

Seamlessly exchange drug safety information dsExchange

Solution at a Glance

dsExchange is the market-leading solution for keeping your adverse event information moving quickly, accurately and efficiently in your clinical research and safety monitoring networks.

Features

- Turn-key system with automated tracking of receipt and processing acknowledgements
- Validation capabilities preview electronic submissions process
- Domestic and international monitoring for regulatory changes

Drug safety issues monopolize headlines around the world. The pharmaceutical industry faces increasing demands to improve processes that enable the sharing of accurate and timely information with trading partners, clinical research organizations (CROs) and other affiliates, particularly when it comes to adverse event reporting.

Couple the large volume of information to track with the challenge of meeting the broad spectrum of worldwide regulatory compliance requirements, and organizations have all of the necessary ingredients for delay, inefficiency and confusion.

Our *dsExchange* solution will help you avoid many of the data and information exchange pitfalls in today's complex drug safety environment.

dsExchange is the market-leading solution for keeping your adverse event information moving quickly, accurately and efficiently in your clinical research and safety monitoring networks.

Next generation

We designed *dsExchange* to provide you with quick, easy and cost-effective access to an extraordinarily powerful communication and workflow application.

Your organization can have this turn-key system, with solutions for both enterprise and ASP models, up and running in a matter of hours, and *dsExchange* has a proven track record of reliability.

A global leader

Cerner, a global leader in healthcare information technology, is solving healthcare's many challenges by connecting the right people with the right information at the right time.



Key Benefits

- Enables quick, efficient and accurate exchange of safety data with trading partners, CROs and regulators
- Supports U.S. Food and Drug Administration and the European Medicines Agency regulatory requirements
- Offers quick implementation and easy-to-use interface

Building on more than 30 years of experience and our partnerships at more than 8,500 sites worldwide, we are finding new and innovative ways to deliver value to our clients.

Let us put our expertise to work for your organization. For more information, please call 703.245.8100, e-mail us at info@drugsafety.com, or visit our website at www.drugsafety.com.