

eTMF

Real-time Quality, Completeness, and Timeliness

LifeSphere electronic Trial Master File (eTMF) system streamlines inspections, ensures compliance readiness, and improves efficiency from study start to close-out. As the first system to support the open eTMF exchange mechanism standard, LifeSphere eTMF brings real-time quality, completeness, and timeliness to the cloud. The system is unified with LifeSphere Clinical Trial Management System (CTMS) and LifeSphere Clinical Platform.

Streamline Inspections

Quickly and effectively respond to inspection requests at any point in the trial, including study closeout and archiving, with on demand access to all trial-related information and a clear view of expected documents by milestone status.

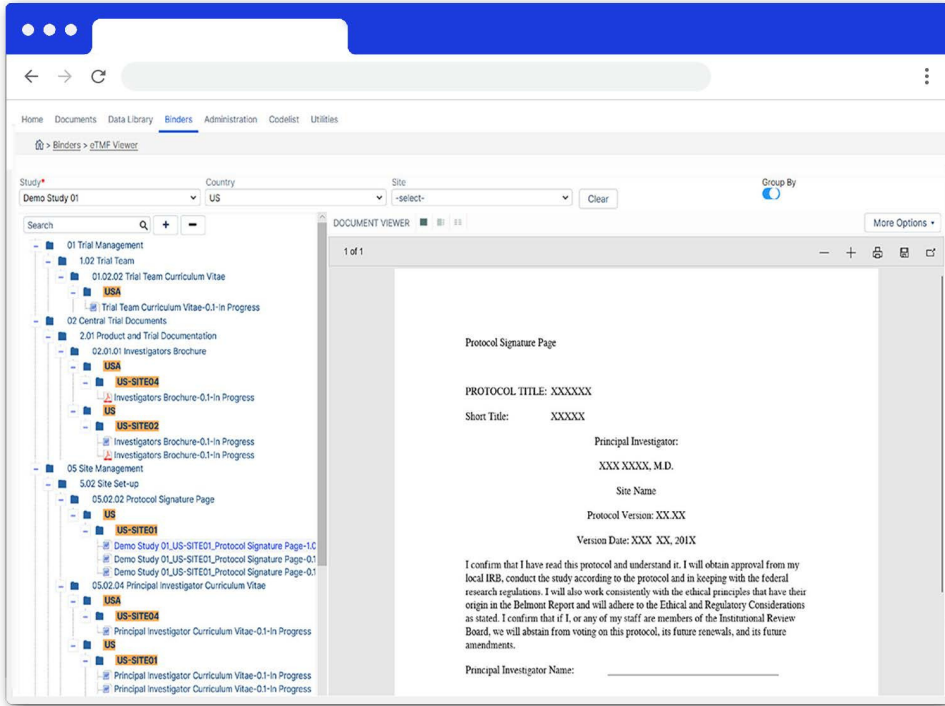
Future-Proof Compliance

Ensure GCP compliance with out-of-the-box embedded workflows and dashboards that allow auditors to easily reconstruct the trial. Leverage industry best practice to ensure your eTMF tells the complete story.

Save Time and Effort

Replace time-intensive manual tasks with automated workflows, giving stakeholders confidence in existing documents, processes, and data so they have time to focus on what matters most.





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

Complete visibility into all clinical trial documents

Fast, Easy TMF Planning

Our template-driven approach expedites TMF planning by ensuring alignment on expected document types. Quick and easy process to apply refinements to Yes/No and expected document counts.

Inspection-Ready, Intuitive TMF Viewer

Easy and intuitive hierarchy and viewer make browsing LifeSphere eTMF a breeze for internal consumers and auditors alike.

Global Search for Increased Productivity

As easy and quick to use as Google or Amazon, our global search returns your related documents and data records ranked by relevance.

Flexible and Easily Configured

Configurable watermarks, readers and footers, and the ability to have multiple rendition options per document type, provide limitless options to meet all your business needs.

Unified eTMF and CTMS Platform

CTMS project plan milestones drive expected documents. Document progress drives milestone statuses. With a unified eTMF and CTMS you have a real-time understanding of both your eTMF and your study status

Streamlined Collaboration

Unified with LifeSphere EasyDocs for content management, enabling real-time eTMF visibility across all departments. Share content with any other system using the eTMF Exchange Mechanism Standard.

Complete Compliance

With full audit trails and 21 CFR Part 11 compliant eSignatures, you gain a turn-key solution ready for Life Sciences companies of all sizes.

Worry-Free Implementation

Fast, knowledgeable teams adhere to industry best practices to get your teams up and running within 4-6 weeks.

Enterprise-Grade Security

User-, role- and group-based security rules ensure consistent, reliable security across organizations of any size or outsourcing and/or service model.

Easy Integration

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

About ArisGlobal

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

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

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