eCaseLink 8.2™

DSG Drug Safety

Streamline Safety

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

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 thousands clinical trials for hundreds of clients, with millions of patients across 93 countries, for over two decades.

dsgedc+ Innovation+Experience

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Genuinely Innovative. Truly Integrated.

FULL SERVICE & TECH TRANSFER

eCaseLink 8.2™

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

We know that you need rapid startup with easy to use study build tools, 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 8.2 and industry leading data management services will help you to reach your goal.

eCaseLink 8.2 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 clinical study build technology makes initial EDC startup *easy and fast.*

DSG knows the initial investment and challenges involved in building your eCRF system to your exact specifications. With **DSG Designer™** we have developed a fast, easy-to-use and cost effective solution for studies hosted by DSG or through our technology and knowledge transfer option.

DSG Designer's intuitive step-by-step flexible approach to clinical study builds ensures you are one step ahead of the competition, creating studies that are attuned to your business needs and generate measurable ROI. DSG's support will guide you and your team through the build phase as your trusted partner.

Genuinely Innovative. Truly Integrated.

eCaseLink 8.2™

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, fast screen creation 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 eCaseLink 8.2™ 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 proprietary technology validates information field-by-field, including complex cross-visit edits, instantly. These immediate field validations decrease the number of times the user must submit a form and reduce the amount of queries that are generated. The result is clean data and a rapid workflow.