

Reactions  
Pharmacovigilance  
Service

Customized literature monitoring service



Wolters Kluwer  
Pharma Solutions

# Streamline your Adverse Drug Reactions (ADR) compliance with confidence

**Reactions Pharmacovigilance Service (RPS) is designed to facilitate your ADR reporting obligations to government organizations like the FDA and EMEA.** Through the most comprehensive, reliable, worldwide literature research available, you gain full confidence that all case reports are identified. So, you save time and resources while reducing the possibility of omissions or errors in monitoring.

**RPS is backed by an unsurpassed editorial staff, highly skilled in medical science and pharmacovigilance, delivering accurate, insightful summaries of all case reports.** You can rest assured that all summaries contain data as specified by the CIOMS II guidelines for periodic safety update reports to deliver the most pertinent information in an easy-to-use format. Our expert editorial staff are available to you to answer questions and help you make the most of our service.

## Most exhaustive literature coverage

- Over 4,000 journals plus their companion supplements\*
- Proceedings from more than 100 major scientific meetings
- Newsletters from more than 80 national centers participating in the WHO International Drug Monitoring Programme
- Regulatory agency and pharmaceutical company websites
- Media releases

## Content customized to your specific preferences and needs

- Choose the drugs or drug classes you want monitored and update them at any time
- Elect to receive first or serious case reports only, or all case reports
- Select the frequency with which to receive content
- Opt for an alert delivered to your desktop or an XML feed delivered to your safety database

## Comprehensive, accurate and timely

- Daily updates available via email alerts
- Full reference information for the original published case report
- Access to a large historic database of more than 35,000 published adverse event case reports dating back to 1992

## New for 2007 – E2B compatible content

Now you can save the time and expense of reworking a report with E2B compatible case reports based on the ICH E2B(M) guidelines. RPS E2B compatible case reports come with the relevant E2B fields already populated, including the latest version of MedDRA coding where required. These case reports are available as a customizable XML feed imported directly into your safety database.

### For more information, please contact:

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\*From January 2007 includes all relevant Medline-indexed journals, including non-English language