NOVARTIS

(Senior) Medical Safety Physician

Job ID
REQ-10088068
Сен. 16, 2026
Китай

Сводка

Act as the deputy of the Local Qualified Person for Pharmacovigilance in Novartis Country Organization to fulfill medical safety oversight and provide safety deliverables for assigned therapeutic areas/ products in China in full drug life-cycles.

About the Role

Major accountabilities:

- Act as Qualified Delegate of the Country Patient Safety Head, functionally (in terms of responsibility for PV system) for assigned therapeutic areas/ products.
- Ensure robust oversight and compliance in terms of reporting/submission/distribution of safety reports/updates/information (e.g., SAE, SR, IN, SUSAR, PSUR, DSUR, changes in risk-benefit profile) to Local Health Authorities according to regulatory requirements and Novartis procedures.
- Provide medical safety support, including safety deliverables in CTA/ (s)NDA/ License Renewal/ Reimbursement dossier is done according to the timeliness described into the respective procedures or as committed with line functions. Play a joint role with global safety leads for safety relevant issues or requests.
- Represent PS in CTT for Post Approval Commitment (PAC) studies and China bridging clinical trials, with the support by global safety lead if needed.
- Work in close collaboration with other local and global medical safety functions to ensure accurate evaluation of safety data.
- Lead local RMP and RMP China addendum creation and approval, based on local regulatory or LHA requirement, if applicable.
- Conduct local safety signal detection and escalate to global medical safety for potential safety signals identified from all local post-marketing sources per local regulatory requirements.
- Provide scientific expertise during review of all Phase IV Clinical Trial and NIS protocols safety sections including Research Collaborations and if a Contract Research Organization (CRO) is conducting the trial or study, review safety relevant sections of the contract.
- Responsible for responses to inquiries from LHA on safety issues related, involve in the communication on safety topics related to responsible products with the LHA.
- Ensure support for and close-out of audits, corrective action plan, investigation, and Health Authority inspections.

Essential Requirements:

- Medical background
- At least 3 years experience in pharmacovigilance or equivalent field, or at least 2 year safety physician experience.
- Project management skills

Desirable Requirements:

- Excellent communications and negotiation (networking) skills
- Quality and results oriented
- Business mindset

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.
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Дивизион
Development
Business Unit
Development
Место
Китай
Сайт
Beijing (Beijing)
Company / Legal Entity
CN06 (FCRS = CN006) Beijing Novartis Pharma Co., Ltd
Alternative Location 1
Shanghai (Shanghai), Китай
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No

Accessibility and accommodation

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