

# Medical Safety Physician

Job ID  
REQ-10072141  
апр 14, 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

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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

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### **Accessibility and accommodation**

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