

Global Regulatory Affairs Associate Director (Cardio-Metabolic)

Job ID

REQ-10064681

May 28, 2026

LOC_GB

About the Role

Major accountabilities:

- Lead the implementation of regulatory strategies and operational activities across major global regions.
- Provide strategic input into global regulatory plans, identifying risks and contributing to key planning documents.
- Align regional regulatory approaches with global objectives through collaboration with cross-functional and regional teams.
- Define and manage Health Authority (HA) interaction strategies, including preparation of briefing materials.
- Oversee the planning, coordination, and submission of regulatory dossiers (e.g., CTAs, INDs, Risk Management Plans).
- Serve as a liaison with local HAs (e.g., FDA, EMA) and lead or support negotiations for regional approvals.
- Develop and implement strategies to minimize review delays and regulatory clock stops.
- Ensure timely and compliant responses to HA queries and requests.
- Contribute to departmental goal setting and lead initiatives to improve regulatory processes.
- Ensure adherence to internal policies, SOPs, and global regulatory requirements.

Minimum requirements:

- Bachelor's or Master's degree in Life Sciences, Pharmacy, or a related field.
- Significant experience in regulatory affairs within the pharmaceutical industry.
- Proven track record in project management and regulatory operations.
- Experience representing the organization in cross-functional and cross-cultural settings.
- Strong knowledge of clinical trials, drug development, and regulatory compliance.
- Excellent problem-solving, negotiation, and communication skills.
- Detail-oriented with the ability to manage complex regulatory projects.
- Skilled in risk management and working with cross-functional teams.
- Ability to navigate and influence Health Authority interactions.
- Fluency in English (written and spoken) is essential.

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.

Role Requirements

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>

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.

[Read our handbook \(PDF 30 MB\)](#)

Division
DIV_GD
Business Unit
Development
Location
LOC_GB
Site
London (The Westworks)
Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.
Alternative Location 1
LOC_GB
Functional Area
FCT_RD
Job Type
Full time
Employment Type
Regular
Shift Work
No
[Apply to Job](#)

Job ID
REQ-10064681

Global Regulatory Affairs Associate Director (Cardio-Metabolic)

[Apply to Job](#)

Source URL: <https://jobapi.novartis.com/req-10064681-global-regulatory-affairs-associate-director-cardio-metabolic>

List of links present in page

1. <https://jobapi.novartis.com/req-10064681-global-regulatory-affairs-associate-director-cardio-metabolic>
2. <https://www.novartis.com/about/strategy/people-and-culture>
3. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/London-The-Westworks/Global-Regulatory-Affairs-Associate-Director--Cardio-Metabolic-_REQ-10064681-3
5. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/London-The-Westworks/Global-Regulatory-Affairs-Associate-Director--Cardio-Metabolic-_REQ-10064681-3