ArisGlobal

A trusted partner with CROs in Clinical Research

Driving Efficiency & Excellence with ArisGlobal's Comprehensive CRO Capabilities and Partnership Initiative

Staying Ahead of the Curve with ArisGlobal for CROs

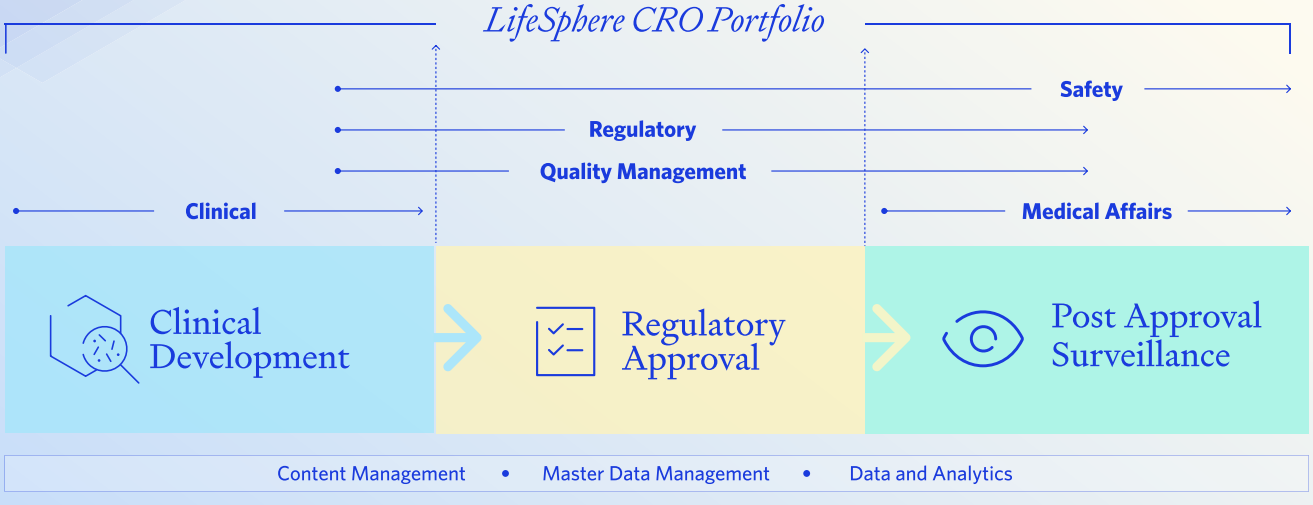
Drug research and development can be highly complex, demanding the delivery of safe and efficient patient therapies. ArisGlobal is committed to providing solutions that empower CROs in achieving success through the utilization of the LifeSphere platform suite of products. Recognizing the significance of timing, cost, and research quality, ArisGlobal focuses on navigating regulatory requirements and embracing industry trends to ensure continued excellence.

Why partner with ArisGlobal?

By partnering with ArisGlobal and harnessing our advanced technology, CROs can elevate their operational efficiency, enhance data quality and integrity, navigate ever-evolving regulations, and expedite regulatory submissions. Dedicated to innovation, partnership, and operational excellence, ArisGlobal is fully committed to assisting CROs in driving growth in their sponsor drug development.

Partnership Benefits

- Partner stewardship
- A dedicated CRO focused team
- Automation & self service
- Training resources & enablement programs
- Commercial growth incentives
- Competitive pricing



LifeSphere
by ArisGlobal

LifeSphere Portfolio for CROs

LifeSphere Safety for CRO

LifeSphere Safety empowers CROs to deliver cost-efficient pharmacovigilance services through the implementation of production-ready intelligent automation. With its flexible, no-code configuration of workflows and reports, LifeSphere Safety can effortlessly adapt to client-specific requirements. Furthermore, the platform rises to the challenges posed by decentralized, multinational trials by offering up-to-date global compliance capabilities, guaranteeing adherence to the latest local and global regulations. Additionally, LifeSphere Safety facilitates a seamless transition to post-market drug surveillance, leveraging its scalable cloud platform to efficiently monitor and analyze real-world data. This gives CROs greater autonomy in putting configurations in place to effectively meet the unique requirements of sponsors.

- **LifeSphere Multivigilence (LSMV)** enables one schema for multiple sponsors, with easy and efficient deployment and configuration for new sponsors.
- **LifeSphere Reporting and Analytics (LSRA)** automates data entry, collection, report generation, and result communication, saving time on manual workflows.

LifeSphere Regulatory for CRO

LifeSphere Regulatory is a comprehensive platform that enables efficient global product and project management throughout the regulatory process. With its open architecture design, it promotes effective collaboration with cross-functional teams and CRO partners. Additionally, LifeSphere Regulatory allows seamless integration with existing eDMS systems. This platform provides unique value by automating processes and linking content across multiple systems, ensuring streamlined planning, tracking, and content management.

- **LifeSphere RIMS** is a comprehensive regulatory information management solution that enhances efficiency through intelligent automation, speeds up time to market, mitigates risk, and facilitates collaboration via cloud-based software.
- **LifeSphere Investigational Product RIMS** is a specialized solution for precise automation and activity triggers in investigational submission and dossier planning.
- **LifeSphere Publishing** simplifies the compilation, publication, and validation of regulatory submissions for teams.
- **LifeSphere IDMP** ensures compliant collection, maintenance, and enrichment of data in accordance with IDMP and xEVMPD regulations.

LifeSphere Clinical for CRO

LifeSphere Clinical supports CROs by offering right-sized clinical solutions that grow as organizations scale – complete with out-of-the-box configurations to start quickly with no vendor lock-in. These integrated clinical solutions are not only user-friendly but also ensure complete transparency, enabling sponsors and site team partners to stay aligned and well-informed throughout the trial process. By streamlining study management and accelerating the trial lifecycle, LifeSphere Clinical empowers CROs to efficiently carry out their crucial work while maintaining optimal collaboration and productivity.

- **LifeSphere CTMS** provides a comprehensive clinical trial management solution in one application, simplifying processes and ensuring stakeholder alignment.
- **LifeSphere eTMF** offers a flexible document management system based on the TMF Reference Model, ensuring inspection readiness and compatibility with LifeSphere CTMS or external systems.
- **LifeSphere Safety Document Distribution** reduces the workload of global drug safety document management for CROs with manual processes.

To learn more, contact us today!