Fact Sheet



Clinical Trial Management System

Connected Intelligence

Enjoy the ease and transparency achieved through digital automation designed around persona experiences for traditional and decentralized trials. Leveraging IQVIA's deep expertise in clinical trial operations, IQVIA Technologies Clinical Trial Management System improves subject safety, data quality and operational efficiency, producing faster trials to database lock and speeding time to market. CTMS, part of IQVIA Technologies' Digital Trial Management Suite, is Intuitive, Intelligent, and Interoperable.

THE CHALLENGES

Traditional monolithic CTMS presents users with challenges to **find meaningful data buried** in cumbersome UI and confounding reports. This is compounded by **third party integrations with EDC**, **eTMF and other point solutions or partners**, which are difficult to use, expensive and require an army to build and maintain. Additionally, upgrading to a more intuitive approach may require purchasing an entirely new suite of products, even if you prefer to retain some of your current investments.

OUR SOLUTION

CTMS is part of IQVIA Technologies Orchestrated Clinical Trials, the end-to-end project-site-subjectcentric platform that:

- Provides an unparalleled data infrastructure, seamless connectivity, and intuitive design to drive smarter, faster trials with a configurable, scalable, SaaS-based solution
- Supports the needs of emerging BioPharma through global mega-trials in one integrated application built on a health-cloud platform
- Configures quickly using pre-built connectors and templates
- Intuitively tracks and logically structures
 information to enable life science teams to
 monitor and report on clinical trial progress
 and make better informed decisions, providing
 the best quality data while ensuring subject safety
- Uses AI and ML and is engineered for flexible business models
- Recommends the next-best action using built-in predictive intelligence
- Ensures your clinical processes, security, and compliance with regulatory guidelines are always current with several annual upgrades

IMPROVES DATA QUALITY AND DATA WORKFLOW

- Produces a single source of clinical trial data throughout the trial lifecycle
- Real-time tracking and event-driven actions for faster throughput
- Enables the management of more efficient overall trial timelines with data

IMPROVES CUSTOMER AND SITE RELATIONSHIPS

- Expedites site and vendor payment cycles
- Removes administrative burden to improve relations and preferred site status
- Provides greater transparency and increased compliance with FDA and EMA



Designed to intuitively track and logically structure information, CTMS enables life science teams to monitor and report on clinical trial progress and make better informed decisions, providing the best quality data while ensuring patient safety.



Configurable and easy to use

- Guided workflows and processes
- Intelligent assisted analysis
- SaaS-based application built on a health-cloud platform



Interoperable and scalable

- Adaptable components
- Adaptive to any trial size

 Emerging BioPharma to

 Mega Trials
- Global scale



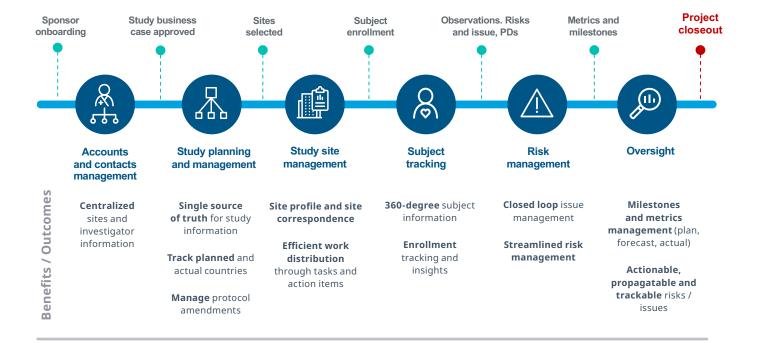
Better Clinical Trial Intelligence

- Provides holistic dashboard view
- Facilitates intelligent decision making
- Enables propagation and management of observations, issues and risks enabling predictability and compliance

CTMS benefits

Enjoy the ease and transparency achieved through digital automation designed around patients and sites for traditional and decentralized trials. IQVIA Technologies Clinical Trial Management System **improves productivity**, **study reconciliation**, **and data quality**.

OPERATIONAL VALUE STREAM



CTMS is one of three applications within IQVIA
Technologies' Digital Trial Management Suite.
Intuitive, Intelligent and Interoperable, our Digital
Trial Management applications work well on their

own but are even more powerful together, improving patient safety, producing faster trials to database lock, and speeding time to market. Visit IQVIA.com/OCT to learn more.

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