

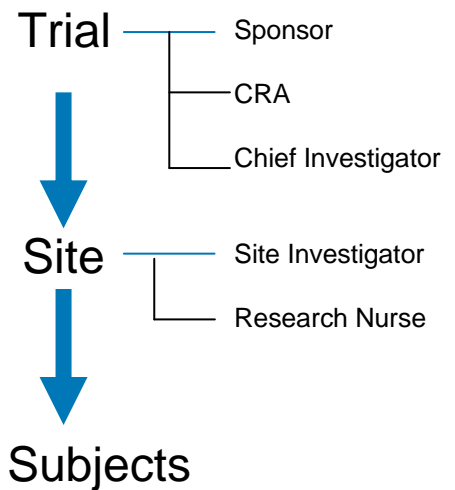


t+ Clinical – Website Users

All website users are allocated a role which controls the level of access they have to various functions and the scope of the data they can access. The roles are arranged in a hierarchy allowing certain users to administer lower levels of users.

Feature	Role			
	Sponsor or CRA	Chief Investigator	Site Investigator	Research Nurse
Secure website	●	●	●	●
View trial statistics & KPI's	●	●		
Manage trial staff		●		
Manage site staff		●	●	
View subject diaries	●	●	●	●
Add subject notes		●	●	●
Message subjects		●	●	●

User Hierarchy



Users can view items at their level and lower level of the hierarchy



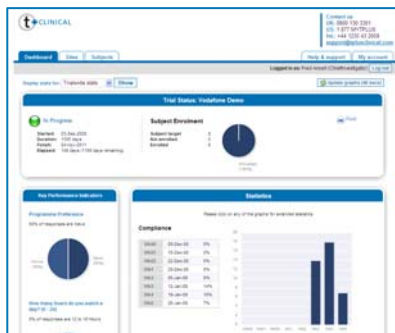
mPRO™



Overview Of Functionality

Secure website

Website access is encrypted by 256 bit SSL, username and password required, and password change every 60 days is enforced.



View trial statistics & KPI's

Compliance, sign up rate and number of diaries submitted can be viewed as bar graphs for the entire trial. Also key performance indicators (KPI's) can be setup at trial initiation and displayed as pie charts. User who can access trial level statistics also have the ability to drill down into details on a per site basis.

Name	Email	Telephone	Address
Research Nurse	ResearchNurse@tplusmedical.com	0800 389 3205	
Principal Investigator	PrincipalInvestigator@tplusmedical.com	0800 389 3205	

Manage trial staff

A chief investigator for the trial can add and manage staff for sites within the trial.

Manage site staff

Administration of staff at a specific site can be managed by a site investigator, allowing them to add more users as required.

Name	Date	Summary
Event 1	2010-01-01	Summary 1
Event 2	2010-01-02	Summary 2

View subject diaries

Diaries submitted by a subject can be easily accessed, and filtered by date range then viewed individually to see all questions and responses.

Add Subject Notes

Textual notes can be added to the log for a subject.

Message Subjects

The system supports instant SMS or secure t+ messaging to a subject.



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