NOVARTIS

Expert Scientific Writer

Job ID
REQ-10074550
апр 08, 2026
Ирландия

Сводка

#LI-Hybrid Hybrid Dublin

As an Expert Scientific Writer you will be responsible for the creation of high-quality complex scientific content, such as publications and foundational core content elements, in line with priorities and scientific narrative defined in SCP. You will also manage the ownership of content from brief to publication or presentation, for first-time right delivery.

About the Role

Major Activities

- Demonstrate a command of assigned therapeutic areas and expertise with assigned products.
- Research and write original content for publications activities (primary and review manuscripts, abstracts, posters), slide presentations and other materials.
- Prepare meeting materials for satellite symposia (agenda, slide content, speaker briefings etc), and reports from advisory boards and other internal or external meetings.
- Develop content that is scientifically accurate, evidence-based, grammatically accurate, referenced using appropriate sources, and consistent with quality standards for author review, customer review as appropriate, and scientific peer review.
- Perform internal scientific reviews to ensure quality in line with the scope and scientific messages.
- Mentor internal team members and help onboard new joiners.
- As needed, perform quality control (QC) checking / proof reading of the above-mentioned documents to meet stakeholder expectations.
- Clearly communicate medical scientific concepts in a condensed, audience-appropriate way.
- Follow all internal processes and procedures with regard to workflow, development of deliverables, and adherence to industry best practices, including GPP.
- Demonstrate the flexibility/adaptability necessary to function on different therapeutic teams as needed and to work on projects across multiple brands at any given time.
- Provide input and aid in troubleshooting/problem-solving.
- Participate in strategic and tactical publications planning and related research.
- Maintains records for all assigned projects including archiving in line with global SOPs
- Maintains audit, SOP and training compliance.
- Performs additional tasks as assigned.

Key Performance Indicators

- Adheres to quality, compliance to SOPs, timeliness, and productivity of deliverables as per KPI targets.
- Adheres to Novartis values and behaviors.

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
Development
Business Unit
Development
Место
Ирландия
Сайт
Dublin (NOCC)
Company / Legal Entity
IE02 (FCRS = IE002) Novartis Ireland Ltd
Functional Area
Research & Development

Job Type
Full time
Employment Type
Regular
Shift Work
No

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

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