



**OmniComm**  
eClinical Solutions for Life™

## TrialMaster®

Electronic data capture (EDC) has become the preferred method of capturing and cleaning data for Phase I – IV clinical trials. TrialMaster®, OmniComm's flagship EDC solution, provides depth of functionality wrapped in a user-friendly, web-based system. Leveraging the robust capabilities of a full clinical data management system, TrialMaster provides improvements in productivity and a more responsive site experience, backed by state-of-the-art hosting facilities and customer service and support.

TrialMaster, a solution designed with the site-user in mind, is both feature-rich and easy to use. A simple drag and drop study designer, an intuitive data capture web interface, on demand importing and exporting capabilities, task-driven workflow, streamlined query management, configurable metric-loaded standard reports, and a wizard driven Ad Hoc report builder complement this eClinical solution.

From a time and cost perspective, bio-pharma companies, both large and small, are struggling to efficiently manage early and later phase studies with a single EDC solution. OmniComm's TrialMaster has a proven track record of successfully addressing this challenge. Whether your objective is to deploy TrialMaster on a large scale global project or a single site Phase I study,

TrialMaster is a cost effective and efficient solution. For prospective CRO clients, OmniComm offers a "no money down" CRO fixed-pricing program, making it cost effective for CRO's and other partner companies to offer TrialMaster.

OmniComm is a global provider of customer-driven eClinical solutions to organizations that conduct life changing clinical trial research. OmniComm's growing base of satisfied customers is a direct result of the company's commitment to deliver products and services that ensure ease of use, faster study build, interoperability and better performance.



# TrialMaster®

## OmniComm's premier EDC solution

Key Features	Key Benefits
OmniConnect™ API.	Easy integration of TrialMaster to other clinical trial systems; IVR, ePRO, CTMS. Our validated API makes maintaining integration between point solutions more manageable, reducing IT costs.
Integrated AutoEncoding codes adverse events and concomitant medications against MedDRA and WHO Drug dictionaries.	Clean and simple interface speeds coding of verbatim terms while improving overall quality of output.
Ad-hoc Reporting.	A simple, wizard driven tool for creating and scheduling custom reports allowing authorized users to quickly and conveniently drill down into study data.
Export Utility which maps data to export domains creating repeatable custom exports, including ASCII, SAS and CDISC SDTM.	Integrated utility for creating custom exports and submission ready data in less time and cost than current methods.
Integrated, self-paced eLearning to train sites, monitors and data managers.	Swift user adoption as a result of self-paced training, a natural user interface, and task lists that streamline workflow.
Meets requirements for 21 CFR Part 11 and GCP.	Confidence that your data can be submitted for any global product registration study; backed by a complete audit trail.
Responsive edit checks that execute in real-time rather than waiting for the user to submit the form.	Reduction of inaccurate data: Real time edit checks result in fewer errors and queries for the data management team and streamlines data cleaning.
Scalable end-to-end solution for all study phases.	Cost effective EDC solution for early as well as late phase studies offering economies of scale and lower total cost of ownership.
Configurable standard reports.	Easy to access metrics and data listings with over 35 out of the box reports. Increases efficiency for site and sponsor reporting.

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