# Medical Affairs Director

Job ID
REQ-10036028

4月 10, 2025

United Kingdom

## 摘要

The Global Medical Affairs team acts as enterprise medical voice across the asset lifecycle and leads the medical strategy for the therapeutic area.

The Medical Director role is responsible for the implementation of medical strategies for early programs globally with focus on innovative evidence solutions including interventional studies, non-interventional studies (NIS) and real-world evidence (RWE) studies and implementation science projects.

With extensive experience in drug development, they will be able to lead Integrated Evidence Packages (IEP) in situations with higher scientific complexity and potential regulatory challenges. They will also act as a subject matter expert in the development of the overarching strategies, providing inputs during design and along the end-to-end execution of programs across different disease areas.

## About the Role

As the Medical Director, you will provide proactive input to development on potential new therapeutic indications, to enrich Registration Programs and to consider new therapeutic opportunities. You will also ensure that Patient Access programs are supported for all brands within Global Medical Affairs (GMA) and that GMA activities are designed and executed in compliance with company policy guidelines and highest medical quality standards.

**MAJOR ACCOUNTABILITIES:**

- Lead development and execution of medical affairs strategy for TA/Asset priority programs including transformative tactics such as: research/population health, innovative partnerships and integrated evidence plans
- Co-develop plans for evidence generation, Medical Science Liaison (MSL) / Field Medical Affairs strategy, medical education programs, scientific publication planning and Medical Expert network development with therapeutic areas (TAs)
- Co-own the development and implementation of innovative education and scientific communication plans for external stakeholders
- Prepare Scientific Research Committee (SRC) submissions for TA assets within remit
- Partner with Development, Sales & Growth (S&G), US and International cross-functions to shape portfolio early and diversify evidence to achieve broad access at launch and to enhance impact on clinical practice for priority programs
- Represent GMA around prioritized portfolio with internal and external audiences, in collaboration with TAs including the investment, medical and regulatory communities, as well as pharmaceutical or biotechnology industry collaborators/partners
- Provide direction and input into the development and implementation of successful reimbursement and market-access strategies

**REQUIREMENTS:**

**Essential:**

- MD (Preferred) or PhD/PharmD in Health Sciences. Specialist Degree or specialist qualification related to discipline for which you will be responsible is an advantage.
- 5+ years in Pharmaceutical Industry experience in Medical Affairs and/or Clinical Development
- Fluent oral and written English; Other relevant languages are an advantage.
- Strategic mindset and able to establish credibility and influence across a range of diverse stakeholders in a matrix organization to drive change
- Ability to truly collaborate across functions and markets: serve-partner-co-create
- Deep understanding of health care systems and key external stakeholders
- Strong track record of delivery focus for time and quality in medical affairs projects
- Firm working knowledge of Good Clinical Practice (GCP), scientific and clinical methodology, protocol designs, management and regulatory requirements for clinical studies designated for review by regulatory authorities.

**Preferred**

- Highly preferred: TA expertise, significant medical affairs early asset lifecycle, pre-launch and launch experience in Global organizations
- Experience in developing and executing "Best in Class" processes at scale
- Clinical trial research experience conducted in a pharmaceutical or equivalent academic environment in TA of interest is strongly desired.

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部门
Development

Business Unit
Pharma Research

地点
United Kingdom

站点
London (The Westworks)

Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.

Alternative Location 1
Barcelona Gran V í a, Spain

Alternative Location 2
Hyderabad (Office), India

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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