PHARMASEAL

Engility™ CTMS

An innovative clinical trial management platform for unified governance.

Product Fact Sheet

Benefits for Biopharmaceutical organizations

- **Better trial oversight, allowing you more control over your trial**
- **Affordable enterprise-class technology with reduced overall cost of ownership**
- **Simplified access control encouraging partners to work within your system**
- **Rapid deployment and implementation**

Reasons to contact us for a demo

Engility™ CTMS is the ideal technology if you're a small business currently using other methods to support your trials (i.e. paper or spreadsheets) or a medium to large organization looking for a replacement **CTMS**.

Our team of industry experts understand your trial challenges and have developed a new **CTMS** platform, allowing complete visibility over your trial data allowing you to make better decisions to improve your trial management whilst ensuring trial compliance.

For an informal product demonstration and to see the simplicity of **Engility™ CTMS** contact us at

info@pharmaseal.co

Benefits for CRO's

- **Flexible configuration for different sponsor requirements**
- **Ability to manage globally diverse trials**
- **Flexible access control and role based security**
- **Data driven for greater insight into performance and quality**

Engility™ CTMS, the future of clinical trial technology

Engility™ CTMS, the current imperative in clinical trial management, has been developed for global clinical operations teams within Biopharmaceutical, Clinical Research Organizations and Medical Device companies. With these teams in mind the technology offers its users an intuitive and thoughtful experience allowing ease-of-use unlike some existing clinical technologies with difficult navigation and workflow. With one version of the truth, it can easily replace multiple unvalidated spreadsheets used throughout an organization.

Here are just some of the capabilities it can provide global clinical groups:

- Enterprise Cloud System
- Flexible and configurable to meet complex trial requirements
- Industry standard reference data reduces setup time and increases data standardization
- Modern, fresh and intuitive user experience
- Interoperable - built to ft into a clinical data ecosystem
- Focused on structured data capture and workflow for better insights, reporting and analysis

Benefits for Medical Device Companies

- **Flexible and cost-effective for trials of any complexity**
- **Manage investigator-initiated research on medical devices**
- **Oversight and governance to support MDR initiative**
- **Management and insights for on-going real world evidence and observational research**