FACT SHEET

CTMS by Octalsoft

Clinical Trial Management System

- Improve Data Quality
- Enhance Operational Efficiency & Productivity
- Get Actionable Insights
- Lower Trial Costs

To know more, email us at
info@octalsoft.com
inquiry@octalsoft.com

+1 (240) 547-4400
+91 (760) 098-3010

Schedule a **Demo** with our Product Specialist today!

Connect with us

www.octalsoft.com • INDIA | USA

CLINICAL TRIAL MANAGEMENT SYSTEM BENEFITS

Site Management

- Track study progress from start-up to close-out
- Standardize CRA trip reports and correspondence letters
- Automate email notifications to stakeholders for critical actions
- Track open action items, and protocol deviations
- Track subject visits and milestones with an integrated calendar
- User specific-to-do list tracker

Finance & Budget Management

- Manage and track trial budget
- Automate Investigator milestone payment based on contractual terms and SDV of the subject data
- Integrate and customize invoice approval workflow for site payments, travel expenses, and monthly project budget

Rapid Deployment & Quick Support

- Configurated and pre-validated system with abbreviated UAT for faster deployment
- Proactive focus on remaining available and customer-focused throughout the study execution
- Integrated support model with online feedback ticketing system, email & support team contact numbers specific for each deployment

> Octalsoft CTMS has transformed the way we manage our late-phase clinical trials. The system provides a comprehensive set of features that allow us to streamline trial processes and make informed decisions. From tracking study progress, managing site and patient data to generating reports, the system has everything we need to conduct successful clinical trials. What sets Octalsoft CTMS apart is its simple user interface. It is intuitive and easy to use. The ability to access the system from anywhere, at any time, has been a game-changer for us. The support we receive from the Octalsoft team has been excellent. They continue to be responsive and helpful whenever we have questions or need assistance. In addition to CTMS, we also leverage their IWRS system and we are quite satisfied with the tool and their delivery method

Dr. Dharmesh Domadia,
VP Global Client Operations
| Clantha Research

Integrated Monitoring Tools & Reports

- Track overall project status with the help of GANTT charts & study progress reports
- Generate activity plans and allocate resources
- Flexible reporting interface and dashboards to provide data summaries, charts, and visualizations
- Create and customize reports with filters on projects, tasks, subject status, and site visit reports

eTMF & Document Management

- Create a standardized & customizable DIA compliant eTMF structure
- Store and organize eTMF & non eTMF documents with version control
- Visualize overall document completion status through an intuitive color-coded dashboard
- Manage complete document creation life cycle online through eDOCS
- Map storing of site visit reports and protocol deviations into the respective eTMF folders

Seamless Integration

- End-to-end system integration with EDC and IWRS
- API with third-party applications/systems

Better Resource Allocation

- Make better decisions throughout the clinical trial process, leading to more efficient use of resources and faster time to market
- Increase productivity of CRAs by automating repetitive tasks and allowing real-time information sharing
- Create a sustainable competitive advantage

Mobile Application

- Interactive dashboards available for strategic decision making

OUR UNIQUE CUSTOMER SUPPORT STRUCTURE

Detailed User Manuals/Guides

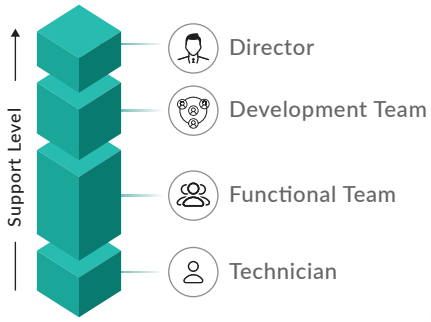
One Stop-Hassle Free Support

- Detailed user guides & training sheets for every product module
- Guided video tutorial sessions to address user queries
- Exhaustive FAQ sheets per product module

