

CaseXport™

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

Eliminate expensive time-consuming re-engineering of trial data into PDFs

Fully integrated with DSG's eCaseLink™ software, CaseXport provides real-time access to clinical data, enabling review of site status and patient data at any time. CaseXport embeds the clinical data in your study within eCRFs together with all of the visual indicators of changes, closed queries, signature status, source document verification status and other data. It then prepares eCRFs for off-line viewing, along with audit trail, queries and reports.

CaseXport is used by worldwide regulatory authorities in the US, Europe and Japan. Additionally, the FDA has audited eCaseLink sites and reported very positive comments to sponsors regarding its ease of use.

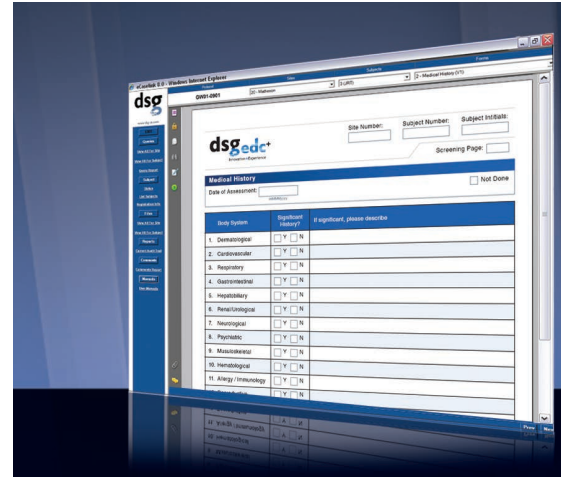
At any point in a study DSG can provide CaseXport CDs or DVDs and individual discs for each site's archive, enabling portable review and analysis of each or all sites.

Easy review of data

CaseXport is designed so that auditors can easily and quickly move from data entry screen to query threads and to the audit trail with a click of a button. Data is searched by subject, keyword or date.

The only fully electronic study archive available

Everything involved in your study — eCRF's, reports, images, audit trail, queries — is captured and stored electronically, saving cost and time. At any point, DSG can create a copy of your study on CaseXport, so you never will lose critical information.



Security

CaseXport's sophisticated encryption option protects clinical data from unauthorized access.

Submission-ready study data available shortly after final database lock.

Once your study is completed, CaseXport can be ready for FDA submission within hours.

No expensive re-engineering of data into PDFs

When you are ready for FDA submission, CaseXport makes it easy to deliver data efficiently and cost-effectively. With DSG's ground-breaking proprietary technologies, eCaseLink files can be saved into a PDF format. The FDA has audited eCaseLink sites and reported very positive comments to sponsors regarding ease of use of the CaseXport site archive product.

About DSG

DSG Inc. supports clinical trial data collection and management with innovative technology solutions including Electronic Data Capture with specialized Clinical Data Management services, Electronic Patient Diaries, Clinical Trial Management Systems and digital on-demand Case Report Form publishing management software. DSG has successfully supported thousands of clinical trials for hundreds of clients, with millions of patients across 93 countries, for over two decades. Founded in 1992, DSG is a global company headquartered in Malvern, Pa., with additional offices in the U.S., Japan and India.

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