

# Expert Scientific Writer

Job ID

REQ-10071592

Apr 09, 2026

LOC\_IE

## About the Role

Key Responsibilities:

### Scientific Content Development

- Develop high-quality, scientifically rigorous materials - including slide decks, congress/symposia content, and advisory board resources - aligned to the overarching scientific narrative. Ensure scientific precision, clarity, and IMACE-level quality standards across all materials, supporting review processes with strong input on messaging, data accuracy, and consistency.
- Own end-to-end delivery of scientific content, ensuring first-time-right execution across multiple concurrent projects and brands.
- Research, interpret, and synthesize complex scientific and clinical data into clear, accurate, strategically aligned content for diverse audiences.
- Provide scientific insight to address data-related queries, support problem solving, and ensure precise messaging.

### Matrix Collaboration & Stakeholder Engagement

- Partner closely with IMA, GMA, and cross-functional clinical, and commercial teams to understand priorities and refine scientific messages.
- Integrate feedback efficiently and contribute to publications planning, research activities, and team capability building through mentoring and onboarding.
- Support continuous improvement of content formats, tools, and delivery approaches across channels and therapeutic areas.

### Content Excellence, Governance & Harmonization

- Uphold scientific quality, accuracy, consistency, and adherence to templates, SOPs, and governance processes, ensuring all content is compliant and audit-ready.
- Promote harmonization of scientific communications across therapeutic areas and markets to support an integrated and impactful communications strategy.
- Apply structured content and digital platforms to maintain, update, and optimize scientific materials, enabling lifecycle management, reuse, and improved efficiency.

Essential Requirements:

- Education and Experience: Minimum BSc with 10+ years relevant experience, Preferred Advanced degree (PhD/Postdoc/MD) with 5+ years of relevant experience.
- Extensive experience developing scientific or medical content within pharmaceutical, biotech, or healthcare communications environments.
- Over 2 years of experience in at least one therapeutic area: Oncology, Cardiovascular, Renal, Neuroscience, or Immunology.
- Proven ability to interpret, synthesize, and communicate complex scientific and clinical data with high scientific rigor.
- Strong experience working in global, matrixed, cross-functional teams (Medical Affairs, Clinical Development, etc.). Demonstrated ability to manage multiple concurrent scientific projects under tight timelines across brands or therapeutic areas.
- Solid understanding of medical review processes, documentation management, compliance standards, and version

control.

- Proficiency with digital and modular content platforms and structured content approaches.
- Fluent oral and written English; additional languages desirable.

Benefits & 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>

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>

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>

## 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\_IM

Business Unit

Marketing

Location

LOC\_IE

Site

Dublin (NOCC)

Company / Legal Entity

IE02 (FCRS = IE002) Novartis Ireland Ltd

Functional Area

FCT\_RD

Job Type

Full time

Employment Type

Regular

Shift Work

No

[Apply to Job](#)

Job ID

REQ-10071592

## Expert Scientific Writer

[Apply to Job](#)

---

**Source URL:** <https://prod1.jobapi.novartis.com.cn/req-10071592-expert-scientific-writer>

**List of links present in page**

1. <https://prod1.jobapi.novartis.com.cn/req-10071592-expert-scientific-writer>
2. <https://www.novartis.com/careers/benefits-rewards>
3. <https://www.novartis.com/about/strategy/people-and-culture>
4. <https://talentnetwork.novartis.com/network>
5. <https://www.novartis.com/about/strategy/people-and-culture>
6. [https://www.novartis.com/sites/novartis\\_com/files/novartis-life-handbook.pdf](https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf)
7. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Dublin-NOCC/Expert-Scientific-Writer\\_REQ-10071592-2](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Dublin-NOCC/Expert-Scientific-Writer_REQ-10071592-2)
8. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Dublin-NOCC/Expert-Scientific-Writer\\_REQ-10071592-2](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Dublin-NOCC/Expert-Scientific-Writer_REQ-10071592-2)