



# Flex Databases Platform

Unified eClinical platform for CRO, biotech, and pharma

## GENERAL

 HR Database

 Project Catalogue

 Report Tool

## DOCUMENTS

 Trial Master File

 Document Management System

## CLINICAL

 CRA Activity Management

 Investigators & Sites Management

 Subject Tracking & Invoicing

## PROJECT & FINANCE MANAGEMENT

 Project Management & Budgeting

 Time Sheets & Utilization

## QUALITY & COMPLIANCE

 Learning Management System

 Quality Management System

## SAFETY

 Pharmacovigilance

## About Flex Databases

We develop a comprehensive clinical trial management solution since 2011.

- **10+ years** on the market
- **4 offices** in Europe & the US
- **All-in-one platform** from study planning to safety database

## What sets us apart

Easy implementation

Full compliance

Complete data safety

Flexible solution

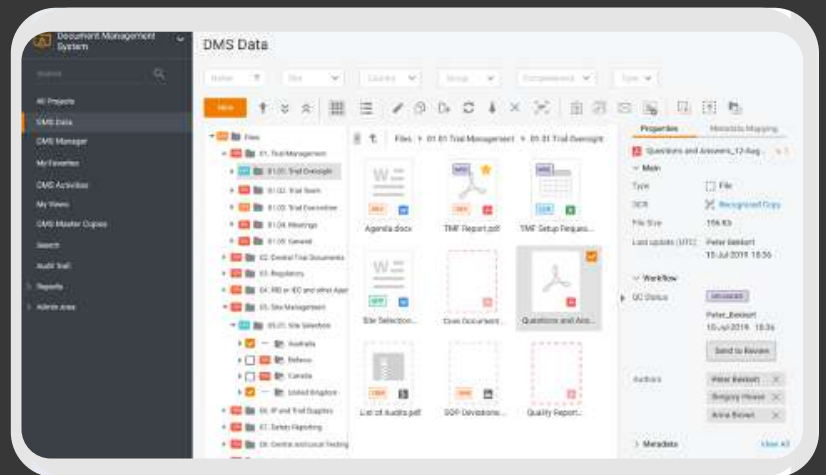
- System implementation in **3 to 10 weeks**
- A robust **backup & disaster recovery and data protection strategy**, including distributed data storages around the world
- All major **international and local regulations** covered, including FDA 21 CFR Part 11, GDPR, etc.
- Reports, workflows, trackers are **under user configuration**. Web-based solution, no installation required

## Some of our clients

Leading CROs, pharma, and biotech



# Document Management System Module Overview



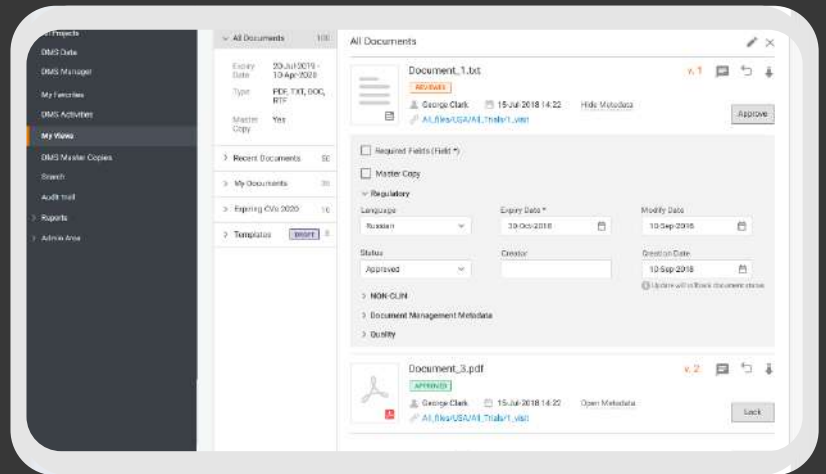
A comprehensive system designed to enhance efficiency and compliance throughout your document lifecycle

- eSignatures that comply with all legal requirements are an in-built part of the system. With a clear indication of signatory identity, signature reason, and timestamp, you can confidently rely on the authenticity of your documents.
- Fully validated for regulatory compliance: our system adheres to the highest standards. It is developed in alignment with 21 CFR Part 11, ensuring data integrity and security in FDA- and EMA-regulated environments.
- Seamless integration for data reusability: unlock the true potential of your information. Our Document Management System (DMS) seamlessly connects with other systems, facilitating easy data exchange and reusability for future needs.

## A ready-to-use system with extreme flexibility

- Tailored access control: effortlessly manage user permissions by differentiating access levels for team members, external auditors, or inspectors based on your preferences and security needs.
- Choose from built-in or customize your structure templates: our system offers a wide range of pre-designed templates to suit various project structures. Alternatively, you can create your own templates, ensuring that your data is organized in a way that aligns perfectly with your workflow.
- Create your own workflow: empower your organization with a fully customizable workflow. Design and implement workflows that mirror your specific processes, streamlining document management and ensuring efficiency at every step.

# Document Management System Module Overview



## Have a user-friendly system that can be navigated intuitively without excessive training

- Effortless communication options: infinite communication tools at your fingertips. Leave comments directly on the documents, initiate chats with team members, and raise queries.
- Full document tracking and archiving: keep a complete trail of all document actions and changes. Our system ensures comprehensive tracking, enabling easy extraction of data when needed. Additionally, utilize our archiving function for organized data storage and retrieval.
- Smart search and filtering: find what you need in seconds. Our powerful search function allows you to locate specific documents with ease. You can also filter documents by status, milestone, completeness, and more, enabling efficient organization and access.
- Send in and out: use any of your favorite ways to upload a document into the system – drag and drop, upload it from your computer, or even send a document into the system. And it goes both ways: email your documents directly from the system.