# CTMS

## Best-in-Class Clinical Trial Management

Our CTMS empowers every stakeholder—from CRAs and study managers, to IT teams and business leadership—to be more efficient. By simplifying study planning and oversight, TI CTMS removes the friction from day-to-day clinical operations. Learn how you can unlock real-time insights, mitigate global regulatory compliance risks, and accelerate therapeutic breakthroughs with a modern CTMS.

**DESIGNED FOR CLINICAL TEAMS**

**WORLD CLASS USABILITY**

**FLEXIBLE AND EASY TO CONFIGURE**

**BUILT IN eTMF AND SITE PORTAL**

## TRIAL INTERACTIVE CTMS: FOR CLINICAL, BY CLINICAL

### Features Overview

- Mobile App for Site Management and Monitoring Activities
- Visual Reporting through Dashboards
- Clinical Portfolio, Program, and Product Management
- Sponsor, Site, Vendor, and Lab Management
- Country and Region Management
- Better Process Adherence through Templates
- Activity Plans with Milestones
- Subject and Subject Visit Management
- Site Visit Related Letters and Report Authoring
- Protocol Deviation and Issue Tracking
- Clinical Trial Performance and Reporting
- Configurable Reports with Custom Fields
- Flexible Support for Many Trial Designs
- Built-in, Best-in-class Content Management and eTMF
- Site Activation and Study Start-Up
- Document Workflows with eSignature Approvals
- Powerful Web Services and Digital Interoperability
- EDC, IXRS, and Payment Integrations
- 21 CFR Part 11 and EU Annex 11 Compliant
- User Defined Experience
- Site Visit Calendar View

## FEATURE DETAILS

### Study View

Study View provides a clear layout of countries, sites, organizations, contacts, activities, milestones, documents, and much more.

### Study Management

Manage the study within the TI CTMS, with simplified access rights. A comprehensive set of metadata may be defined in the study for a single source of truth.

### Milestones

Milestones for a simple view of the overall study progress, showing both planned and actual milestones by date, priority, and risk.

### Activity Plans

Publish and apply plan templates across multiple countries, sites and studies.

### Virtual Dashboards

Dashboards with visual insights on subjects, enrollment, milestones, activities, documents, and plans. The study manager can assess deviation or safety trends, visit report cycle times, and other key performance indicators. (KPIs)

### eTMF / Essential Document Tracking

Track site documents, upload and reference critical documents, and seamlessly publish to the TI eTMF.

### Site View

Offers a comprehensive overview of all the clinical trial sites by region. Provides a centralized location for managing site-specific data and site users. Includes organizations, contacts, activities, milestones, plans, documents, communications, tasks, deviations, issues, and much more with the site view.

### Site Selection

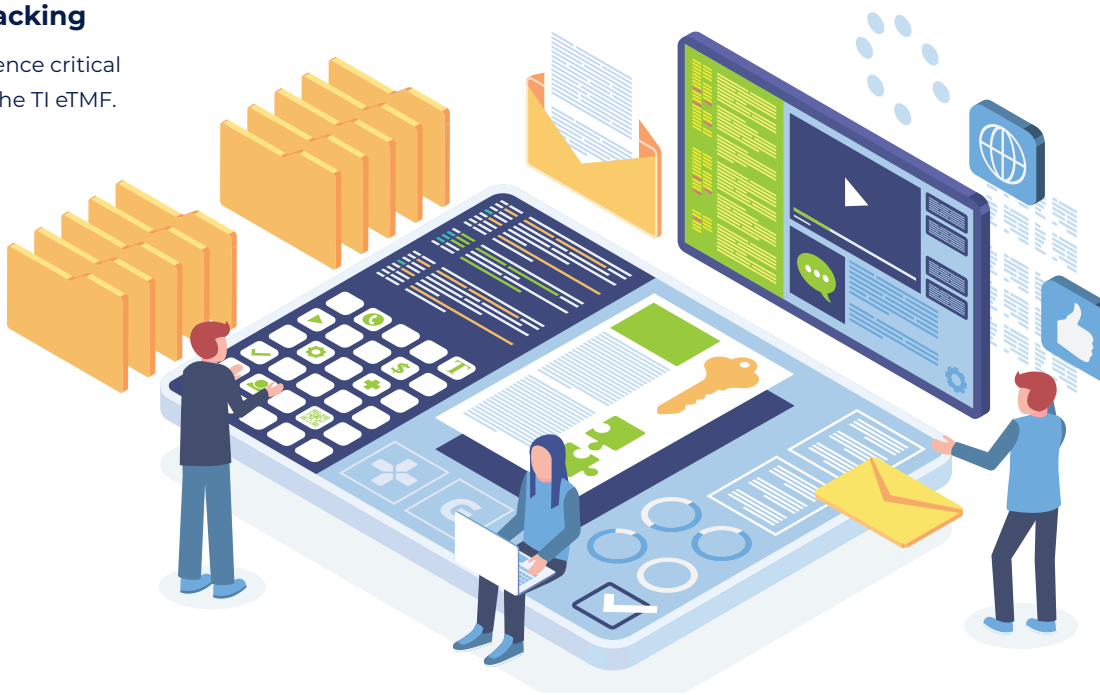
Site Selection to read the following: Provides increased oversight of the site selection process and start-up status. Allows for idenficiation, evaluation, and selection of potential clinical trial sites. TI CTMS Inter-operabililty with TI SSU module helps user expedite site timelines.

