NOVARTIS

Global Medical Affairs Director, Cardiovascular

Job ID
REQ-10058469

9月 23, 2025

United Kingdom

摘要

#LI-Hybrid
Location: London, United Kingdom & Barcelona, Spain

Are you ready to shape the global medical voice across the asset lifecycle? Our Global Medical Affairs team is at the forefront of enterprise-wide medical leadership, driving strategy across therapeutic areas and ensuring a unified international and U.S. medical perspective. We lead the development and ownership of Integrated Evidence Packages (IEPs), guiding innovative evidence generation through interventional studies, non-interventional studies (NIS), real-world evidence (RWE), and implementation science projects.

This is a unique opportunity to bring your deep expertise in drug development and scientific strategy to some of the most complex and high-impact programs in the industry. You ’ ll act as a subject matter expert, influencing design and execution across disease areas, and providing strategic leadership in collaboration with NIBR, Development, and Medical Affairs teams globally. If you're passionate about advancing medical innovation and thrive in high-stakes, cross-functional environments–this role is for you.

About the Role

Key responsibilities:

- **Lead** development and execution of medical affairs strategy for Asset priority programs, including transformative tactics such as research/population health, innovative partnerships, and integrated evidence plans.
- **Co-develop** plans for evidence generation, MSL / Field Medical Affairs strategy, medical education programs, scientific publication planning, and Medical Expert network development.
- **Co-own** the development and implementation of innovative education and scientific communication plans for external stakeholders.
- **Track** financials to ensure timely and cost-effective development and execution of medical activities.
- **Prepare** SRC submissions for TA assets within remit.
- **Partner** with Development, US and International cross-functional teams to shape the portfolio early and diversify evidence to achieve broad access at launch and enhance impact on clinical practice for priority programs.
- **Represent** Global Medical Affairs 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.
- **Advocate** for "the voice of the patient" internally and evaluate factors relevant to a patient's informed decision-making.
- **Provide** direction and input into the development and implementation of successful reimbursement and market-access strategies.
- **Contribute** proactive input to Development on potential new therapeutic indications to enrich Registration Programs and explore new therapeutic opportunities.

Essential Requirements:

- Fluent oral and written English.
- MD or PhD/PharmD in Health Sciences. Specialist Degree or specialist qualification related to discipline for which you will be responsible is an advantage.
- Experience working in the cardiovascular therapy area is preferred
- Above country medical affairs experience either in a global or regional multi-country role.
- Experience in Pharmaceutical Industry experience in Medical Affairs and/or Clinical Development.
- Strategic mindset and able to establish credibility and influence across a range of diverse stakeholders in a matrix organization to drive change.
- Deep understanding of health care systems and key external stakeholders.
- Strong track record of delivery focus for time and quality in medical affairs projects.
- Understands unmet medical needs, generates the right evidence to fulfil them, uses innovative, multichannel communication formats for effective evidence dissemination.
- Firm working knowledge of GCP, scientific and clinical methodology, protocol designs, management and regulatory requirements for clinical studies designated for review by regulatory authorities.

Commitment to Diversity and Inclusion / EEO

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

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

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: https://talentnetwork.novartis.com/network

Benefits and Rewards: Read our handbook to learn about all the ways we ' ll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

部门
Development

Business Unit
Universal Hierarchy Node

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

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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