Fact Sheet Embase®



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8

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Enhanced pharmacovigilance and post-market drug monitoring with Embase

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- Dedicated search form for building searches using the PV Wizard to facilitate vastly quicker creation of the complex search strings needed for high-recall, high-precision searches focused on adverse events.
- Enable precise retrieval of adverse event information with direct linking of drugs, diseases and devices to associated events mentioned in the article,

Drug and medical device safety research

- Capture the most relevant and up-to-date information from comprehensive journal and conference coverage.
- Avoid missing relevant articles given in-depth, full-text indexing of drug and device trade names, linked to manufacturer names and related procedures.
- Find any mention of your and your competitors' drugs and medical devices.



Systematic reviews of medical literature

- Generate comprehensive, systematic reviews using trusted biomedical information and clinical evidence.
- Refine searches and get results faster using specific search limits, such as Cochrane review, controlled clinical trial, meta analysis, randomized controlled trial and systematic review.

Regulatory compliance

- Track all published and peer-reviewed biomedical information on products to quickly review reported side effects and adverse events information.
- Compare therapeutic effects of two or more substances on a disease, look up reported economic evaluations of specific drug therapies or disease management, track a specific drug trade name in the literature and retrieve information needed to comply with legislation and regulations.
- Use in-depth information to assist in filing of regulatory documentation, such as New Drug Applications (NDA), clinical trial research and program management, including writing protocols and study reports.

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