

# eClinical Dashboard



Study Setup &  
Configuration



Patient Recruitment &  
Management



Electronic Data Capture



Data Analysis &  
Reporting

To know more,  
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Schedule a **Demo** with  
our Product Specialist  
today!



Connect  
with us

# eCLINICAL DASHBOARD BENEFITS

## ● Data Centralization

- Centralize various data sources, providing a single location for clinical trial data
- Researchers and stakeholders can access near-real-time data.
- Data visualization tools that help users quickly understand and interpret complex clinical trial data, trends, and patterns.
- Create customized reports and metrics tailored to specific research needs.

## ● Efficient Site Management

- Monitor clinical trials remotely, reducing the need for physical site visits and improving efficiency.
- Track protocol adherence, ensuring that the trial is conducted in accordance with the study plan.
- Manage and coordinate activities at multiple trial sites more efficiently.

## ● Streamlined Trial Setup

- Streamline the setup of clinical trials, including defining study protocols, eligibility criteria, and randomization processes.
- Our eClinical Dashboard offers end-to-end trial management capabilities, from study setup and patient recruitment to data collection and analysis.

## ● Efficient Patient Enrollment and Data Capture

- Identify eligible patients, manage enrollment, and track their progress throughout the trial. We believe that technology should be accessible to everyone.
- Our user-friendly interface is designed with simplicity in mind, making it easy for both professionals and newcomers.
- Collect patient data electronically, reducing errors and enhancing data quality. Stay informed with real-time access to critical trial data. Monitor progress, track patient participation, and make informed decisions on the fly, improving the efficiency of your trials.

## ● Data Analysis and Reporting

- Generate insightful reports and analytics to aid in decision-making and data-driven insights. We understand the importance of data security and compliance in healthcare.
- Octalsoft's eClinical Dashboard adheres to industry-leading security standards and complies with all relevant regulations.

## ● Integration Capabilities

- Seamlessly integrate with other clinical trial systems and databases for a unified experience. Every clinical trial is unique, and your technology solutions should reflect that.
- Our dashboard is highly customizable to adapt to your specific needs and can scale as your research projects grow.

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Monitor progress, track patient participation, and make informed decisions on the fly, improving the efficiency of your trials. Our eClinical Dashboard is designed to empower healthcare providers, researchers, and organizations with the tools they need to streamline clinical trial management and data analysis. Octalsoft's eClinical Dashboard adheres to industry-leading security standards and complies with all relevant regulations, ensuring the confidentiality and integrity of your data

Krunal Bhatt  
Technical Manager | Octalsoft

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# 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



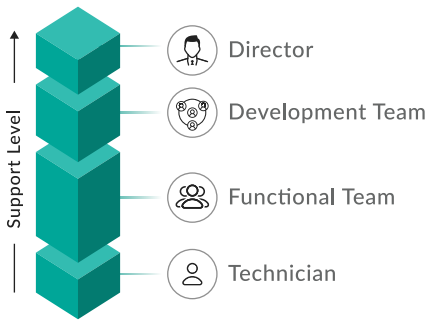
Exhaustive FAQ sheets per product module



## 24 X 7 Help Desk Support

Our Specialists are Always Available to Resolve Your Queries

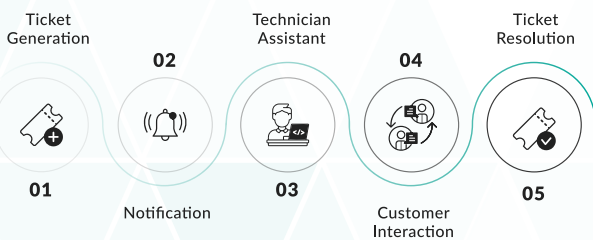
- 1 | Customized screen for every user
- 2 | Integrated online & offline support
- 3 | Comprehensive repository of common queries
- 4 | Module embedded chatbot



## 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 Emily, Clinical Research Manager



She works at a mid-sized pharmaceutical research organization

Emily's primary goal is to efficiently manage all aspects of clinical research, from study initiation to closeout. She seeks to streamline processes, reduce administrative overhead, and ensure that trials run smoothly and within budget.

As a research manager, she is responsible for data analysis and reporting. She is keen on having access to real-time data, analytics tools, and dashboards to make informed decisions and accelerate research progress.

Octalsoft's eClinical Dashboard streamlines clinical research management, automating workflows and reducing administrative tasks. It provides a centralized platform for overseeing trial operations, helping Emily optimize processes and allocate resources effectively. The eClinical Dashboard offers real-time data access, analytics tools, and interactive dashboards that empower Emily to make informed decisions quickly. It accelerates data analysis and reporting, ultimately expediting research progress.

In summary, Octalsoft's eClinical Dashboard is perfectly suited to help Emily efficiently manage clinical research, enhance data analysis and decision-making, maintain regulatory compliance, and streamline communication with research teams and stakeholders.



# 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:



## 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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