A modular end-to-end approach to literature monitoring for pharmacovigilance

Our medical literature monitoring solution offers a modular end-to-end approach to literature monitoring for pharmacovigilance.

It gives any organisation tasked with drug safety reporting a streamlined, efficient and compliant approach to the entire literature review process.

Our approach to drug safety reporting allows us to improve every element of this process. Taken together, our content, Technology and support services allow us to provide a complete drug safety monitoring workflow.

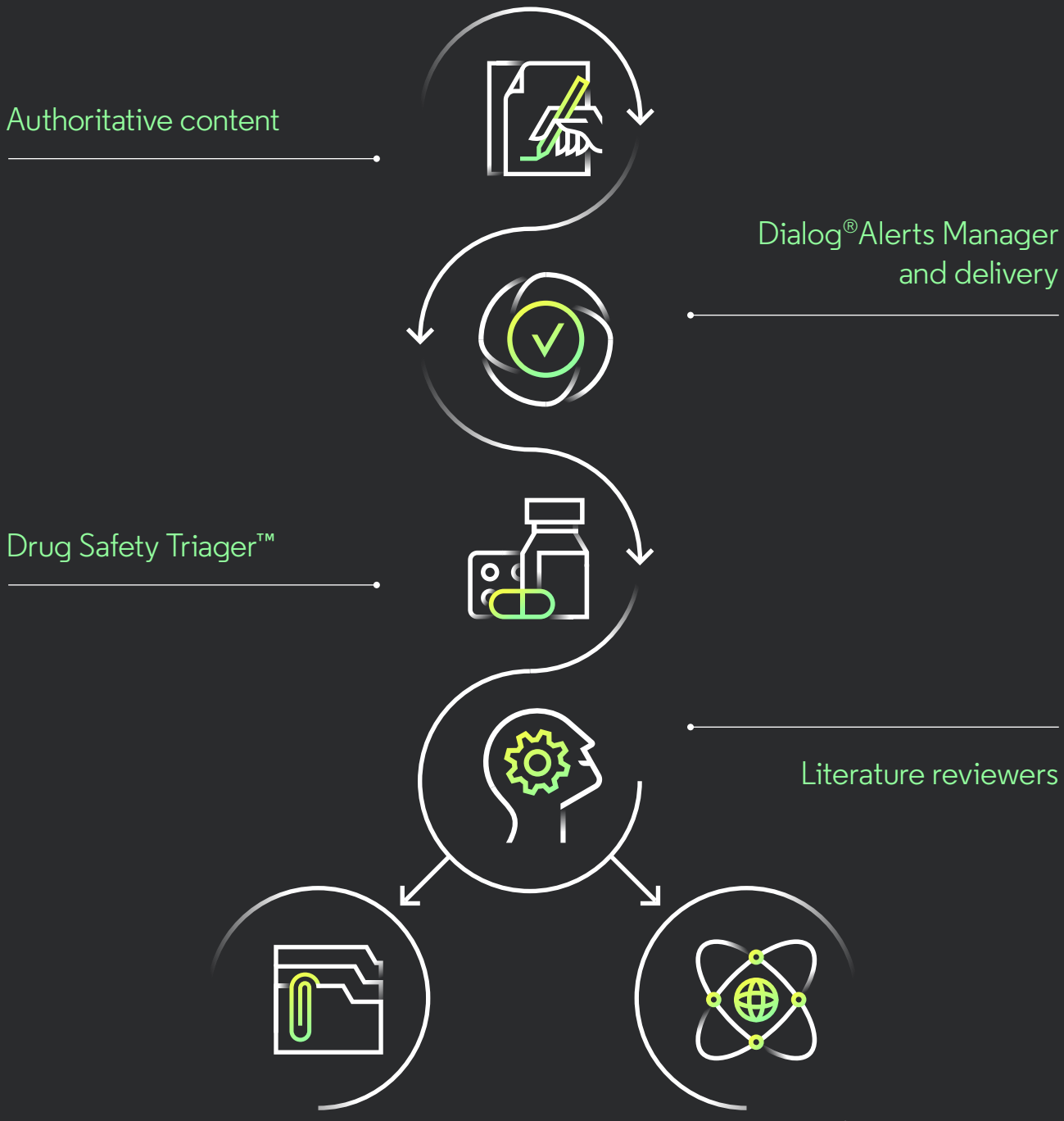
Pharmaceutical companies of all sizes, contract research organisations, medical device manufacturers, biotech and biopharma firms use our medical literature monitoring solution to identify and review articles for relevant ICSRs, aggregate reports (includingPBRER, PSUR and DSUR) and safety signals.

70%

reduction in literature review workflow "noise" delivered by the Drug Safety Triager System.

Clarivate™

The Medical Literature Monitoring process



Our medical literature monitoring approach

Dialog®

Optimised for pharmacovigilance and proven to minimise 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. The 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.

Literature Search Services™

Clarivate has an experienced team of scientific search specialists that can create tailored search strategies for any business. For drug safety monitoring, we can deliver drug safety searches which balance precision of results with recall. As part of the search service, the Literature Search Services team will set up and manage alert search strategies using the Dialog Alerts Manager, ensuring their is an audit trail of all actions taken on each alert.

Dialog 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 process 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 organisations of all sizes. Get inspection-ready confidence in your product safety monitoring and achieve significant cost and time savings. 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.

Literature Review Services

Our long-established and highly qualified literature review services team can support your time consuming and resource intensive tasks within the drug safety workflow. Our team members have extensive experience of performing medical literature scanning for adverse events reporting and many of them have gained their experience while working for large pharmaceutical companies.

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Find out more

Learn about our end-to-end approach to medical literature monitoring at:

clarivate.com/products/dialog-family/

If you'd like to discuss how our medical literature monitoring solution can be deployed at your company, get in touch with your sales rep or contact us at:

dialog.sales@clarivate.com

To learn more about how Clarivate can help you, please visit:

clarivate.com