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 & COMPLINCE
- 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 the US, Europe, and Asia
- **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

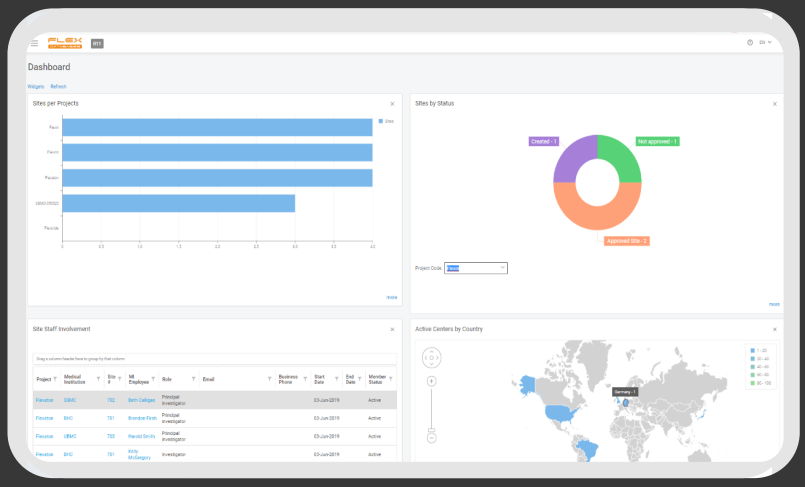
Leading CROs, pharma, and biotech

novo nordisk, CMIC, FGK CLINICAL RESEARCH, Dr.Reddy's, ERGOMED, OPTIMAPHARM, WORLDWIDE CLINICAL TRIALS

FLEX DATABASES

Investigators & Sites Management

Module Overview



Organize all the information on investigators, medical institutions, sites, regulatory authorities, and vendors

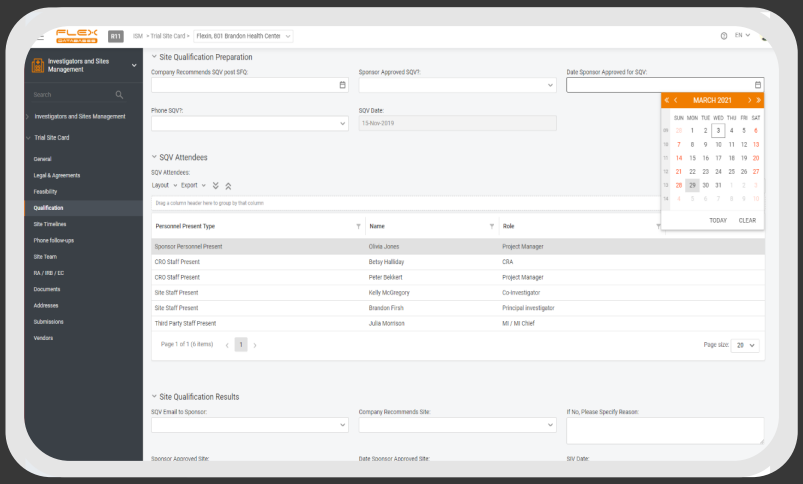
Sites statuses, addresses, documents, contracts, qualification, vendors, submissions, site staff, timelines and so much more at hand.

Find all the related information in few clicks
Keep all your Medical Institutions, Sites, and Sites Staff information in a clear organized manner.

Manage submissions packages across countries, studies and sites in one interface.

Investigators & Sites Management

List of Features



Regulators and investigators

- RA, IRB, and EC information: address, contact person, periodicity of session, terms of submission, and other key information
- Investigators information: contacts, CVs, certificates, and licenses, experience, studies, therapeutic areas, etc.

Hospitals, clinics, and vendors

- Hospitals and out-patient clinics (addresses, description, equipment, capacity)
- Vendor information (address, pricing, services)

Site management

- Sites information: address, accreditation certificates, study team, and studies
- Site team assignment
- Site performance
- Capturing and tracking of site communication

Tracking and reporting

- Documents tracking: contracts, site regulatory documents, licenses, and certificates
- Centralized IRB/LEC submissions and approval tracking
- Ad-hoc reporting tool for cross-project and cross-module reporting (graphs, widgets)

Contact us to learn more about Flex Databases Platform: contact@flexdatabases.com