### Site Management

TI CTMS provides the ability to monitor sites anytime, anywhere, making it easier for CRAs and Study Managers to monitor clinical trial sites through the use of visual dashboards and site views.

### Site Visit Reports

Review individual sites progress with scheduled visits. Review planned and completed activities, protocol information, ICF tracking, deviations, and safety reports. Review screen failure rates, protocol visit windows, and required activities.



# THE eCLINICAL PLATFORM FOR GLOBAL PRODUCT DEVELOPMENT

Trial Interactive is a suite of integrated products that help study teams work efficiently and reduce risk across the clinical trial lifecycle, including:

- Flexible and Configurable to Requirements
- Expedites Site Activation and Document Work Streams
- Comprehensive, Easy-to-Use Workflows, Mobile eTMF
- Fast Implementation and User Onboarding / Adoption
- 21 CFR Part 11, Annex 11, ERES, GxP, and GDPR Compliant

## eTMF

A practical, secure, compliant, single access point for TMF documentation, supporting all essential document processes and reducing time, costs, and risks of TMF management.

## Collaborate

Collaborate on clinical and R&D documentation with shared workspaces for sponsors, CROs, and sites. Enforce project workflows to build and finalize study documentation such as TMF and eISF with remote monitoring and mobile document processes.

## CTMS

CTMS provides mobile-first trial management for planning and tracking, with specific attention to monitoring requirements.

## Study Start-Up

Accelerate site activation by simplifying the collection, completion, and finalization of critical regulatory documentation necessary for approval to bring a site online.

## GlobalLearn

Train site personnel and study teams, orchestrate virtual investigator meetings and ensure SOP compliance to reduce risks and increase team efficiency with a compliance-focused learning management system.

## eFeasibility

Speed site selection with a configurable survey module to contact, assess, and prequalify sites. Screen sites for inclusion and save surveys and collected data for future use.

## Site Portal & eISF

Reduce administration and improve speed and compliance for site personnel and study teams with an investigator site solution that facilitates digital investigative site binder processes.

## myTI

The mobile TI eTMF experience enables site document capture and key e-clinical workstreams anytime, anywhere.

## Quality Management

Handle your SOPs, CAPAs, and deviations, as well as clinical trial and R&D documentation, with an EDMS focused on quality and R&D document workflows.

## Safety

Simplify safety letter communication between sponsor, CRO, countries, and sites with a shared investigator site and safety team solution. Enforce safety reporting deadlines with a database of regulatory requirements.

## EXPERIENCE AWARD WINNING TECHNOLOGY

2023
Best Technology for Sponsors

TRIAL INTERACTIVE

AI-Enabled and Mobile:

**CTMS | eTMF | Site Solutions | LMS**

2023
Best Content Management Innovation

TransPerfect's Trial Interactive solution provides a collaborative, web-based platform for study start-up and eTMF that enables sponsors, CROs, IRBs, central laboratories, and other vendors to maintain and update clinical trial documentation in a secure online environment. Trial Interactive streamlines study timelines and reduces the administrative burdens of global clinical trials.

As part of TransPerfect's Life Sciences division, Trial Interactive is dedicated to working collaboratively with clients on a global scale, supporting the clinical trial life cycle with platform offerings including investigator database, e-feasibility, study start-up, eTMF and TMF management services, mobile eTMF, clinical document management and collaboration, compliance-focused learning management, pharmacovigilance and safety management, endpoint adjudication, and product licensing and alliance management.

For more information, please contact:
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