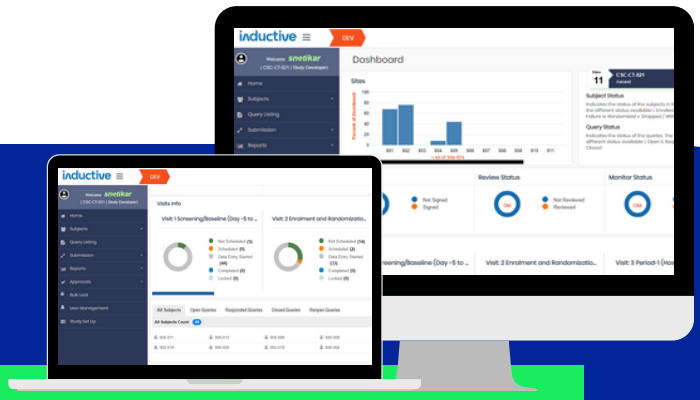
Inductive Quotient

INDUCTIVE EDC

ACCELERATE YOUR CLINICAL RESEARCH WITH SEAMLESS DATA MANAGEMENT

Inductive EDC is the most feature-rich and cost-effective software to capture, manage and report clinical trials data across study phases to advance your research without delays. Our intuitive platform is designed to be a reliable data management ecosystem for faster study setup, improved data quality and enhanced data security. Inductive EDC is built on not just technical expertise but also decades of experience in supporting clinical trials.

Inductive EDC will give you single-screen view into your data at any stage of the clinical trial's life cycle to empower you with insights for better decision-making. Our software is highly configurable and supports both - a single clinical trial and a cluster of studies across therapeutic areas. The robust platform facilitates you to scale your study to reach your goals faster and without hassle.



Inbuilt Compliance

FDA 21 CFR Part 11 · EU GDPR COMPLIANT · GAMP · HIPAA COMPLIANCE · ISO 27001 · ISO 9001:2015

SPEED

Design & Build Studies Faster

Build your study setup rapidly with drag & drop option to design visit schedules, case report forms, edit checks and workflows, and achieve FPI (First Patient In) in just days instead of weeks.

PERSONALIZATION

Configure for Your Unique Research

Customize your CRF layouts in a jiffy regardless of complexity of your study requirements. Using extensive library of forms, you can also modify queries and approval steps.

FLEXIBILITY

Make Mid-Study Changes

Empower your data managers and analysts to instantly make mid-study changes like protocol amendments or updated requirements reducing the dependency on tech support.

QUALITY

Set-Up Real-time Edit Checks

Ensure data quality through an ability to setup edit checks in real time at both field-level and cross-form, so that you move ahead with your studies without delays.

ANALYTICS

Automate Report Generation

Gain in-depth insights into your studies across phases, therapeutic areas, indications, and eligibility so that you can efficiently monitor through customized & automated reports.

COMPLIANCE

Built-In Regulatory Compliance

Ensure your data is in compliance with global standards and regulations including FDA 21 CFR Part 11, ICH GCP (HIPAA), GDPR (EU), ISO 27001, and ISO 9001.

INTEROPERABILITY

Integrate with any tool of your choice

Our tool gives you the freedom to work with any clinical tool for data integration and further processing. We also offer an API so that you can streamline your clinical data management.

COST

Full Functionality within Budget

Get access to end-to-end trial data management platform at market-best price with transparent pricing structure and zero hidden or extra costs.

Other Key Features of Inductive EDC

- ✓ Lab Management
- ✓ Audit Trail
- ✓ Digital Signature
- ✓ Document Imaging
- ✓ FDA 21 CFR Part 11
- ✓ Customizable Fields
- ✓ Data Import/Export
- ✓ Alerts/Reminders
- ✓ Activity Dashboard
- ✓ Real Time Notifications
- ✓ API Support
- ✓ Secure Data Storage