

Built leveraging 35 years of Clinical Trial experience

ComplyDocs eTMF is an eTMF solution that helps minimize the risk and complexity of managing clinical trial documents. ComplyDocs eTMF is a software solution provided by [McDougall Scientific](#). For over 35 years, McDougall Scientific has earned a reputation as one of the most experienced CROs in the industry.



ComplyDocs

Powerful eTMF Solution for your Clinical Trial needs

We continue to earn our solid reputation.

FEATURES & BENEFITS

Managing clinical trial documents through paper-based or network-folder TMFs can be time-consuming and may produce costly errors that put your clinical trials at risk. Adopting an eTMF allows for shorter trial start-up and close-out times, cost savings and real-time oversight and management of documents to ensure compliance and audit readiness throughout the trial.



Real Time Dashboards & Reports

Gain real-time insight into the health of your TMF with simple, easy-to-navigate reports. Identify problems as they occur. Upload and share documents, find out what is completed and what is missing.



Review & QC Workflow

Built-in workflow routes documents through a QC review process, reducing manual work and ensuring quality issues are resolved as they arise – not weeks or months later.



Secure & Compliant Role-Based Access

TMF content is maintained in a secure, compliant environment with controlled access, configurable user roles, and a complete audit trail.



Support for TMF Reference Model

ComplyDocs eTMF has built-in support for the DIA TMF Reference Model; identifying & classifying trial documents, providing a standard set of categories and metadata.



Easy to use



Quick to Implement



Affordable to deploy

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