



# Target e\*CTMS®

For more information,  
Contact:

Jules Mitchel, MBA, Ph.D., President  
Target Health Inc.  
261 Madison Avenue, 24<sup>th</sup> Floor  
New York, NY 10016  
Tel 212.681.2100  
[julesmitchel@targethealth.com](mailto:julesmitchel@targethealth.com)

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## Target e\*CTMS®

### What Is It?

Target CTMS™ is a secure, **USER FRIENDLY**, web-based clinical trial management system which enables users, depending on their roles and responsibilities to manage a clinical trial, all within a web browser and without installing any software. Target CTMS® reduces the need to distribute documents via email and is ideal for companies wanting their own system to manage sponsors, sites, IRBs, study startup, and study progress. Together with Target e\*CRF® and Target Document, Target CTMS™ fulfills the trilogy of products needed to manage a clinical trial from cradle to grave.

Clinical trial management involves tracking of regulatory documents and clinical trial operations. For example, all clinical trials require the same regulatory documentation (e.g. signed Form 1572, signed Investigator CV's, Ethics Committee approval of protocols, protocol amendments and Informed Consent Forms, annual Ethics Committee approval of protocols, etc.). Regulatory documents also have project management related information such as completion date, submission date, approval date, and notes such as who's in charge, activities, and etc. Tracking of documents involves knowing where a document is in terms of the project progression and taking actions at appropriate time. Clinical trial management also involves tracking of the overall status of patient enrollment and other metrics associated with a clinical trial; managing personnel scheduling; as well as tracking communications between the sponsor, CRO and clinical sites.

### Advantages

- Friendly user interface without any clutter or useless features.
- Completely web-based, no need to install any software by the end-user.
- Eliminates the need for meetings to identify tasks and milestones.
- Speeds up information distribution.



## Target CTMS™ Feature List

- Web-based solution
- Login ID/password security (also used for sign-offs)
- Architected for easier system integration
- User self-registration with administrator approval
- Separate administration module
- Permissions-based access
- Database of:
  - Sponsors
  - Sites
  - IRBs
  - Projects
- Project Milestones
- Project deliverables
- Essential documents
- Investigational product tracking
- Dedicated CRA module

### **Administration Module**

- User management
- Password reset
- Sponsor, clinical site and IRB management
- User access control auditing.
- Password expiration configuration.