Clinical

cara LIFE SCIENCES PLATFORM

eTMF Archive

Streamline your process and cut costs with the AI-Enabled CARA Life Sciences Platform

generis

www.generiscorp.com

The CARA Life Sciences Platform offers a comprehensive Electronic Trial Master File (eTMF) Archive, designed to enhance the management, accessibility, and compliance of your clinical trial documentation. By integrating advanced automation, seamless collaboration, and robust compliance features, CARA empowers life sciences companies to efficiently manage trial records from inception through long-term archival.

With CARA, organisations benefit from a user-based pricing model rather than being charged per site or per study—significantly reducing costs. CARA's AI Assistant also makes it easier to find the information you need, with the ability to 'ask' the system a question in natural language.

eTMF Archive Process

Trial Complete

Finalise eTMF and ensure completeness

Archive

Lock eTMF and import with CARA Archivist

Retrieve

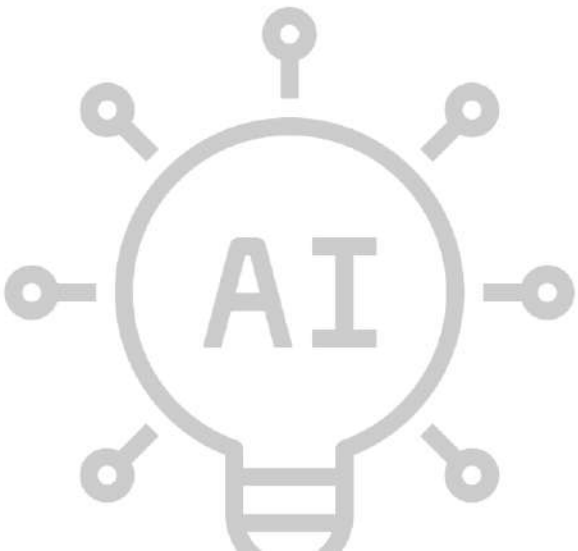
Leverage AI for inspection ready information search on demand

Store

Secure, readable, accessible archive for 15-25+ years

Decommission

Shut down active archives when this is due



Benefits

Secure Long-Term Archival

Ensures documents are stored securely with controlled access, maintaining integrity and compliance over required retention periods.

Comprehensive Metrics and Analytics

Offers dashboards and reports, including missing document reports, to monitor trial documentation status and ensure completeness.

DIA Reference Model Compliance

Aligns with industry standards for structured documentation, ensuring consistency and regulatory adherence.

User-Friendly Interface

Provides a consistent and intuitive user experience across all modules, reducing training time and enhancing user adoption.

Seamless Information Flow

Provides a unified platform for clinical trial management by connecting eTMF with CTMS, enabling real-time data sharing and reducing redundancy.

Simple Integration

Connect CARA to your other systems and easily pull the TMF data from external environments to archive in CARA.

Why CARA Life Sciences Platform?

Enhanced Compliance

Stay aligned with regulatory requirements through standardised processes and comprehensive audit trails.

Operational Efficiency

Streamline document management processes, reducing administrative burden and accelerating trial timelines.

Improved Collaboration

Facilitate seamless communication and document exchange between internal teams and external partners.

Scalability

Adapt to the evolving needs of your organisation with a flexible platform that grows with you.

Experience a transformative approach to clinical trial documentation with CARA's eTMF Archive module, designed to optimize your trial management processes and ensure regulatory success.

Contact us for a demo or evaluation
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