REMARQUE SYSTEMS

Clinical Trial Management System (CTMS)

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A simple, cost-effective, and easy-to-implement clinical trial management system (CTMS)

Remarque Systems CTMS makes monitoring sites easier through an intuitive interface designed with Clinical Research Associates (CRAs) in mind.

From one straightforward, at-a-glance home page, you can:

- Quickly access and track information like upcoming monitoring visits and tasks, as well as documentation status and deadlines
- Manage protocol deviations and action items
- Easily create, review, and approve monitoring visit reports

For site monitoring, Remarque Systems CTMS provides a simple way to:

SCHEDULE	MANAGE	TRACK
Schedule remote/central/ on-site monitoring visits for all global sites — from pre-study through site qualification, site initiation, interim monitoring, to closeout.	Manage all monitoring visits and visit documentation on one user-friendly platform for letter/report completion, review, and approval. Efficiently manage investigator contacts and site information.	Effortlessly track upcoming visits, tasks, and deadlines.

Remarque Systems CTMS features include:

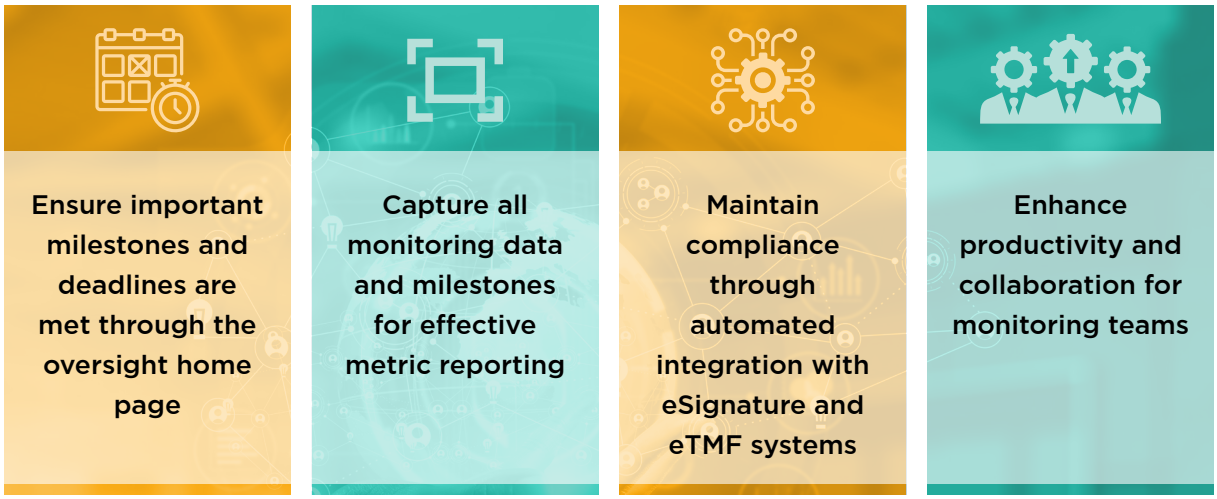
- Monitoring visit schedule management
- Template-based on-site, remote, and central monitoring visit documentation
- Report and letter completion/review/approval in one system
- Captures all monitoring data and milestones for effective metric reporting
- Protocol deviation, action item, and communication log management
- Site creation, contact management, and investigator database
- Online/offline functionality
- Electronic signature and eTMF integration
- Direct integration with Remarque's Study Start-Up (SSU), RealTime Monitoring, and Risk-Based Study Execution (RBX) functionality

The go-to solution for simple and effective monitoring oversight and management

Remarque CTMS is a centralized location for every aspect of site monitoring. With all the documentation you need for streamlined on-site, remote, and central site monitoring in one place, you can reduce report review and approval times by 75% when compared to standard methods.

Getting the data you need is easy. Quickly view crucial information like upcoming visits and track documentation deadlines, tasks, and metrics — as well as review and approve reports — all in one application.

For monitoring oversight, Remarque Systems CTMS provides a simple way to:



A simple system for monitoring sites

Remarque Systems CTMS is easy to implement and use. It integrates as needed with your current systems, allowing you to get going quickly without spending months on a complex setup. It is also fully configurable to match your existing templates and processes so you can optimize for faster, more efficient monitoring.

The easy path to better study oversight

Seamlessly combine Remarque CTMS with Remarque's other study management products (Study Start-Up/Real Time Monitoring/RBQM) to aggregate and analyze all site documentation and study data in one centralized location. Remarque Systems provides a simple, elegant, and complete solution for clinical trial professionals to manage their studies.

Looking for an easier way to manage all aspects of monitoring?

Book a demo today

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