FACT SHEET

CTSM by Octalsoft

Clinical Trial Supply Management

- Real-time Inventory Insights
- Supplies Accountability
- Enhanced Visibility
- Process Automation

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!

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CLINICAL TRIAL SUPPLY MANAGEMENT BENEFITS

Inventory Optimization

- Octalsoft's system allows for real-time tracking and optimization of clinical trial supplies, ensuring that sites have the right amount of investigational products and ancillary supplies at all times.
- It minimizes overstocking, reduces waste, and prevents supply shortages, ultimately saving time and resources.

Forecasting and Demand Planning

- The system includes advanced forecasting and demand planning tools, which use historical data, patient enrollment predictions, and other factors to accurately anticipate future supply needs.
- It ensures that sponsors can proactively address potential supply challenges.

Site Management

- Octalsoft's platform provides comprehensive site management capabilities, enabling users to efficiently distribute supplies to various clinical trial sites.
- Users can easily track shipments, monitor site inventory levels, and manage site-specific requirements, improving overall trial efficiency.

Regulatory Compliance

- The Clinical Trial Supply Management System is designed to assist sponsors and sites in adhering to regulatory requirements.
- It helps with the management of labeling, documentation, and quality control, ensuring that clinical trial supplies meet the necessary regulatory standards and compliance.

Shipment Tracking and Management

- Users can track the status of shipments, including the receipt, storage, and distribution of supplies.
- The system helps ensure timely and accurate delivery to trial sites, reducing the risk of delays or errors.

Real-time Reporting and Analytics

- The system offers a comprehensive reporting and analytics module that provides stakeholders with real-time visibility into supply chain performance.
- Users can generate customizable reports and dashboards to monitor key performance indicators, make data-driven decisions, and adapt to changing trial dynamics effectively.

> Efficiently manage the complex tasks of end-to-end clinical trial supplies and logistics processes while ensuring compliance with regulations and Good Manufacturing Practices (GMP). Robust inventory management tools automate the depot management processes of tracking and tracing both regulated and unregulated inventory and shipments, across multiple clinical sites and warehouse locations.

Krunal Bhatt

Technical Manager | Octalsoft

OUR UNIQUE CUSTOMER SUPPORT STRUCTURE

Detailed User Manuals/Guides

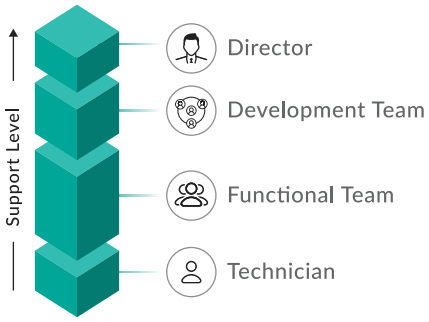
One Stop-Hassle Free Support

- 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

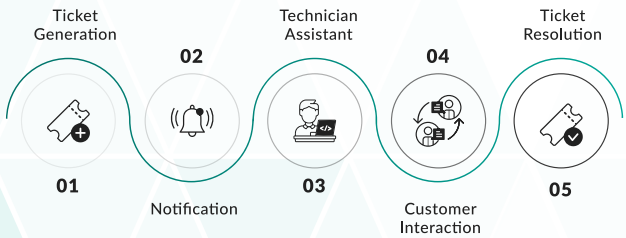
1. Customized screen for every user
2. Integrated online & offline support
3. Comprehensive repository of common queries
4. Module embedded chatbot



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 **Indira,**

Clinical Trial Manager

She works as a Clinical Trial Manager at a mid-sized pharmaceutical company specializing in clinical research

Indira's primary goal is to efficiently manage all aspects of clinical trials, from study initiation to completion. She is constantly looking for ways to optimize processes, reduce administrative overhead, and ensure trials run smoothly and within budget.

Octalsoft's CTSM system streamlines clinical trial management, automating workflows and reducing administrative burdens. It provides a centralized platform to oversee trial operations, helping Indira optimize processes and allocate resources effectively. It offers version control, audit trails, and secure data storage to reduce the risk of compliance issues. The system's robust security features, including encryption and user access controls, align perfectly with Indira's commitment to protecting sensitive clinical trial data. It ensures data security and patient confidentiality.

Octalsoft's CTSM system is well-suited to help Indira efficiently manage clinical trials, maintain regulatory compliance, ensure data security, and streamline communication with research teams and stakeholders. This allows her to focus on strategic decision-making and contribute to the success of clinical trials at BioPharm Innovations.

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.

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