

Edge Archive

Stress-free regulatory compliant digital preservation solution

Life science companies are struggling with decentralized data storage and non-qualified environments, costly on-premise or legacy systems, and rapidly growing data needs, all while needing to demonstrate compliance with industry regulations.

Edge Archive is a cloud-based digital preservation platform that combines and maintains your organization's information in a central location, keeping content accessible and usable regardless of where the content originated. Edge Archive provides long-term content integrity, authenticity, and accessibility in a secure and compliant environment.

As part of the Medidata Clinical Cloud®, Edge Archive reduces your archival complexities and costs with an easy-to-use and effective content management system for all stakeholders.



Reduce Risk

Highly secure and compliant

Edge Archive's digital preservation imports from original sources, such as paper or DVD, eliminating the need to rely on legacy-based media. Edge Archive maintains compliance per data handling and content retention regulatory guidelines, safeguarding the integrity and authenticity of all content.



Reduce Cost

Eliminates legacy and on-premise costs

Avoid unnecessary costs for licenses and maintenance fees on multiple—or inefficient—legacy systems. Edge Archive consolidates all content and provides optimal data usage, saving time and money.



Reduce Complexity

Scalable & unified easy-to-use solution

Easy and automatic bulk uploads from external systems allow unification of all content in a cloud-based repository. More importantly, Edge Archive provides comprehensive metadata management, making information easy to find in an intuitive user interface.

KEY FEATURES

- ≠ 21 CFR Part 11 compliant
- ≠ Fully validated and highly secure cloud content platform
- ≠ Rapid and easy implementation
- ≠ One-touch content import, drag-and-drop support
- ≠ Preservation of source file and folder structures
- ≠ Automatic generation of migration reports, full audit logs, and signatures
- ≠ Audit trails and reports
- ≠ Roles based system for content search
- ≠ Electronic signatures
- ≠ File access control and metadata

“The addition of Medidata’s Edge Archive capabilities will allow us to collaborate across our clinical trials with greater ease and compliance.”

Dr. Albert Collinson
President and CEO of Theracos

45%

of life science CIOs ranked Enterprise Content Management as a top priority based on company spending plans in 2016-2017

Manage content with ease



Unlike on-premise or general enterprise content management systems, Edge Archive allows for easy bulk content migration, search and access, as well as structural preservation of regulated content coming from external sources (CRO, M&A related events, etc.).

Highly integrated with Medidata Clinical Cloud, Edge Archive provides a single source of truth for regulated content and data including TMFs, contracts, CVs, and IRB letters with full access and advanced search capabilities.

- Seamless**

Medidata's pre-built and pre-validated functionality makes content migration seamless, allowing you to implement quickly and confidently.
- Simple**

Clean, easy, and intuitive user experience makes working with content straightforward, even for the occasional user.
- Accessible**

Advanced search and retrieval capabilities on a unified platform provide access to all content anytime, anywhere.
- Secure**

Built and managed to the highest standards of validation and security, Edge Archive allows you to be audit ready.

Revolutionizing Life Sciences with the Edge Ecosystem

As part of the Medidata Edge Regulated Content Management, Edge Archive along with Edge eTMF and Edge Quality are purpose-built for life sciences, delivering a unified solution to manage both regulated and non-regulated content that is effective and quickly deployable to ensure high integrity, traceability, and compliance.

Medidata's proven methodology and track record give you the confidence and peace of mind for a seamless transition to the cloud along with the goal to support more effective cross-enterprise collaboration, our transformative tools and insights aid in clinical trials, supporting R&D to focus on better patient outcomes while reducing costs and creating a more agile operating model.

1. 2017 CIO Agenda: A Life Science Perspective, Feb 2017, Stephan Davies, Michael Shanler

About Medidata Solutions

Medidata is reinventing global drug and medical device development by creating the industry's leading cloud-based solutions for clinical research. Through our advanced applications and intelligent data analytics, Medidata helps advance the scientific goals of life sciences customers worldwide, including nearly 850 global pharmaceutical companies, biotech, diagnostic, and device firms, leading academic medical centers, and contract research organizations.

The Medidata Clinical Cloud® brings a new level of quality and efficiency to clinical trials that empower our customers to make more informed decisions earlier and faster. Our unparalleled clinical trial data assets provide deep insights that pave the way for future growth. The Medidata Clinical Cloud is the primary technology solution powering clinical trials for 18 of the world's top 25 global pharmaceutical companies and is used by 18 of the top 20 medical device developers—from study design and planning, through execution, management, and reporting.

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