cara™
LIFE SCIENCES PLATFORM

Manage Clinical Trials from Protocol Approval to the Study Closeout through an integrated CTMS on the CARA Life Sciences Platform™. CTMS enables the consistent use of the Study Protocol and Study Master Data along the entire Clinical Trial operations value chain.

Our CTMS provides a single source of truth for all operational Clinical Trial data and workflows. It helps Clinical Trial Managers to make operational decisions based on timely, accurate and consistent data consolidated across simultaneous workstreams like study and site initiation, patient recruitment, monitoring and drug supply.

Reach across the organisation
Connect all clinical trial operation workflows through CTMS data & master data.

Align disparate workstreams
CTMS on the CARA Life Sciences Platform is the single source of truth for operational data, integrating multiple systems and processes.

Adapt to evolving regulations
Conduct clinical trails by adapting the latest regulations to the advantage and safety of enrolled patients.

Data Interoperability
The CARA Life Sciences Platform™ supports the industry standard data types and formats for enterprise-wide data exchange.

Features

- Clinical Program/Project Management
- Trial and Site planning
- Trial Design
- Site and Subject Management
- Study Management
- Investigator Management
- Clinical Supply Tracking
- Clinical Trial Performance and Reporting