LIFESPHERE®

ArisGlobal

LifeSphere vs. COVID-19

Deploying a Next-Generation Drug Development Platform to Deliver COVID-19 Breakthroughs. Faster.

As teams race to develop COVID-19 therapies, it has never been more important to own a modern technology stack that can bring efficiency and speed to the drug development lifecycle. LifeSphere is a cloud platform that helps life sciences companies of all sizes accelerate drug development, maintain compliance and streamline collaboration between teams. With LifeSphere, life sciences organizations have a toolkit to scale COVID-19 trials and bring safer products to market more quickly.

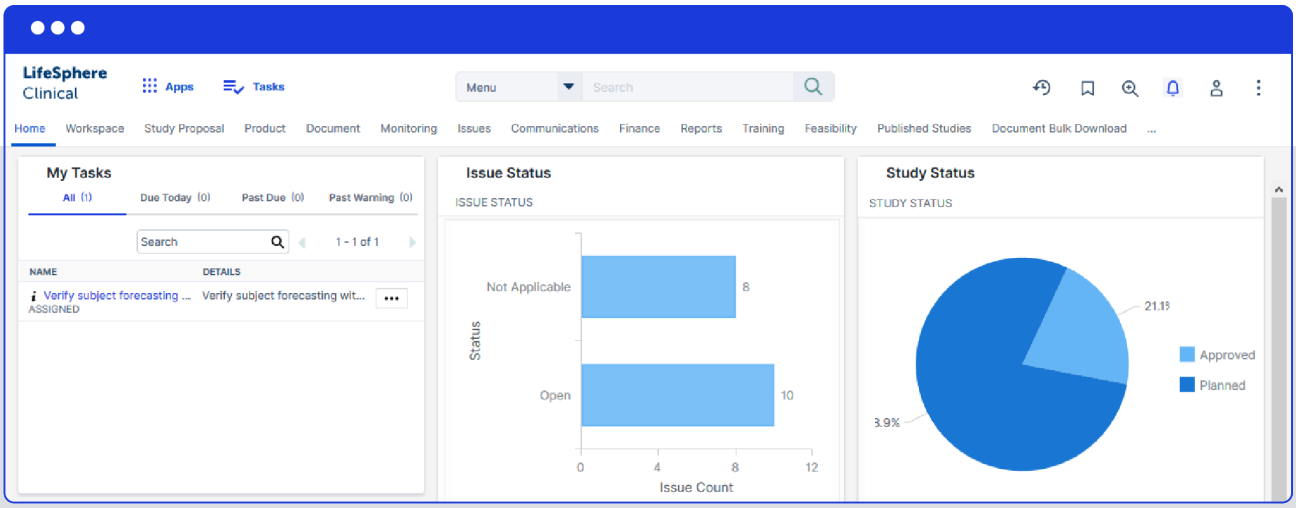
LifeSphere provides a best-in-class software platform for rapid development of COVID-19 therapies.

CLINICAL

LifeSphere CTMS

Everything an organization needs to plan, track and control the set-up, conduct and closeout of COVID-19 trials, delivered in a simple, easy-to-use application that ensures full oversight of global stakeholders.

- Reduces complexity to empower organizations of all sizes with a single, end-to-end application that includes payments and monitoring out-of-the-box
- Automates key activities to enhance productivity and deliver faster COVID-19 trials
- Unified with LifeSphere eTMF for rapid study planning, execution and closeout



CLINICAL

LifeSphere CTMS

Works seamlessly with LifeSphere CTMS to provide a centralized, global electronic trial master file for managing, accessing and archiving all COVID-19 trial-related documents.

