



Clinical • Post-Market • Medical

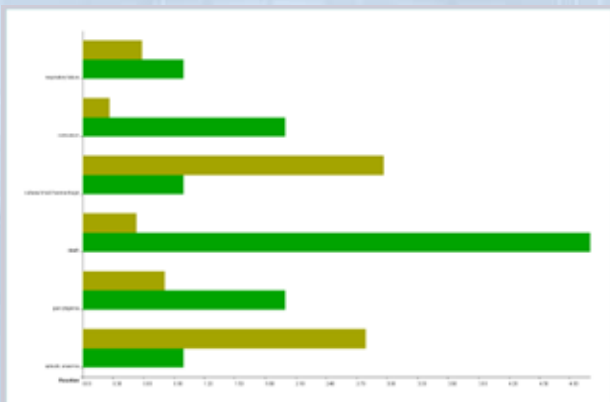
Qscan®-Clinical: The Multi-Database Safety Platform for Risk Evaluation and Mitigation Strategies

What is Qscan?

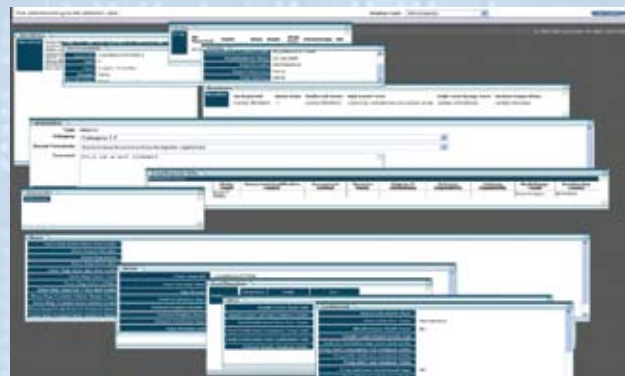
Qscan is an unparalleled safety platform that provides simultaneous analytical views of information from your proprietary safety data, clinical trial data, or medical records data as well as public databases such as FDA/AERS and VAERS and WHO-Vigibase. Drug safety teams benefit from functionality such as auto-alerting, case annotation, signal management, custom reporting and charting, and project management. It does all this in an environment that helps enforce your Standard Operating Procedures and complies with the Food and Drug Administration Amendments Act of 2007.

Why Qscan for your Clinical Trial Safety Teams?

- ✓ Optimize trials for the best safety position
- ✓ See anomalies against important in-class comparators
- ✓ Share results across departments and functional teams
- ✓ Compare clinical trial data to marketed product data



Arm-to-Arm comparison against same background



Clinical trial comparison against various backgrounds

*Qscan puts all
your data in
one place.*

*Let Qscan put
you in control.*

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Qscan-Clinical Approach

Qscan-Clinical was designed to support the people and processes vital to drug safety analysis and risk management strategies. It offers:

- Flexibility to support processes and workflow unique to clinical trial drug safety
- Ability to browse and analyze data from multiples sources simultaneously
- Access to all pertinent analytical techniques including new methods of logistic regression and correlation

How does Qscan-Clinical help your team?

Qscan-Clinical helps your team analyze safety aspects of clinical trials by enabling:

- On-demand characterization of the adverse events as serious or non-serious
- Tailored responses to higher or lower incidences of problems
- Immediate visibility into demographics, label data and other factors
- Constant assessment of time and/or arm differences and differences relative to competitor drugs
- Customized backgrounds selected from a variety of sources such as FDA and competitor label data to detect potential problems early in the process

Let Qscan put you in control:

- ✓ Detect issues as early as possible
- ✓ Integrate your Standard Operating Procedures (SOPs)
- ✓ Document with full audit trail and 21CFR Part11 compliance
- ✓ Satisfy FDAAA requirements

*Drug safety is emerging as a critical operational issue
earlier in the product life cycle.*

Qscan-Clinical...Safety for the Real World.

DrugLogic, Inc. specializes in developing analytical tools and enterprise process support systems for managing risks related to drug safety issues. DrugLogic designs, develops, and delivers products that provide the latest innovations and state-of-the art solutions in support of pharmacovigilance and drug safety surveillance practices for both pharmaceutical and biotech companies. Its proprietary Qscan[®] product monitors both company proprietary adverse event data and publicly available data sources.