

## Global Medical Affairs Director Remibrutinib

Job ID  
REQ-10061831

9月 17, 2025

Spain

### 摘要

LOCATION: London, UK or Barcelona, Spain

ROLE TYPE: Hybrid Working, #LI-Hybrid

The Global Medical Affairs Director leads the medical strategy & tactics for a Remibrutinib program/indication, ensuring the US and International medical perspective is reflected.

They will develop, own and drive the execution of the Integrated Evidence Plan (IEP), ensuring the right evidence is available at the right time to enable access and clinical adoption of our assets.

They will also act as a strategic partner to Biomedical Research (BR), Development, International Medical Affairs, US Medical Affairs, Strategy & Growth and Commercial.

About the Role

## Major accountabilities:

- Development and execution of high quality medical strategy for the asset/ indication and vision throughout its lifecycle at global level.
- Creation and execution of the integrated evidence plan addressing US and International top market needs and supporting clinical adoption of the asset, incl. various evidence generation vehicles, e.g. interventional trials, RWE, data mining, IITs etc.
- Design, set-up and execution of interventional clinical trials
- Serves as disease area medical expert for internal stakeholders from different line functions as well as external customers.
- Partners with Development, Strategy and Growth (S&G), US and International cross-functions to diversify evidence to achieve broad access at launch and to enhance impact on clinical practice for the asset.
- Financial tracking to ensure timely development & execution of medical activities.
- Prepare Specification Review Committee (SRC) submissions for Therapeutic Area (TA) assets within remit.
- Represent GMA around prioritized portfolio with internal and external audiences, including the investment, medical and regulatory communities, as well as pharmaceutical or biotechnology industry collaborators/partners.
- Provide proactive input to asset lifecycle management on potential new therapeutic indications to consider.
- Ensure that Patient Access programs are supported and delivered with full compliance.
- Ensures GMA activities are designed and executed in compliance with company policy guidelines and highest medical quality standards.

## Education:

- MD or equivalent (preferred), PhD, or PharmD degree required
- Specialist Degree or specialist qualification related to discipline for which you will be responsible is an advantage

## Key Skills & Experiences:

- 5+ years in Pharmaceutical Industry experience in Medical Affairs and/or Clinical Development
- Experience working cross-functionally
- Immunology experience & expertise
- Successful development and execution of innovative medical strategies, a broad of range of medical tactics and Integrated Evidence Plans
- Track record of successful high-quality evidence generation projects
- Firm working knowledge in Good Clinical Practice (GCP), evidence generation activities, such as interventional or non-interventional studies and Real World Evidence (RWE) projects, including scientific communication of their results
- Sound experience in a broad range of typical medial tactics, e.g. advisory boards, steering

committees, sponsorships, congresses, symposia, publications or various forms of external (academia) partnerships

- Deep understanding of health care systems and key external stakeholders such as Health Care Professionals (HCPs), payers, medical societies and guideline committees
- Understands unmet medical needs, generates the right evidence to effectively address them, uses innovative, multichannel communication formats for effective evidence dissemination
- Strategic mindset and able to establish credibility and influence across a range of diverse stakeholders in a matrix organization to drive change

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部门  
Development

Business Unit  
Innovative Medicines

地点  
Spain

站点  
Barcelona Gran V í a

Company / Legal Entity  
ES06 (FCRS = ES006) Novartis Farmac é utica, S.A.

Alternative Location 1  
London (The Westworks), United Kingdom

Functional Area  
Research & Development

Job Type  
Full time

Employment Type  
Regular

Shift Work  
No

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representative of the patients and communities we serve.



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