



THE FIRST TRULY UNIFIED CLINICAL TRIAL SOFTWARE

Understand why oomnia beats the competition with ease.

Save time and money while increasing data quality, simplifying study tracking, and achieving better outcomes.

INFINITE CAPABILITIES IN ONE SOFTWARE SOLUTION

More than just a clinical trial management system



All-In-One Solution

All professional clinical trial tools you will ever need fully integrated into one webbased software solution.



Infinite Scalability

Broaden research with limitless trials, storage, patients, users, sites, data points, documents, queries, ensuring expansion.



User-Friendly Interface

One simple user interface, no coding skills needed, drag-and-drop-functionality with a steep learning curve.



Data Security

You determine data storage and access preferences. We meet all regulatory standards, ensuring maximal security for data storage.



Real-Time Reporting

Enter data one time only for real time data analysis. Gain instant insights for infinite biomedical statistics.



Access From Everywhere

Access data anywhere & work offline. Minimal software costs with our browserbased SaaS solution, ensuring flexibility & efficiency.



Interoperability

Fully interoperable with other clinical trial systems. Import and export data with ease & connect third party solutions.

Professional

Assistance

All professional trial services

to ensure optimal trial setup,

management and trial

member training.



Infinite Trial Types

Realize all trials - Hybrid, decentralized, synthetic, real world data, global and even custom trial projects.



Convenient Customization

Customize EDC with dragand-drop eCRF-Builder in oomnia for effortless creation of documents, variables, and functionalities.

WATCH YOUR RESULTS





Better patient outcomes







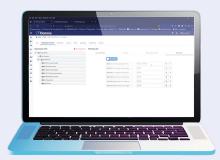
EVERYTHING INCLUDED IN ONE SOFTWARE

All tools included:

EDC | eCRF | RTSM | eTMF | ePRO | eCOA | CTMS | eConsent | Lab Management | Imaging | eArchive | eSource | and many more

Simple and competitive pricing. Pay per trial per month.





PROFESSIONAL CLINICAL TRIAL TOOLS

Discover the first all-in-one software solution enabling infinite clinical trials.

This integrated suite of tools will immensely simplify all processes involved in clinical trials. The following tools and numerous functions are fully integrated in the software. Using oomnia you will see that improving efficiency is easy.

EDC

Configuration · User access · Automated data integration and processing · IMP randomization directly within the eCRF document · Expert support team

RTSM

Complete lifecycle coverage for kit shipment and current status · randomization, allocation and accountability · Reduced manual data entry · Automated real-time reconciliations · Export of end-of-trial randomization report

eTMF

Insights through real-time reports · Rapid startup timeline · Exportable to fully validated eArchive · Convenient document management

Trial Conduct Data Entry eCRF **ePRO** eCOA electronic Patient Reported Outcome Medical Supply **RTSM** Interface Management EDC Electronic Data Capture Documentation **eTMF** electronic Trial Master File **CTMS** Clinical Trial Management System Trial Managenent **Professional Trial Services** Multi-Trial Planning, Set-Up and Real Time Monitoring

ePRO

Possibility for patients to directly report data · Involving electronic devices such as smartphones, tablets, or web-based platforms · Higher involvement of patients · Improved data quality

eCOA

Collection and documentation of clinical outcome assessments from patients, physicians or nurses · Involving electronic devices such as smartphones, tablets, or web-based platforms · Improved data integrity and efficiency

CTMS

Customized dashboard per user role · Standard study templates and logs · Key performance indicator (KPI) reports · Monitoring visit tracking · Timeline and milestone tracking · Contract management · Budget and payment management

PROFESSIONAL TRIAL SERVICES THROUGHOUT THE COMPLETE TRIAL LIFE CYCLE

Plan

- ∞ Protocol Development
- ∞ CRF Development
- ∞ Clinical Advisory and Scoping

Conduct

- ∞ Clinical Data Management
- ∞ Pharmacovigilance
- ∞ Risk-Based Monitoring

Close

- ∞ Biostatistics
- ∞ SDTM and ADaM datasets and define.xml
- ∞ Medical Writing



