NOVARTIS

Associate Director, GMA Study Management - Neuroscience

Job ID
REQ-10076843

4月 30, 2026

United Kingdom

摘要

Ready to shape how medical evidence influences patient care on a global scale? As Associate Director, Global Medical Affairs Study Management, you will lead the end to end delivery of high impact medical studies across a Neuroscience disease area portfolio. This is a strategic, hands on role where scientific insight meets operational leadership - driving non interventional studies, research collaborations and investigator initiated trials that matter. Working in a highly collaborative, matrix environment, you ’ ll partner closely with global stakeholders to ensure studies are delivered with rigour, quality and purpose, ultimately supporting better decisions and better outcomes for patients worldwide.

Location: London, UK #LI-Hybrid

Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

About the Role

Responsibilities:

- Lead end - to - end planning, execution, and reporting of Global Medical Affairs studies across the Neuroscience disease area.
- Ensure studies are delivered on time, within budget, with high quality and full regulatory compliance.
- Drive delivery of non - interventional studies, research collaborations, and investigator - initiated trials.
- Partner with the Study Management Director on resource planning, prioritisation, and capability deployment.
- Lead matrix teams, including internal associates and external service providers, to ensure capacity and performance alignment.
- Identify operational risks early, implement mitigation strategies, and provide clear progress updates to senior stakeholders.
- Represent Global Medical Affairs Study Management in programme governance forums and cross - functional decision - making bodies.
- Oversee contract research organisation selection, contracting, and performance in partnership with vendor management.
- Build and maintain strategic partnerships with institutions, key opinion leaders, and external collaborators.
- Champion a culture of quality, compliance, process simplification, and operational excellence across study delivery.

Essential for the role:

- Master ' s degree in a scientific discipline; doctorate or Doctor of Pharmacy qualification preferred.
- At least eight years ' experience planning, executing, and reporting complex clinical or medical studies in pharmaceutical or research settings, with proven end-to-end delivery accountability.
- Strong experience leading and delivering complex, international programmes within a matrix, cross - functional environment.
- Proven knowledge of clinical development, Good Clinical Practice principles, global medical affairs processes, and experience within Neuroscience or closely related therapeutic areas.
- Demonstrated expertise in project leadership, budget management and oversight, resource planning, and operational risk management.
- Ability to build effective partnerships with internal stakeholders, external service providers, and scientific collaborators to deliver high - quality study outcomes.
- Experience driving quality, compliance, and inspection readiness across regulated study environments.
- Excellent communication, problem - solving, and leadership skills, with confidence influencing senior stakeholders.

Desirable for the role:

- Experience in Medical Affairs and non-interventional study design
- Prior involvement in Health Authority inspections or audit readiness activities

Commitment to Diversity & Inclusion:

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

Benefits and Rewards: Learn about all the ways we ' ll help you thrive personally and professionally. Read our handbook (PDF 30 MB)

部门
Development

Business Unit
Development

地点
United Kingdom

站点
London (The Westworks)

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

Functional Area
Research & Development

Job Type

Full time

Employment Type
Regular

Shift Work
No

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