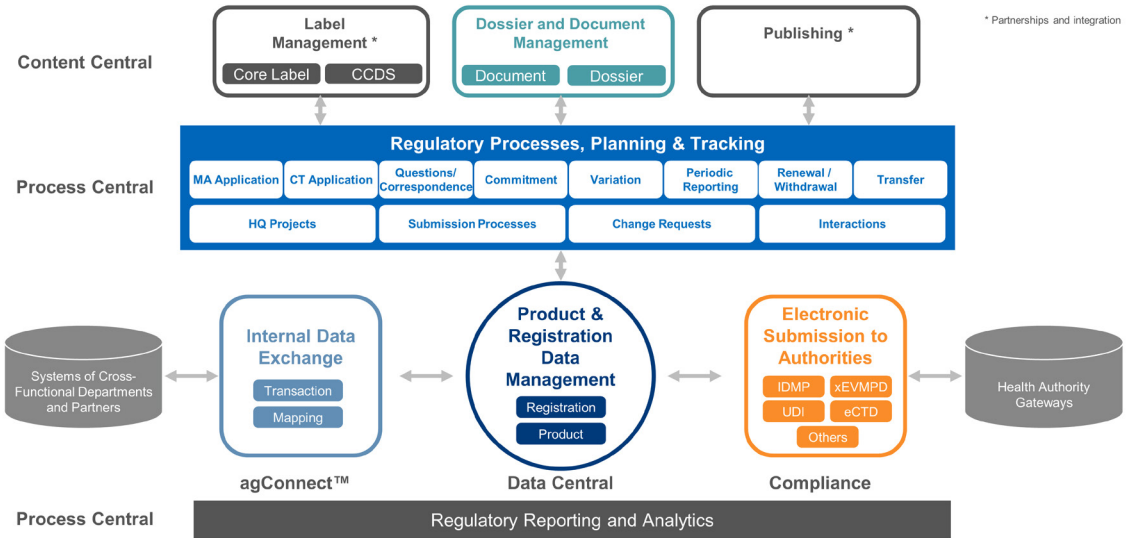
LifeSphere RIMS improves visibility across the organization, providing a global view of issues while enabling faster, informed decision-making at a lower cost.

A process layer on top of regulatory information enables the entire regulatory department to effectively plan, track and have quick status visibility into all phases of regulatory submissions.

Single Solution for Global Tracking

Core regulatory affairs modules give users a single solution for tracking all RA business processes. Real-time visibility of ongoing submissions, authorizations, commitments, open agency correspondence and upcoming renewals ensure your continued compliance.

A Single Solution for Tracking Regulatory Affairs Processes



Intelligent Insights

Comprehensive business process workflow automation enables intelligent tasks assignments, submission planning and performance tracking for improved e-fficiencies and informed decision making. Comprehensive business analytics capabilities allow RA leaders to visualize regulatory resource productivity and measure affiliate performance against expected KPI's with comprehensive business workflow tools and graphing capabilities.

Built on Best Practices

Users can leverage collective industry intelligence and standard practices for faster deployment and harmonized process implementation thanks to LifeSphere RIMS' built-in Industry Standard Practices.

ABOUT ARISGLOBAL®

ArisGlobal is a visionary technology company that's transforming the way today's most successful life sciences companies develop breakthroughs and bring new products to market. The ArisGlobal LifeSphere® cognitive technology platform integrates machine-learning capabilities to automate the core functions of the product lifecycle. Designed with deep expertise and a long-term perspective that spans more than 30 years, our cognitive platform delivers actionable insights, boosts efficiency, ensures compliance, and lowers total cost of ownership through multi-tenancy. Headquartered in the United States, ArisGlobal has regional offices in Europe, India and Japan. For more information, visit arisglobal.com or follow ArisGlobal on LinkedIn and Twitter.

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