PERCEPTIVE MyTrials®

A PAREXEL® eClinical Platform

IMPACT® CTMS

PULLING IT ALL TOGETHER

KEY FEATURES

- Full-featured CTMS providing trial planning, start-up, monitoring, financial administration, supplies tracking and investigator management functionalities
- Global deployment including international language and currency support
- Flexible monitoring tool, available both online and offline
- Advanced site selection capabilities including investigator profiling and search functionality based on historical trial performance and customer-defined metrics
- Configurability by trial type, allowing the creation of multiple trial types, controlling the visibility of menus and milestones, determining the level of data entry in use and enabling which fields are in use on each screen
- Application Program Interfaces (APIs) to enable automated data import from any clinical system

KEY BENEFITS

- Different study types can be set up to reflect different levels of tracking for different phases, therapeutic areas and partners
- Single, centralized system containing all operational trial data
- Accurate and consistent information to facilitate timely and informed decision making
- Eliminates informational silos and promotes efficient cross-departmental processes resulting in accelerated drug development cycles
- Automated integration with ClinPhone® RTSM and DataLabs® EDC applications eliminates need for re-keying data and discrepancy resolution
- Available as a software-as-a-service (SaaS) application enabling rapid and flexible deployment without high upfront costs or ongoing infrastructure investments
- Highly configurable and adaptable to sponsor workflow and terminology
- Active user community that provides input for product direction and promotes knowledge sharing among IMPACT CTMS users

Part of the Perceptive MyTrials® framework, enabling integration with clinical trial software applications to help users plan, design and conduct clinical trial programs in a single place

Value of a Single Centralized System

Clinical trials constitute the single most expensive component of the whole drug development process. In particular, clinical trial operations represent one of the most resource-intensive areas within a biopharmaceutical company and often involve tedious, manual processes for collecting, aggregating and rationalizing information from a wide variety of data sources. A CTMS yields immediate efficiencies and cost savings for clinical operations.

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IMPACT® CTMS

By providing a single, centralized system to orchestrate operational and administrative activities, a CTMS allows biopharmaceutical companies and CROs to intelligently manage the complexities of clinical trials. Instead of relying on disparate "islands" of information, often in direct conflict with each other, CTMS users benefit from sharing consistent trial data across the entire enterprise. This allows the sponsor or CRO to streamline processes, appropriately respond to current statuses and proactively manage their trials. IMPACT CTMS is an enterprise-wide solution used by many top global pharmaceutical companies and CROs to plan, administer and track every aspect of clinical trials. It is available as an on-demand, software-as-a-service (SaaS) application, enabling customers to enjoy rapid and flexible deployment of our leading CTMS solution without high upfront costs. Customers can also choose to implement IMPACT CTMS as an on-premise solution and install it within their operating environment. With one of the largest user bases and over 16 years of proven success, you are in the hands of true CTMS experts when you join the IMPACT solution community.

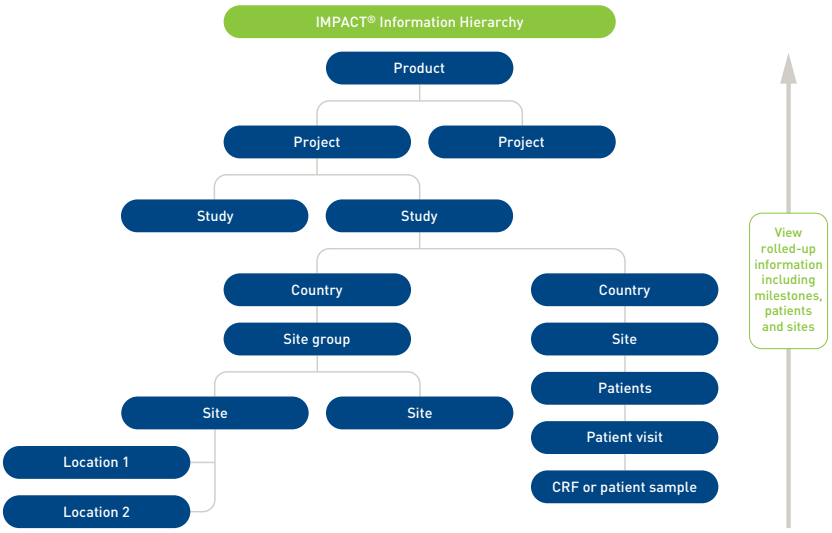
Making It Easier for You

IMPACT CTMS has been developed with flexibility and configurability in mind. The exceptional breadth of functionality has been designed to adapt to sponsor-specific workflows and terminologies. With IMPACT CTMS, clients do not have to modify their working practices and compromise with inflexible or rigid functionality. The system can manage trials of any complexity and support organizations of any size and global reach with complete flexibility to configure different study types for the different needs of each customer.

IMPACT CTMS has a large, global user organization—the IMPACT User Group (IUG). The IUG is an independent organization run by users for the benefit of the user community. It is a forum for exchanging knowledge on IMPACT CTMS usage and a direct channel for feeding back product needs and direction to the Perceptive MyTrials eClinical Suite. The IUG annually holds two formal, in-person meetings—one in Europe and one in North America. In addition, there are special interest groups that meet throughout the year to exchange information on specific subject areas and provide targeted feedback. For us, customers are not just nameless faces; they are part of the close-knit community of biopharmaceutical professionals in a meaningful, long-term partnership.

The IMPACT CTMS solution is a key part of our leading integrated eClinical Suite. The automated integration with the ClinPhone RTSM and DataLabs EDC applications allows users to seamlessly share trial data. Enabled through our unique integration platform, IMPACT CTMS users enjoy accurate and timely data without having to duplicate activities. The integration of IMPACT CTMS with RTSM and EDC enables your entire organization to benefit from the true power of CTMS.

The IMPACT CTMS solution is an integral part of Perceptive MyTrials® through which we are able to converge our integrated suite of clinical trial software applications.



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