Adis Pharmacovigilance
The expert solution for regulatory literature monitoring

Life in a pharmacovigilance department is fast-paced and demanding. You are constantly working against the clock to meet your regulatory obligations and with something as important as patient safety at stake, there can be very little margin for error.

In order to meet these challenges, your team needs enough people with the right level of expertise to make the process run smoothly, and to keep up to date with ongoing changes in regulations.

By outsourcing some of the work to us, you can free up your internal resources and have peace of mind that you are securing a reliable and cost-effective service for your organization.
Literature monitoring is a necessary activity for all pharmacovigilance departments. Most likely, you have staff on your payroll dedicating their time exclusively to manually searching and reviewing bibliographic databases. But this process doesn’t have to be such a drain on your internal resources.

When you work with us, we become your partners, an add-on to your existing team that will give you even greater capabilities. Our experts work closely with you to understand your individual situation, current set-up and requirements, and together we will design a solution that makes the most of our combined manpower and expertise.

Structured literature monitoring and assessment are at the very heart of what we do, and we offer a wide range of related services that can be tailored to meet your precise needs.

Our literature monitoring services include:

- Annual Reports
- Periodic Safety Update Reports
- Aggregated Reports
- Local Literature Monitoring
- Special Situations ICSRs
- Customized Source Coverage
- Signal Monitoring
- Ongoing Safety Monitoring
- ICSR Monitoring
Personal support every step of the way

We fully appreciate how precious your time is and we don’t believe that you should have to spend it chasing down the right person to answer your queries. That’s why when you work with us you will have a dedicated contact on hand to help keep things running smoothly, backed up by our global support network.

Our continuous process of support

Feedback

Assess Needs

Devise Solutions

Support Transition

Audit Support

Day-to-day Support

We work with your team throughout the whole process, from designing the right solution to helping you prepare for audits. We understand that your business needs can change during the course of the year, as can regulatory requirements, and so we meet with you, our customers, on a regular basis to ensure that we are always delivering the best possible service to your organization.
The expert solution for regulatory literature monitoring

From the industry-leading provider of pharmacovigilance content

We know that it can be a big step to even consider outsourcing some of this vital pharmacovigilance work, but you can be confident that you are in safe hands with Adis.

With more than 30 years’ experience of delivering pharmacovigilance content and solutions, Adis is the unrivalled industry leader.

More than 6,000 customers globally purchase drug safety information and pharmacovigilance solutions from Adis.

“It would be difficult for us to replicate the expertise and automation that an external vendor such as Adis can offer. The beauty of working with them is that we no longer have to worry about literature monitoring as this part of our pharmacovigilance process is taken care of. We can present our workflows during inspections and audits and be confident that we are complying with regulations”

If you are ready to take the first step and explore the possibility of working in partnership with Adis, please get in touch. Together we will review your current processes and determine how we can best support you.

Email pharmacovigilance@adis.com or visit adis.com/pharmacovigilance for further details.

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- Your partners in pharmacovigilance
- Personal support every step of the way
- Industry-leading provider