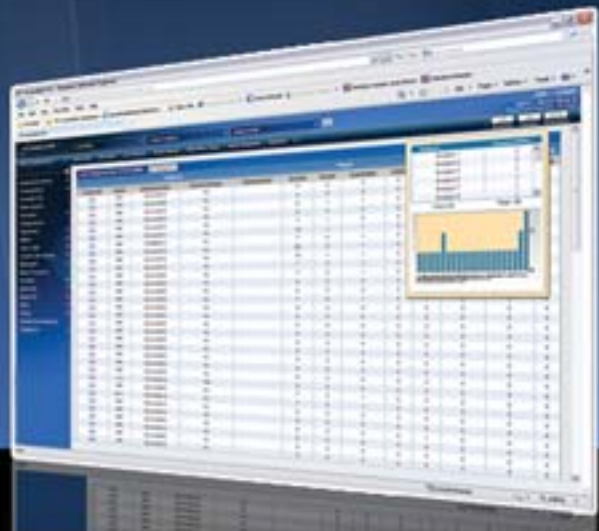




Genuinely Innovative
Truly Integrated



DSG is the oldest and most trusted provider of EDC software solutions in the clinical trial industry. Since our founding in 1992, DSG has strived to develop the most innovative software solutions for clinical trials in the industry.

Our corporation, software and services have grown through our understanding of technology and our collaborative approach with our clients.

DSG has pioneered CRF (Case Report Form) document management software, setting an industry standard. DSG offers a comprehensive product suite including electronic data capture (EDC), electronic patient diaries (ePRO), data management, robust reporting and data analysis tools. DSG has successfully supported over 1,000 clinical trials for more than 325 companies, at over 18,000 sites in over 68 countries. DSG is a global company headquartered in Malvern, Pa., with additional offices in the U.S., Japan and India. DSG has over 250 employees globally and continues to grow.



“ DSG is leading the move forward on clinical trials in the healthcare arena, providing electronic data capture (EDC) for clinical trial data collection and management on a global basis. DSG is accelerating the change from paper data collection and management to the highly efficient electronic data system of eCaseLink™, making the transition seamless, efficient and cost-effective.

~ Inc.500, August 2009

» Superior Technology

For more than a decade, DSG has provided the most innovative software applications such as eCaseLink electronic data capture (EDC), eDiaryLink electronic patient diaries (ePRO), CaseXport Data Archiving, CaseView Imaging, reporting, data integration and document management to the life sciences industry, including the top four pharmaceutical companies.





Your clinical trial goal is the same as ours: to bring drugs and devices to market faster, with measurable ROI. We would like to work with you to achieve your goal, just as we have for some of the world's top pharmaceutical and device companies, for nearly two decades.

We know that you need rapid startup, you need to know what's happening at every stage of your clinical trials, you must effectively manage drug/device supply, you need an efficient way to manage safety and achieve a faster database lock – all resulting in peace of mind, a safer study, faster submission and maximum return on investment.

DSG's innovative eCaseLink 8.0 and industry leading data management services will help you to reach your goal.

eCaseLink 8.0 is a truly integrated eClinical solution that seamlessly combines CTMS, EDC, IWRS and Safety into a single system, cutting costs and giving you control over your entire study. Unlike other systems that attempt to force completely unrelated systems to work together, eCaseLink was developed from the ground up as a truly integrated system whose parts work effectively as a whole. eCaseLink helps you to see everything that's happening with your study in real time, at all times. This unmatched level of control helps you to make mid-study changes, contain costs, improve safety and increase ROI.

DSG's eClinical solutions expedite trials, bring drugs and devices to market faster and provide significant ROI.

We make the transition from paper to EDC *easy*.



At DSG, we understand the challenges of moving from a paper-based system to an electronic data capture process, and we have developed successful and proven best practices through our years of industry experience.

DSG anticipates your challenges and needs so that we can deliver strategic and proactive services and support, as opposed to reactive support once a problem occurs. By planning strategically up front, DSG minimizes problems and challenges once the study has been deployed.