- Maintains visibility, status and control of trial documentation
- Enables faster and more efficient site setup and initiation
- Ensures file structure and organization quality with support for the DIA TMF Reference Model

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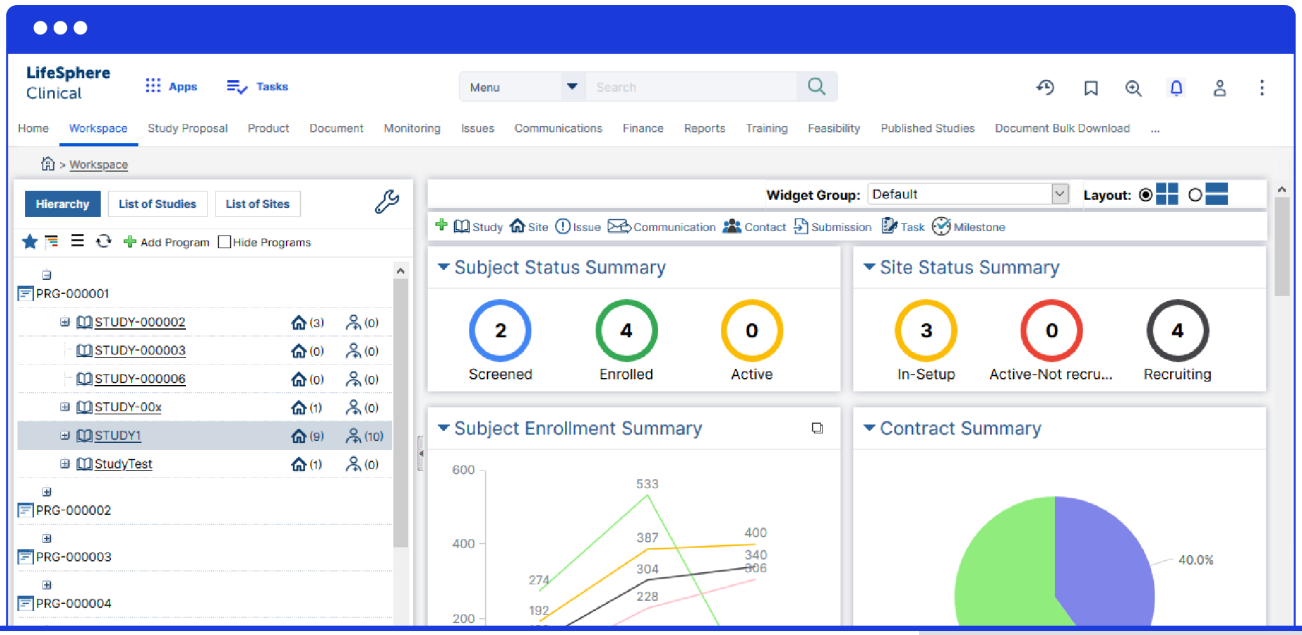

REGULATORY

LifeSphere RIMS

Provides intuitive handling of all regulatory affairs processes by headquarters, regulatory operations, regional managers and local affiliates. Delivering the necessary scalability, automation and single source of truth that regulatory teams need to navigate COVID-19 therapy approvals.

- Includes planning, tracking, document and dossier management, publishing, reporting, and data standards compliance in a single platform
- Easy to use interface designed for simplicity for every role in the regulatory process
- Seamlessly connected with LifeSphere Clinical solutions to give teams confidence that data and documents are accurate and up-to-date, and provide a single source of truth

SAFETY

LifeSphere MultiVigilance

An easy-to-use, all-in-one drug safety solution delivering compliance and efficiency, with rapid implementation to quickly meet the safety needs of present and future COVID-19 trials.

- Reduces overhead, easily scales with your needs and provides future-proof compliance
- Automation technology enables you to maintain a lean safety organization and focus key resources on development of COVID-19 therapies
- Supports end-to-end clinical and post-marketing safety needs in one simple suite that provides a single source of truth for safety data

DOCUMENT MANAGEMENT

LifeSphere EasyDocs

Streamlining enterprise document management between teams and ensuring the continuity of business processes across the end-to-end COVID-19 therapy development lifecycle.

- Ensures that your organization has a central, cloud-based repository for regulatory documents
- Connects seamlessly with LifeSphere solutions to provide a single source of truth for documents across development functions like Clinical, Regulatory and Safety
- Purpose-built for the industry to bring efficiency to cross-functional document workflows

For more information on how LifeSphere can enable rapid COVID-19 trial execution, visit our website or contact us today.

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We Bring the Future to Life