eTrialTrack

Streamlining Clinical Trials

Speeding up the start and completion of clinical trials, reducing costs, and increasing accuracy & efficiency with **decades of industry experience and one seamless solution.**

Unified Electronic Data Management Platform

Sponsor/CRO-driven build-and-deploy flexibility

Always audit-ready, by design

Built to scale—from pilot studies to global, multi-site trials

www.quantum-quip.com

info@quantum-quip.com

About eTrialTrack

We Didn't Just Build Another EDC. We Fixed What Others Missed

eTrialTrack is a unified, fully integrated EDC platform designed to simplify and accelerate clinical trials across geographies. Created by global clinical technology experts with over 15 years of domain experience, eTrialTrack supports trials of any size, phase, or complexity—from early-phase studies to large, multi-country programs.

Its intuitive no-code study builder with drag-and-drop functionality enables local teams and global sponsors alike to independently design and deploy studies—no programming or vendor dependence required. eTrialTrack consolidates eSource, RTSM, eConsent, ePRO, Medical Coding, CTMS & eTMF in a single interface, removing operational silos and improving data consistency across borders.

With built-in capabilities for risk-based monitoring, adaptive trial designs, and custom real-time reporting, the platform offers unparalleled visibility and control. Teams can generate data correction reports, access AllPrint functionality, and ensure compliance with international regulatory standards, including 21 CFR Part 11, GDPR, and ICH GCP.

Trusted by multinational sponsors and CROs alike, eTrialTrack empowers clinical operations with true oversight, centralized control, and audit readiness—anytime, anywhere.

Key Features and Benefits

- Customizable eCRFs with drag-and-drop design
- ODM export capabilities for seamless interoperability
- Auto encoding of medical terms using MedDRA & WHODD dictionaries
- Real-time analytics and reporting dashboards
- Automated data validation and compliance checks
- Integrations with other EHR and Lab systems
- Global Compliance & Data Security
- One build, One Database

SINGLE CONFIGURABLE BUILDER FOR ALL

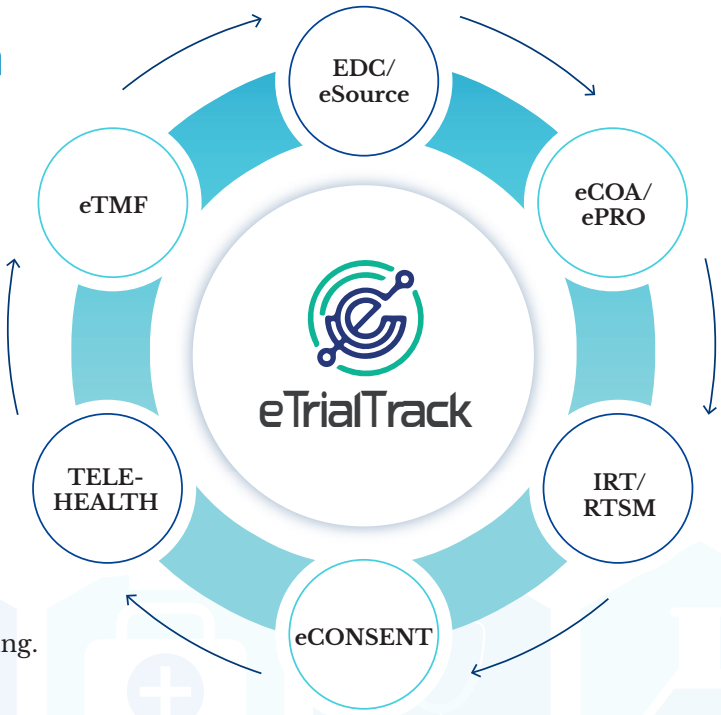
Our unified platform is supported by our configurable designer. Utilize one tool to configure any combination of eTrialTrack platform solutions through a highly configurable build, with no software coding.

The simple-to-use editor, its underlying CDISC ODM database, along with our library of preset forms, reusable study layouts and standards, makes it easy to deliver studies rapidly – typically in 4-6 weeks.

This unified platform is easy for sites and sponsors, with a single access point, role based access control per user, and a converged guided workflow allowing users to move seamlessly between solutions.

Our Unified Solution

eTrialTrack is a single unified platform containing all the key technology components for any clinical trial. Take advantage of some or all of the eClinical technologies in this singular, unified platform including fully integrated EDC, eCOA, eConsent, RTSM, Tele-Health, and Participant Tracker applications. Being part of the same unified platform means a streamlined experience for sites and sponsors: a single place to login, and a single set of access credentials; simpler, converged workflow across individual platform components; and a single database enabling consolidated oversight and reporting.



EDC
Conversion of data into electronic from paper/ source documents.

eSOURCE
Real time data entry per visit/ each update. Source data validation checks are not required .

IRT
IWRS/ IRT helps clinical trial sponsors and sites manage the patient and drug supply logistics throughout a clinical trial.

ePRO
Capturing outcomes data electronically. Employs handheld devices or the web to allow trial participants, physicians to directly report information.

ENCODER
A software platform that selects medical codes based on a set of terminology.

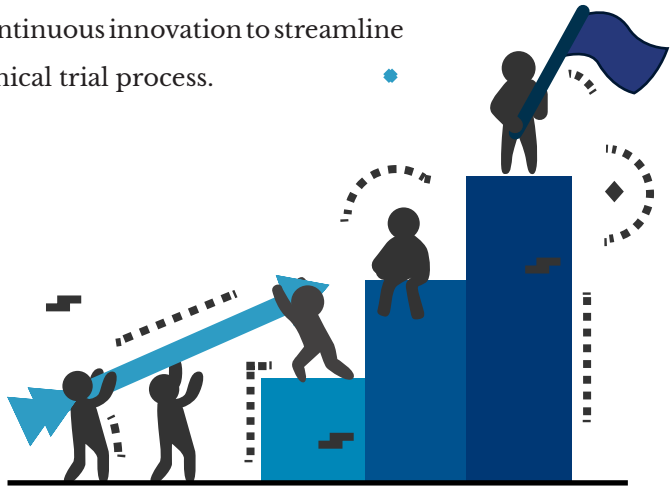
eCONSENT
Documenting information exchange between the researcher and the prospective subject, that consent was obtained. Audio video consenting is also available.

Our MISSION

To revolutionize clinical trial management by providing a cutting-edge electronic data capture solution that ensures efficiency, accuracy, and compliance, ultimately accelerating the path to groundbreaking medical discoveries and improved patient outcomes.

Our VISION

Empowering clinical trial professionals to efficiently manage trial data, ensure regulatory compliance, and enhance data integrity. We are committed to supporting our users with exceptional service and continuous innovation to streamline the clinical trial process.



WHY US

- **Built by Experts with 15+ Years of EDC Experience**
- **Scalable to any size of study**
- **Rapid Implementation**
- **24*7 global support**
- **Risk-Based Monitoring & Adaptive Trial Ready**
- **Flexible Engagement Models**

Contact Information

+91 987 35 14525 / +91-11-6879 6542

info@quantum-quip.com

https://www.etrialtrack.com/

NOIDA MUMBAI MD, USA