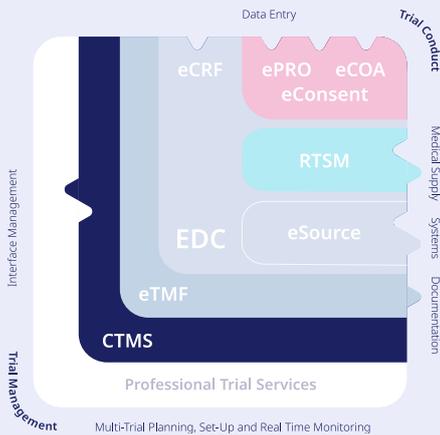


omnia CTMS



ADAPTIVE OVERSIGHT AND INSTANT INSIGHTS WITH OOMNIA CTMS

Navigate clinical trial complexities with confidence and precision

omnia integrates CTMS features designed to optimize your clinical trial management. omnia CTMS supports you in the planning, execution, monitoring, and reporting of clinical studies. Gain unparalleled insights and comprehensive oversight across your clinical trials.

- ✓ All data in one place
- ✓ Easy data management and tracking
- ✓ Adverse event monitoring
- ✓ Efficient budgeting

BENEFITS OF OUR UNIFIED CTMS

Discover the advantages of streamlined trial management



ALL DATA IN ONE PLACE

omnia features a clear and comprehensive overview of participant, investigational site, and trial information. All essential components – eCRF, RTSM, budgeting, eTMF, and monitoring – are conveniently accessible with a single login.



EASY USER MANAGEMENT

Information from contact lists is quickly accessible, and investigational sites and investigators can be easily selected. Study staff information can be prepopulated from existing profiles within the system.



INTERACTIVE OVERVIEW

omnia CTMS allows an easy oversight of the study with real-time data feeds and downloadable reports. Metrics include for example adverse event information, protocol deviations, or source data verification progress.



ERROR REDUCTION

All data and analytics are in one place, easily accessible, and automated. omnia can pull data from various forms and auto populate, reducing errors from manual input and reconciliation.



TIME SAVINGS

Monitoring visits can be scheduled directly through omnia CTMS, providing rapid access to information from contact lists. The tool allows for the efficient creation of budgets prior to the commencement of a study.



COST SAVINGS

Budgeting with CTMS allows sponsors to see how their funds are being used and if they could be used more efficiently.



POWERED BY

Wemedoo
Clinical Information Specialists

www.omnia.io/ctms

omnia
Infinite clinical trials

ADVANCED FEATURES OF OOMNIA CTMS

Unlock the full potential of your clinical trials

STUDY MILESTONE AND ENROLLMENT MANAGEMENT

FEATURES	DESCRIPTION
Key milestone and enrollment dates tracking	<ul style="list-style-type: none">Define key milestones and schedule planned completionTrack timelines and make adjustments as required
Recruitment information	<ul style="list-style-type: none">Track participant disposition by site, country, and trial, directly from the EDC

COMPREHENSIVE FINANCE MANAGEMENT

FEATURES	DESCRIPTION
Budget creation and management	<ul style="list-style-type: none">Create a study budget and budget grid, with expenses for various visits, examinations, and other trial related costs
Invoice processing and payment tracking	<ul style="list-style-type: none">Streamline invoicing with efficient invoice entriesAutomatic invoice generation from completed patient examinations
Cost tracking and analysis	<ul style="list-style-type: none">Track expenses against the budget in real-time

DOCUMENT MANAGEMENT

FEATURES	DESCRIPTION
Essential documentation tracking	<ul style="list-style-type: none">Automatically track essential documents with seamless eTMF integration
Site staff management	<ul style="list-style-type: none">Tracking of trial staff in an including activation and removalDefine trainings and track completion status
Site delegation management	<ul style="list-style-type: none">Efficiently delegate responsibilities with a site staff and delegation log
Automatic filing to the eTMF	<ul style="list-style-type: none">Save time by seamlessly filing documents directly into the eTMF

MONITORING MANAGEMENT

FEATURES	DESCRIPTION
Monitoring visits scheduling and tracking	<ul style="list-style-type: none">Monitoring visits can be scheduled via the CTMSFile essential documentation into the eTMFSet up automated workflow templates for appropriate users
Action item log	<ul style="list-style-type: none">Generate, assign, and track action items across, trials, sites, and staff members
Quick monitoring visit report creation	<ul style="list-style-type: none">Automatically create monitoring visit reports with data entered into the eCRF and other documents

EASY PARTICIPANT PROGRESS TRACKING

FEATURES	DESCRIPTION
Study visit tracking	<ul style="list-style-type: none">Tracking notifications for upcoming or out of window visits
Recruitment source tracking	<ul style="list-style-type: none">Optimize recruitment by tracking enrollment sources for each patient is recruited

EFFICIENT STUDY PLANNING AND DECISION MAKING

FEATURES	DESCRIPTION
Interactive study calendars and timelines	<ul style="list-style-type: none">Efficiently organize monitoring visitsCentral view of all scheduled visits and tasksCreated milestones with their expected dates
Fully customizable visualizations	<ul style="list-style-type: none">Customize visualizations for the specific needs and preferences
Adverse event tracking	<ul style="list-style-type: none">Track AEs to resolution with integrated graphical reports, facilitating analysis and critical safety information
Protocol deviation tracking	<ul style="list-style-type: none">Support action items with central accessible protocol deviations and real-time reporting