Dialog Solutions

Medical Literature Monitoring

Join the growing number of companies - large and small - who rely on us to make sure they don't miss any relevant article when it comes to drug safety.

We offer an innovative, end-to-end suite of biomedical content, software and services designed to help pharma, biopharma, generics, biotech and medical device organizations fulfill their adverse events reporting mandate.

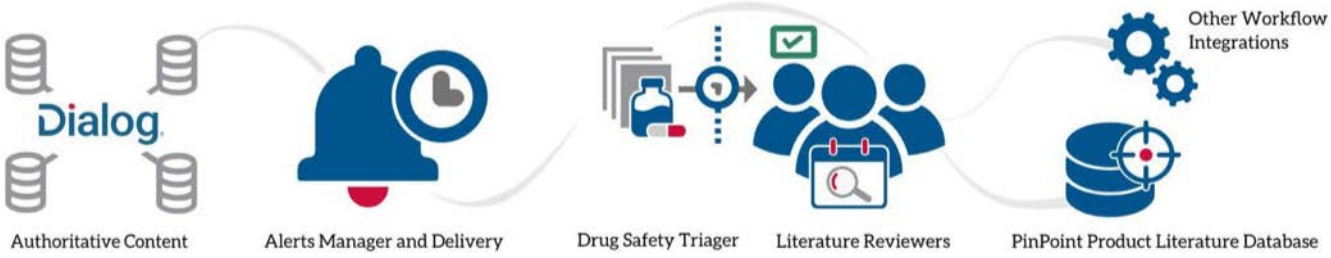
Drug safety monitoring is a continuous and tedious process and yet is subject to intense regulatory scrutiny. Our flagship product, Drug Safety Triager, reduces the laboriousness of this process and ensures you remain compliant and efficient.

Precise, reliable discovery and retrieval of relevant content…

+ Validated, secured and compliant tools to manage the literature workflow

+ Expert medical literature review services

= Improved drug & patient safety enabling you to focus on your core competencies.



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MEDICAL LITERATURE MONITORING: BENEFITS FROM START TO FINISH

DIALOG PRECISION SEARCH

Optimized for pharmacovigilance and proven to minimize the volume of duplicated references delivered into the review process, Dialog's precision search enables you to discover and retrieve the references you need to meet your drug safety monitoring obligations. Dialog's high-quality XML output integrates seamlessly into your drug safety ecosystem where the pool of articles is reviewed and routed appropriately to case processing, periodic safety reports writing and signal evaluations. Our trusted Dialog search specialists can help further improve compliance by constructing audit-ready search strategies and assist in their ongoing maintenance.

ALERTS MANAGER

Plan your work effectively by scheduling your pharmacovigilance alerts to run automatically on a day and time of your choosing with results delivered directly to the Drug Safety Triager. When you need to change search strategies, recipients, frequency or formats the Alerts Manager lets you update your alerts easily, individually and in bulk. An audit history is generated for every change you make to an alert, with the option to add your reasons for each change

DRUG SAFETY TRIAGER

The Drug Safety Triager streamlines the literature review processes for ICSR, Aggregate Reports and Safety Signals submission. This is a fully validated, configurable workflow tool that drives compliance, quality and efficiency. Our system can be scaled to suit organizations of all sizes. Get inspection-ready confidence in your product safety monitoring and achieve significant cost and time savings.

LITERATURE REVIEW SERVICES

Our long-established and highly qualified Literature Review Services Team can support you with critical, time-consuming and resource-intensive tasks within the drug safety workflow. Our team members have extensive experience of performing medical literature screening for adverse events reporting and many of them have gained their expertise while working for large pharma companies. Our quality and timeliness of ICSR review are outstanding. The best practice literature review workflow we developed through the Drug Safety Triager system results in a 70% reduction of the literature "noise" for product leads and safety scientists, enabling them to complete their review for aggregate reports and signals with greater efficiency.

Save time, reduce costs and be audit-ready whilst increasing productivity, compliance and quality.

Learn more about Medical Literature Monitoring at
https://dialog.com/product-service/literature-review-monitoring/
We look forward to connecting with you soon.
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