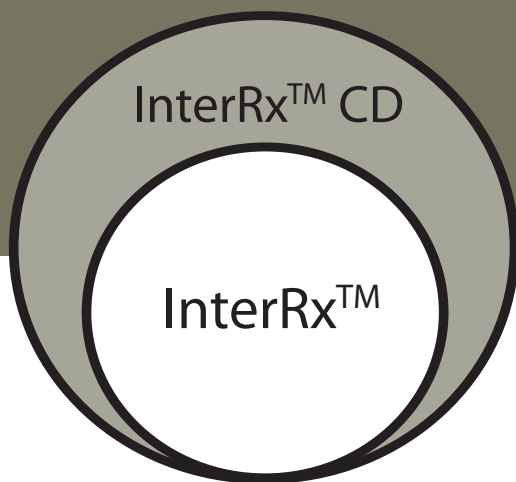


## InterRx™ CD

InterRx™ CD utilizes the best practices of an experienced clinical team and InterRx innovation to move your trial from protocol to FDA submission quickly, safely, and effectively. InterRx CD delivers a complete digital trial organization to biotechnology and to small pharmaceutical companies that need a clinical trial that is completed by outside resources and defines the financial and time requirements of the trial from day one.



### InterRx CD In The Field

Site investigators and coordinators enter data through Web-enabled dCRFs installed on a Tablet PC which is provided as a part of **InterRx™ MobileTrial** system. When cradled, the tablet connects to MobileTrial site server, which replicates and synchronizes data with the central database. This allows multiple coordinators and investigators to work online or offline and to pass their threaded data securely to the central trial database with the highest data integrity. All source from laboratories, imaging, or associated devices can be acquired with the Tablet PC and shared across the study teams, eliminating the requirement for redundant paper or physical source. InterRx CD's clinical team of field and medical monitors review data and source, ensuring efficient source verification, query resolution, analysis, and electronic packaging for further biostatistical review and eventual submission.

// **InterRx CD combines the best of InterRx with a streamlined study team to create a fully outsourced clinical research organization that is available on a moment's notice.** //

### What is Digital Trial Management?

Digital Trial Management (DTM) goes beyond electronic data capture and basic trial management systems by initializing, capturing, managing and delivering trial results in a digital format. In DTM, all source documents related to a trial, along with secondary capture and trial operational metrics are part of the solution - DTM reduces the need to keep secondary systems to visualize the trial's operational and clinical effectiveness.

### Fully Outsourced Digital Trial Management

InterRx CD even goes beyond our standalone digital trial management solution to fulfill the need of a readily available digital research organization at a moment's notice -

- Effectively control costs through planned trial start, team, management, and close.
- Plan capacity from the beginning to anticipate trial changes and additional sites.
- Eliminate slow trial start times and study initiation.
- Start quickly, with auditable, quality results. InterRx CD combines experience and an unbeatable research platform to save time and lower costs through efficient, effective work.
- Audit-ready at study start.
- Get exactly what you need to complete the trial safely and effectively so you can focus your internal resources on what you do best.

# THE DIFFERENCE

## Realtime Visibility

Sponsors have complete visibility to the trial's projected enrollment completion, patient safety, trial team activities, and site effectiveness with **InterRx™ TrialControl** dashboard which tracks the health and progress of the trial. Points of risk are identified and can be avoided in order to keep the trial going forward, or to stop a poorly executed trial.

## InterRx CD Clinical Development Organization

InterRx CD does not require a large clinical team to complete the trial. The InterRx CD team's expertise combines decades of clinical experience on hundreds of clinical trials with InterRx know-how to get to submission. The InterRx CD team remains a part of the solution for the life of the trial. As a result, you reap the benefits of the best service with the best results – a quality trial, in compliance, on time, within a fixed budget.

## Partnership

Our commitment to you does not end with software. InterRx CD immediately engages when the approved protocol arrives and ends with digital submission to the FDA. Strategic, technical, and clinical support is essential to getting to market, and at the core of InterRx CD is a valuable partnership that understands and commits to your need of getting to market.

## One Stop, Anytime, Anywhere

The InterRx platform was created for clinical research professionals by seasoned clinical research professionals who have partnered with leading technologists to leverage the latest technology advances for clinical research. The InterRx platform was designed to be the heart of any clinical trial, with one trial technology, stored and managed in one place, and supported by one company. The InterRx CD team is expertly trained in the use of the InterRx platform to ensure maximum efficiency throughout the life of the trial.

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## InterRx CD Rapid Trial

InterRx Clinical Research Associates (ICRAs) average over ten years of overall experience. Each ICRA is fully trained in FDA regulations, ICH guidelines, GCP/SOP compliance and with the InterRx platform. Rapid Trial combines technology with solid, best practices in clinical management to move a trial swiftly and efficiently to an evaluable conclusion.

### Rapid Trial

- Quickly implements trial initiation activities
- Provides complete trial monitoring
- Implements best strategies in project management
- Focuses on medical monitoring and drug safety
- Supports regulatory services and medical writing
- Reviews trial conduct and site management activities
- Completes trial closeout
- Assures trial quality at start, during lifecycle, and to conclusion

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## Get Started, Get Done

With Rapid Trial, you or your approved IRB can send your protocol electronically and it will be demonstrated to you as a full digital trial – with your sites, professional team, and hardware – in as little as 10 business days. Your InterRx CD team can complete a trial with all source information right at their fingertips, ensuring short timeframes between close, evaluations, analysis, and preparation for submission.

**Contact us today to see your trial in action with InterRx CD.**

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