Genuinely Innovative. Truly Integrated.

eCaseLink8.0™

Collect data accurately and more efficiently.

DSG's award-winning eCaseLink software is the most advanced EDC solution in the industry. eCaseLink helps you to achieve your real-world goals by consistently delivering high site acceptance, rapid and efficient workflow, and clean data.

eCaseLink provides rapid start up, rapid database lock, and more rapid FDA submissions, resulting in high ROI.

eClinDirect™ CTMS

Know the status of your study, at all times.

eClinDirect, DSG's web-based Clinical Trial Management System (CTMS), is a project management application that helps you to capture and share a wide variety of clinical trial project status information across multiple studies. eClinDirect gives you unparalleled control over your study.

Advanced reporting functionality provides easy access to metrics, workflow events, action items, documents collected and protocol deviations. Providing a detailed view of your entire study in real time, at all times.

eCaseLink IWRS (Interactive Web Response System)

Gain control of your supply management.

DSG's award winning IWRS is a seamlessly integrated component of eCaseLink that helps you to easily manage randomization, subject enrollment and drug supply management in any clinical trial.

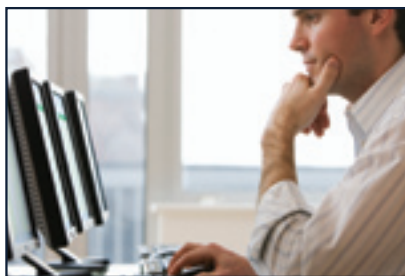
eCaseLink's IWRS functionality completely manages supply chain from shipment to re-supply to ensuring proper kit numbers and the monitoring of defective supply.

eCaseLink's IWRS increases efficiency and productivity with lower error rates, and eliminates the need for costly and inefficient IVR systems. eCaseLink's IWRS delivers savings of over \$100,000 per study, on average.

Best-in-class Data Management Services

Optimize your clinical data management process with DSG's Data Management Division.

Years of experience help us anticipate your challenges and needs so that instead of reacting to data issues, we can design and deliver strategic and proactive data management services and support. We do not wait for problems to occur: we prevent them from happening in the first place. By planning strategically up front, we can minimize problems and challenges further down the line.



The eCaseLink8.0™ advantage

- Control Your Entire Study
- Cuts Trial Costs
- Streamlines Workflow
- Reduces Risk
- Fast Startup
- Rapid Database Lock
- Maximizes ROI
- Award-Winning EDC Software
- Multi-Language Studies
- User-Focused Interface
- Easiest EDC System To Use
- Standard, Custom & Ad-Hoc Reports
- CDISC Standards



eCaseLink's eCRF data entry screens are identical to paper CRFs, providing a look and feel that is very familiar to site users. eCaseLink makes electronic data capture easy & efficient.

eCaseLink 8.0™



➤ eCaseLink Safety System

Streamline Safety

DSG's eCaseLink Safety System streamlines the safety recording and reporting process, eliminates double data entry on SAEs, stores critical documents in a single electronic repository, and generates SAE submission forms.

➤ eCaseLink ScanSource

Accelerate the monitoring process, reduce clinical trial costs

eCaseLink's ScanSource reduces the time spent during monitoring visits, and in some cases may eliminate the need for some interim visits. ScanSource scans study documents at the source and makes them available for review electronically-anywhere, anytime through a secure internet application. This greatly reduces the need for onsite monitoring trips, which saves time and cuts costs.

➤ CaseXport™

Take control with a powerful archive and submission tool

CaseXport is a searchable, standalone archive tool that is easy to use both as a site archive, and for incorporating data into an eSubmission.

CaseXport provides real-time access to clinical data, enabling review of site status and patient data at any time.

CaseXport is trusted by worldwide regulatory authorities in the US, Europe and Japan.

Innovation+Experience

DSG has successfully supported over 1,000 clinical trials for 325 clients, with 1,000,000 patients at over 18,000 sites, in 68 countries, for almost 2 decades.

dsg edc+
Innovation+Experience

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