About DSG

DSG, Inc. is a leading global eClinical provider with a fully unified suite of innovative technology solutions and data management services for the global clinical research community. DSG's eClinical software platform provides competitive advantage that is cost-effective with on-time project delivery. DSG solutions have been used in thousands of clinical trials around the globe with our award-winning eCaseLink™ platform and eCaseLink Designer for enterprise licensing. Founded in 1992, the company is proud to be recognized as the first provider of a fully integrated EDC and IRT Randomization and Trial Supply Management system with the SCDM Data Driven Innovation Award.

DSG, Inc.
325 Technology Drive
Malvern, PA 19355
Office: +1 484.913.0210
Web: dsg-us.com
Email: sales@dsg-us.com

dsg | eClinical Innovation & Experience

eCaseLink™ CTMS

Gain control of your studies with **DSG's**

Clinical Trial Management System

eCaseLink CTMS

Manage clinical trial projects easily with eCaseLink CTMS from DSG.

eCaseLink CTMS, is a powerful web-based application specifically designed for collecting and distributing clinical trial status information rapidly throughout multiple studies. Investigator visit monitoring reports give access to critical status information whenever needed. Sponsors have the ability to make this information available for review by significant stakeholders on the project team. Revolutionary reporting functionality provides seamless access to clinical metrics, workflow events, action items, documents collected, and protocol deviations. Simple configuration, role-based security instills levels of access control including read-only, read-write, and workflow e-signature authority.

eCaseLink CTMS provides

- Comprehensive investigator database
- Visit monitoring forms (*trip reports*)
- Phone contact log
- Action items
- Documents collected log
- Electronic document upload
- Expired documents report
- Expired medical license report
- Site feasibility forms
- Project / Site initiation milestone tracking and reports
- Monitoring activity calendar

Flexible, easy configuration

- Comprehensive access controls
- Customizable role-based security
- Section headings and question wording in Visit Monitoring form configurable per study

Greater control and visibility

Investigator Database

- Investigator status report
- Investigator personnel contact information report

Visit Monitoring Forms (*trip reports*)

- Available form types:
  - Site qualification
  - Site initiation
  - Interim (routine) monitoring
  - Close-out
- Monitoring form headings and questions are configurable
- Both Off-line and On-line trip reports
  - Online implements edits (*question not answered and/ or required comment not provided*)
  - Seamless real-time integration with eCaseLink EDC
- Subject enrollment
- Discontinued subjects
- (SAEs) Serious Adverse Events
- Protocol Deviations
- Generation of a PDF of Visit monitoring forms for printing or electronic file storage

Follow Up Tracking

- Phone Contact Log
- Actions items

Documents:

- Documents Collected Log
- Electronic Document Upload
  - User-definable hierarchy of folders and sub-folders
  - Captures metadata for User, date-time, and site
  - User-defined keywords

eCaseLink™ CTMS advantage

INVESTIGATOR CONTACT INFORMATION

- ISO 27001 : 2013 Certified
- eCaseLink stores investigator and stakeholder contact information
- Available in real-time
- Investigator site information pre-populates in forms reducing data entry

PAYMENTS

- Payment options are established with defined budget milestones
- Budget versus actual reports are available
- Payments are recorded against budget milestones

REPORTING FUNCTIONS

- Real-time standard and customized reports provide seamless and easy access to data
- Provides necessary details that ensure the success of your clinical trial