

# TrialWorks® CTMS

## Are you seeing the full picture?

### Key features

- **Comprehensive enterprise-wide trial management**
- **Web-based, with offline CRA/monitor access**
- **User-friendly and intuitive interface**
- **Rapidly implemented company wide**
- **Over 175 standard reports tailored to all key departments**
- **Export to Microsoft Word, Outlook, Excel and ClinicalTrials.gov**
- **Seamless integration with ClinPhone® RTSM (Randomization and Trial Supply Management) and DataLabs® EDC for sharing of trial data**
- **Robust, flexible security including study and group level permissions**

### Trial Management

TrialWorks® is a powerful, user-friendly application designed to track, manage and report on virtually every aspect of your clinical study programs. TrialWorks is an enterprise solution and gives you full visibility of your clinical trial information.

TrialWorks provides comprehensive trial management data ranging from high level, consolidated metrics appropriate for senior managers to meaningful drill-down reports to support project teams. For example, financial managers can view the total costs and expenditures across clinical programs while a study project manager can view up-to-date enrollment figures and other study details. The underlying relational database allows this information to be grouped and filtered at various levels including trial, country, site/investigator and more.

TrialWorks is used to track thousands of trials by companies around the world. With TrialWorks, clinical trial management is easier, faster and more efficient.

### Flexible Solutions

#### Versatile application

TrialWorks is fully configurable to suit your needs. TrialWorks is used by biopharmaceutical companies, CROs and other specialized research organizations with functionality to fully meet the specific needs of each type of organization. It supports trials of all sizes, from large multi-national to single site studies.

TrialWorks can be used in conjunction with the Monitoring Module, which provides offline capability to monitors/CRAs, and with the Universal Data Transfer, which imports patient data from virtually any other clinical technology, including IVR/IWR, EDC and CDMS.

## TrialWorks® CTMS

### Flexible implementation

Depending on your organization's needs, TrialWorks can be licensed and installed at your facilities as an enterprise solution or implemented as a hosted solution via the TrialWorks e-Service.

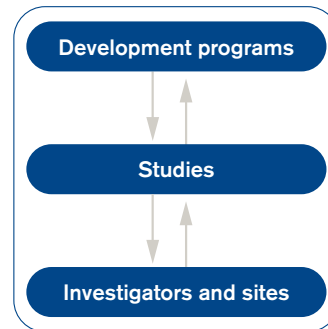
### See the Full Picture

#### Ease of Use

- Tracks and reports on every aspect of day-to-day trial management
- Delivers relevant information to all study stakeholders
- Promotes operational and workflow standards
- Ensures timely investigator payments and accurate financial reporting
- Imports and/or exports data to IVR/IWR, EDC and CDMS

TrialWorks is organized to allow users to rapidly navigate through the system, which promotes consistent and efficient work processes as well as high user adoption. The system is also built to meet the management needs of all study stakeholders. TrialWorks is one of the most user-friendly clinical trials management systems available.

Monitoring • Satellites • CRAs • Alerts • Vendors • Correspondence • Site certifications  
CDAs • Patient visits • IRBs • Coordinators • Investigator payments  
Regulatory documents • CRF pages • Assessments • Schedules • Patients



Outlook • Word • IVR • Reports • Excel • Clinicaltrials.gov