NOVARTIS

Medical Safety Lead

Job ID
REQ-10086248
авг 25, 2026
Индия

Сводка

Responsible for the drug surveillance program including the necessary follow-up, risk assessment, and relatedness to product on adverse reaction reports, oversight of safety in clinical trials and post marketing programs. Participates in the resolution of any legal liability and complying with governmental regulations. Provides and contributes trending and safety signal detection and risk management assessment for the products' life cycle. Provides safety support to the clinical development teams.

About the Role

Working Model: Hybrid (12 days/month in office)

#LI-Hybrid

Key accountabilities:

- Monitor clinical safety data, including literature, case reports, and signal detection activities
- Drive the application of AI-enabled solutions to simplify and enhance pharmacovigilance processes, leveraging knowledge of AI agents and their use in medical safety.
- Conduct medical assessment of individual adverse event cases and ensure accurate evaluation
- Identify, evaluate, and monitor safety signals using single-case and aggregate data
- Contribute to responses for regulatory authorities and healthcare professional safety inquiries
- Support preparation of core safety documents, including clinical overviews and summary reports
- Provide medical input to aggregate safety reports and regulatory submissions
- Collaborate cross-functionally to integrate safety insights across global development teams
- Guide adverse event coding, causality assessment, and interpretation of clinical safety data

Essential Requirements:

- Degree in Pharmacy, Nursing, Pharmacology, Life Sciences, or a medical degree (medical degree required for roles involving medical review of individual case safety reports)
- Fluency in written and spoken English
- At least 3 years' experience in drug development within a pharmaceutical company (including 2 years in patient safety at an operational or medical position is preferred)
- Experience in clinical trial methodology, regulatory requirements, scientific methodology, and statistical principles, including authorship of scientific publications
- Strong ability to analyse, interpret, and clearly communicate clinical safety data to diverse stakeholders
- Experience in safety and cross-functional issue management (e.g., regulatory inquiries, compliance issues, labelling updates, and risk escalations)
- Proven experience contributing to safety reports and regulatory documentation

Desirable Skills

- Experience of using professional AI tools and agents for process improvement is strongly preferred
- Experience managing clinical safety issues

If you are passionate about patient safety and want to make a meaningful impact at scale, we encourage you to apply and join us in reimagining medicine together!

Why Novartis: Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together? https://www.novartis.com/about/strategy/people-and-culture

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

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Business Unit
Development
Место
Индия
Сайт
Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular

Shift Work
No

Accessibility and accommodation

Novartis is committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to perform the essential functions of a position, please send an e-mail to diversityandincl.india@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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