


PHARM

Industry


CANADA

Location


>\$2B

Investment in R&D


>100

Clinical Trial

Client Requirement

- Manage and monitor **multiple clinical trials** across the globe with the highest level of accuracy and precision.
- Ensure patient safety, accurately assess and report outcomes, and **comply with global regulatory requirements**.
- Organize and store documents in a **CDISC-compliant eTMF structure**.
- Customize **country-specific** finance budget sheets.
- Convert **labor-intensive** clinical operations management into an **effortless and seamless** process by deploying CTMS.



Our Solution

Octalsoft provided a customized & centralized system for capturing trial data, monitoring trial progress, generating reports, and addressing open action items. We currently host a database of 1000+ trials, which help maintain clinical operations and subject-specific data in compliance with the highest regulatory standards. With Octalsoft CTMS, the client:

- ✓ Made real-time, data-driven decisions, increased the efficiency of the business processes, and contained costs, resulting in timely trial completion and license filings.
- ✓ Increased productivity of the CRAS by automating tasks, allowing real-time information sharing, and effective time management.
- ✓ Generated an intensive database of verified Clinical Trial Sites and Principal Investigators, giving it a sustainable competitive edge
- ✓ Enabled integration of IWRS & EDC for subject-specific critical data points. The data points included but were not limited to subject statuses & visit progression, recruitment efforts, and subject safety.
- ✓ Accessed flexible exports, smart reports, dashboards, and multiple reporting tools as per the management requirement.
- ✓ Improved document management and quality processes through Octalsoft eTMF.
- ✓ CTMS Finance Module helped with online tracking of pass-through and non-pass-through study costs and investigator milestone payments, and claim management in case of SAES

To know more,
email us at

info@octalsoft.com
inquiry@octalsoft.com

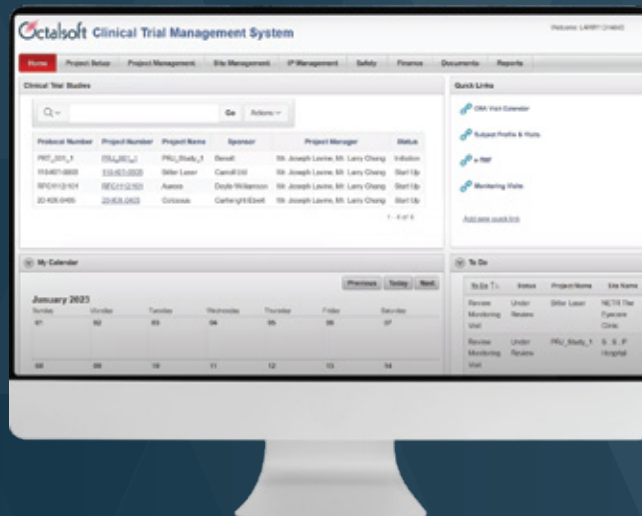
+1 (240) 547-4400
+91 (760) 098-3010

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our Product Specialist
today!



Module Features

- Clinical Site Visit Reports
- Milestones Planning and Tracking
- Resource Planning and Tracking
- Action Items & Deviations
- Finance Management for Trial Budgeting and Payments
- Site Budget and Payments



PRE-DEVELOPMENT

01

SYSTEM DESIGN & DEVELOPMENT

02

SYSTEM VALIDATION

03

SYSTEM DEPLOYMENT & SUPPORT

04

Study Build, Validation & Deployment Process

- Site Monitoring (Site Qualification, Selection, Initiation, and Closure)
- Travel Expenses & Timesheet Tracking
- Regulatory Documents & Compliance Tracking
- Training Management
- Adverse Event Tracking
- Integration with Other eClinical Solutions

Our Key Platform Benefits

Modularized & Integrated Platform With Single Sign In

Compliant With Highest Regulatory Standards

Unparalleled Customer Support

Easily Configurable

Cost Effective

Simple & Intuitive User Interface

To know more,
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