

ClinRT Global Services is a technology-driven provider of Interactive Response Technology (IRT) solutions. We help sponsors efficiently manage clinical trials across all phases and geographies. Headquartered in Pune, India, we are trusted by global pharmaceutical and biotech companies.



Rapid Deployment and Customizability

- 4-Week Commitment, Build to Go Live
- Design to Configure Complex Designs of Clinical Trials in a Short Time
- Robust Subject Management Module to Handle Complex Randomization & Protocol Requirements
- Inbuilt Configuration for Different Randomization Designs, Cohorts, Strata, Subject Subgroups, Complex Visit Schedule
- Flexible to Accommodate Changes Due to Protocol Amendments



Inventory Tracking

- Kit Tracking From Origin to Destruction via Robust Supplies & Accountability Module
- Automatic Supplies With Inbuilt Buffer and Prediction Strategies for Ordering Supplies
- Built in Site to Site Transfer Functionality
- Built-In Retention Sample Management Functionality as per USFDA Guidelines
- Built Functionality for Kit Expiry Update & Kit Status Change



User Experience

- Real-Time Customer Reporting Through Configurable Reports & Dynamic Dashboards
- Robust Module for Notifications and Alerts for Notifying Key Events and Transactions to Respective Stakeholders & Decision Makers During Study Progress



Expert Support

- A Technical Team With Strong Domain Knowledge to Facilitate Expedited Protocol Configuration in the System
- User-Friendly Tool for Technical Troubleshooting
- 24x7 Helpdesk Support

Key Distinctive Features of iClinRT®

Contact us



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Reliable IRT Service Provider for Global Clinical Trials



Customized and Scalable IRT Platform for Managing Global Clinical Trials



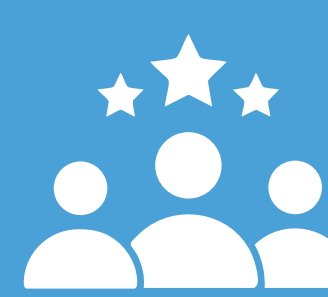
Intelligent Interactive Innovative



40+ Years of Consolidated Experience of Leadership Team in IRT & CTSM Domain

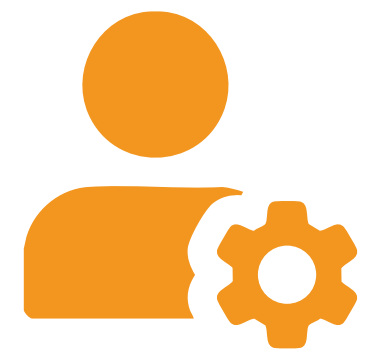


1000+ Clinical Trials Support Experience of Leadership Team



500+ Clients Catered with IRT & CTSM Services

'iClinRT' is a registered trademark of ClinRT Global Services



SUBJECT MANAGEMENT

Effortlessly manage different visit schedules, tailored to specific treatment protocols. Our intuitive interface ensures seamless coordination and communication, enhancing the overall subject experience while maintaining data accuracy and compliance.



Randomization

Ensure unbiased subject assignment with our robust randomisation capabilities. iClinRT® provides advanced stratification options, including site-based stratification, enhancing study precision and integrity. Tailor treatments based on real-time data, optimising the trial's efficiency.



Drug/Kit Assignment

Simplify logistics with our kit management system. Track, manage, and assign kits with precision, ensuring timely delivery to trial sites. Our platform supports both bulk and discrete kit types, providing unmatched versatility in managing your trial's inventory needs.



Drug Supply Management and Auto Shipments

iClinRT® streamlines your entire supply chain from procurement to distribution with real-time inventory tracking and automated shipments. Set predefined triggers and let the system handle timely deliveries, reduce costs, cut wastage, and keep your trials running smoothly without delays or disruptions.



Subject Unblinding

Provides process-mapped feature/functionality for "Unblinding Subjects" in medical emergencies or when trial subject safety is at risk. The feature is built to ensure participant safety, maintaining trial integrity & protecting against unintentional unblinding.



Drug Expiry Date Management

Maintain strict quality control with our kit expiry date management feature. Receive alerts and notifications well in advance, allowing proactive management of expiring kits. Minimise wastage and ensure the availability of vital supplies throughout the trial.



Drug Accountability

The functionality tracks the entire lifecycle of investigational drugs from birth to grave. This includes tracking of investigational drugs throughout the entire supply chain from packaging & labelling, distribution to clinical study sites, dispensing to trial subjects, return from trial subjects, reconciliation & drug compliance and destruction. The module provides a centralised and auditable system. This functionality enhances compliance with regulatory requirements and reduces the risk of errors and discrepancies.



Study Reports and Dashboards

iClinRT® provides access to a wide range of study reports, offering real-time updates on enrolment, randomisation, subject unblinding, patient compliance, site performance, subject visits, drug supply management, and more. These detailed reports support in-depth analysis and regulatory compliance. Dynamic dashboards visualise real-time data, enabling quick decision-making and efficient study management. Both reports and dashboards are customisable to display specific data based on country, depot, site, user role, etc., allowing for focused and flexible monitoring.



Notifications and Alerts

Provides configurable notifications and alerts to applicable stakeholders in the clinical trial process, including sponsor, study sites, PI, CRO, drug depots, etc, about critical and important events such as screening, randomisation, supply shortages, emergency unblinding, cautions & caps to facilitate decision making for smooth conduct of trial. This helps to improve SCM & facilitate emergency responses.



Helpdesk Management

Helpdesk provides a centralised point of support for users and stakeholders involved in a clinical trial. This support covers various aspects, including technical issues with the IRT system, protocol-related questions, and general inquiries about the trial process.