NOVARTIS

Senior Scientific Writer II

Job ID
REQ-10077966
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Великобритания

Сводка

#LI-Hybrid

Location: London, UK.
Assigned therapeutic area: CRM

This role is based in London, UK. Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

The Senior Scientific Writer II develops high quality, accurate, and compliant medical and scientific communications aligned with therapeutic area strategy and the brand's overarching scientific narrative. The role leads the planning, and delivery of a broad range of scientific materials, including medical education slide decks, medical congress including symposia, advisory board materials, and scientific content supporting congress activities and internal medical engagements.

Operating within an International, matrix environment, the Senior Scientific Writer II partners closely with other Scientific Writers and collaborates cross-functionally with colleagues across IMA (IMACE, TAs), Global Medical Affairs (GMA), and additional clinical, and commercial stakeholders. Through these partnerships, the role drives content excellence, governance, and harmonization across therapeutic areas and markets, contributing to a cohesive and impactful scientific communication strategy.

The position reports into the Scientific Writing Lead, CRM.

About the Role

Key Responsibilities:

- Develop a broad range of scientific and medical materials, including slide decks, congress/symposia content, advisory board materials, and internal medical engagement assets.
- Prepare congress-related materials such as satellite symposia agendas, speaker briefing documents, and slide content.
- Research, interpret, and synthesize complex scientific and clinical data into accurate, well-referenced, evidence-based content aligned with TA strategies.
- Ensure scientific precision, clarity, and IMACE-level quality standards across all materials, supporting review processes with strong input on messaging, data accuracy, and consistency.
- Manage multiple concurrent projects, potentially across more than one brand, while maintaining high quality and timely delivery.

Matrix Collaboration & Stakeholder Engagement

- Collaborate with functional and cross-functional partners (IMA, GMA, medical, clinical, etc.) to align on scientific priorities and clarify content requirements. Participate in routine discussions to refine key messages and ensure content is accurate, consistent, and fit for purpose.
- Contribute to enhancements in content formats, delivery approaches, and tools to improve experience and effectiveness across channels.

Quality, Standards & Governance

- Ensure all materials comply with internal policies, external regulations, structured review processes, and governance frameworks.
- Apply established templates, writing standards, QC processes, and documentation requirements to maintain scientific rigor, quality, and audit-ready outputs.
- Maintain robust version control, documentation trails, and content integrity across the lifecycle of scientific materials.

Essential Requirements

- Education minimum: BSc or equivalent, but preferred: Advanced degree (PhD/Postdoc/MD).
- 2-3 years experience in a scientific writing from the industry (pharma or consulting for pharma)
- Strong ability to interpret, synthesize, and communicate complex scientific and clinical data with accuracy and scientific rigor.
- Experience collaborating in matrixed, cross-functional environments. Proven ability to deliver high-quality scientific content under tight timelines while managing multiple parallel projects.
- Familiarity with medical review and approval processes, documentation management, version control, and compliance standards.
- Proficiency with digital content platforms and structured/modular content approaches, with strong grounding in scientific governance, QC processes, and templates.
- Fluent oral and written English; additional languages desirable.

Desirable Requirements:

- Previous experience in Cardiovascular, metabolic or renal medicine.

Commitment to Diversity and 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)

Дивизион
International
Business Unit
Marketing
Место
Великобритания
Сайт
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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