

# **Medical Advisor, Ophtha**

Job ID REQ-10025594 Nov 29, 2024 China

# **Summary**

#### About this role:

In line with overall product strategy, the Medical Advisor is responsible for supporting the design, implementation and execution of Medical Affairs plans for assigned Therapy Area, providing scientific information, helping design and organize clinical studies, building educational dialogue with KOLs and regulatory stakeholders

#### About the Role

Your key responsibilities:

Your responsibilities include, but are not limited to:

- Support country medical affairs strategy in line with the global strategy, country insights and market conditions, & secure implementation of planned Medical Affairs activities within the designated therapy area(s).
- Coordinate scientific meeting, symposia, congresses, Continuous Medical Education (CME) and other medical / scientific exchange and engagement activities which could bring additional value to the relevant therapy area; develop strategic engagement plan(s) for country customer-facing medical activities and events, and ensure timely execution of planned medical affairs activities in an efficient and compliant way.
- Ensure medical enquiries are responded to in a high quality, timely manner, and in accordance with applicable standards; establish standard response documents as appropriate, for frequently asked questions.
- Provide medical/scientific input into the development and execution of clinical trial or clinical research related activities, including initiation and oversight of clinical studies / clinical research within the respective therapeutic area. Support country strategy for Non Interventional Studies/Investigator Initiated Trial activities.
- Coordinate review and approval of medical materials and locally developed promotional materials; ensure medical materials provided from global or region for stakeholder engagement and events are tailored to local needs, and reviewed/approved per local/P3 guidelines.
- Ensure medical insights are provided to cross functional groups, including, but not restricted to: Pharmacovigilance, Regulatory affairs, Market Access, QA, Commercial teams, Brand team and others.
- Responsible for risk identification and assessment, mitigation planning as well as implementation and monitoring of appropriate internal controls within the area of responsibilities.

What you'll bring to the role:

- •Education & Qualifications: Masters' degree or above, medical background
- Languages: English& Chinese
- Collaborating across boundaries
- Medical advisor and new launch experience preferred
- Operations Management and Execution

#### Project Management

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Novartis is committed to building an outstanding, inclusive work environment and diverse team's representative of the patients and communities we serve.

### 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.china@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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Division

International

**Business Unit** 

Innovative Medicines

Location

China

Site

Shanghai (Shanghai)

Company / Legal Entity

CN06 (FCRS = CN006) Beijing Novartis Pharma Co., Ltd

Functional Area

2/3

Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work

Nο

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