24 X 7 Help Desk Support

Our Specialists are Always Available to Resolve Your Queries

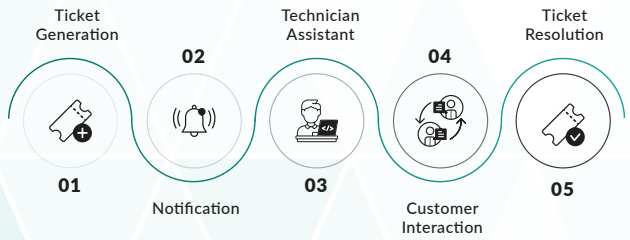
1. Customized screen for every user
2. Integrated online & offline support
3. Comprehensive repository of common queries
4. Module embedded chatbot



Embedded Agile Ticketing Feedback System

Your Query is Our Priority

Issue based (bug, enhancement, data rectification & others) ticket resolution system to address urgent queries on priority



AI Powered Module Specific Chatbot

Personalized Just for You!

1. Specific phone key number allocation to every client
2. Local & international support
3. Integrated online & offline support

Meet **Alex,**

Clinical Trial Lead

He works at a large-scale pharmaceutical company. With more than 10 years of experience in clinical research, Alex oversees the planning, execution, and monitoring of multiple clinical trials across various therapeutic areas. Alex's key responsibilities include managing study timelines, budgets, and resources, ensuring compliance with regulatory requirements, and maintaining effective communication with key stakeholders.

Despite having a team of experienced Clinical Research Associates (CRAs), Alex faces several challenges in managing clinical trials such as tracking study progress and performance metrics, coordinating with vendors and study sites, and ensuring timely and accurate data collection and reporting.

To overcome these challenges, Alex has adopted the Octalsoft CTMS. Our solution provided Alex with real-time visibility into study performance metrics, allowing him to quickly identify areas that need attention and take corrective actions. It also offered advanced reporting capabilities, enabling Alex to generate custom reports and dashboards to meet the needs of various stakeholders. Our solution also helped Alex to manage study timelines, track budgets, and resources with greater efficiency.

Overall, by adopting our CTMS solution, Alex has been able to focus on the big picture and make informed decisions to drive the success of clinical trials.

WHY CONSIDER OUR eCLINICAL SOLUTION?

Single Unified Mobile-Enabled Platform

End-to-end physically & logically secured cloud solution that effortlessly manages data collection, IP supplies, randomization, study compliance, and documents

Regulatory Compliant

Compliant with all current & highest regulatory guidelines:

GDPR | FDA 21 CFR Part 11 | EU COMPLIANT Annex 11 | GAMP 5 | cdisc | HIPAA COMPLIANT | GxP | ICH GCP

Patient Centric

Integrated ePRO & eConsent enables you to conduct patient-centric virtual trials to access diverse population all over the world for smarter and faster trials

Audit Ready Database

End-to-end physically & logically secured cloud solution that effortlessly manages data collection, IP supplies, randomization, study compliance, and documents

Global Mega Trials in Multiple Languages

Experience conducting global trials for all types of studies and trial phases with US FDA Submission | EMA Submission | DCGI Submission

Fast Study Startup

Shortest build times with over 90% of studies deployed within 2-4 weeks. Easy-to-use trial builder allows quicker configuration of eCRFs

Configurable, Customizable, Cost Effective & Easy to Use

The platform can easily integrate with your existing system and can also get designed as per your requirements for increased flexibility, mobility and scalability

Unparalleled Global Customer Support

End-to-end physically & logically secured cloud solution that effortlessly manages data collection, IP supplies, randomization, study compliance, and documents

Our Values

For 15+ years, we've mastered the art of Simplifying Technology so that greater discoveries can happen sooner. As a leading eClinical Solution provider, we've always invested in the right expertise, experts, and technology that keeps us, and you at the forefront of science and medicine. As a trusted tech partner for some of the world's leading clinical trial initiatives, our values of innovation, reliability, and client satisfaction guide everything we do.

To know more, email us at info@octalsoft.com inquiry@octalsoft.com

+1 (240) 547-4400
+91 (760) 098-3010

Schedule a **Demo** with our Product Specialist today!

Connect with us

www.octalsoft.com • INDIA | USA