

# Sr. Scientific Writer II

Job ID  
REQ-10036662  
Ene 30, 2025  
Irlanda

## Resumen

To write, edit and reconcile high quality medical and scientific communications including literature reviews, abstracts, posters and slide sets for submission to journal, congresses and/or clinical teams.

## About the Role

### Location

This role is either based in Dublin (Hybrid office/home) OR UK (Homebased)

### Major accountabilities:

- Prepares literature review, abstracts, posters, and slide sets working from various data sources including clinical study reports, patient profiles etc.
- Performs quality control (QC) checking / proof reading of above documents to meet customer expectation.
- Manages assigned individual projects.
- Obtains feedback from contributors and project teams.
- Complies with and support group's project management tool, standards, policies and initiatives.
- Follows Novartis specifications for documentation, templates etc.
- Maintains records for all assigned projects including archiving.
- Maintains audit, SOP and training compliance.
- Performs additional tasks as assigned.

### Key performance indicators:

- Preparation of the above referenced documents meeting set quality standards and on time for submission to Health Authorities/ Clinical teams / Journals as appropriate. (i.e. complying with standards e.g. CONSORT regarding publication of trial results, complying with journal formatting requirements etc).
- Publications are acceptable to internal and external authors (no issues with authorship).
- Completion of an adequate number of medical and scientific documents (taking into account complexity) per year.
- Adhere to Novartis values and behaviours.

### Minimum Requirements:

#### Education and Work Experience:

- **Minimum:** Life-science degree or equivalent B.Sc./equivalent with 4 years Clinical Research (CR) experience, M.Sc./M.Pharm + 2 years of CR experience **Desired:** Doctoral degree, Qualification in

Medical Sciences (MBBS/MD/equivalent). PhD + 1 year of CR experience, MBBS/equivalent + 1 year of CR experience

## **Skills:**

### **Functional competencies (Fundamental)**

- Results Driven; Customer/Quality Focus; Leadership; Innovative and creative; Action oriented; Show initiative; Empowerment / Accountability; Commitment / Self discipline; Mutual respect / Trust / Loyalty / Candor; Open Communication / Collaboration/Compassion; Drug Development knowledge; Science and Technology; Commercial Proficiency; Operational Excellence; Clinical communication & Info. Mgmt.

### **Leadership Competencies (Fundamental)**

- Sets clear direction and aligns team and others around common objectives
- Energizes the team
- Displays passion for the 3 Cs (Consumers, Customers, Competition)
- Exercises good judgment and drives change for competitive advantage
- Drives for superior results and has passion to win
- Displays analytical and conceptual thinking

### **Functional Experience**

- Scientific/ clinical knowledge of safety aspects, TA, disease, brand (Solid, Critical)
- Writing medical documents and publications (eg., abstracts, literature review, slide sets, posters, manuscripts, meeting reports) (Exposure, Critical)
- Clinical Research/ Drug Development (Exposure, Critical)
- Drug Safety (Exposure, Critical)
- Quality management (Exposure, Critical)
- IT/ web applications, office productivity tools and document formatting skills (Exposure, Critical)

### **Leadership Experience**

- Project Management (Exposure, Critical)
- Third Party (Customer/Vendor/Buyer) Relationship Management Exposure (Desired)
- Driving operational excellence Exposure (Critical)

### **Languages :**

- **Excellent written and oral English skills**

**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>

**Benefits and 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>

División  
Operations  
Business Unit  
Universal Hierarchy Node  
Ubicación  
Irlanda  
Sitio  
Dublin (NOCC)  
Company / Legal Entity  
IE02 (FCRS = IE002) Novartis Ireland Ltd  
Alternative Location 1  
London (The Westworks), Reino Unido  
Functional Area  
Research & Development  
Job Type  
Full time  
Employment Type  
Regular  
Shift Work  
No  
[Apply to Job](#)

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

Job ID  
REQ-10036662

## **Sr. Scientific Writer II**

[Apply to Job](#)

---

**Source URL:** <https://prod1.adacap.com/careers/career-search/job/details/req-10036662-sr-scientific-writer-ii>

### **List of links present in page**

1. <https://www.novartis.com/about/strategy/people-and-culture>
2. <https://talentnetwork.novartis.com/network>
3. <https://www.novartis.com/careers/benefits-rewards>
4. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Dublin-NOCC/Sr-Scientific-Writer-II\\_REQ-10036662](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Dublin-NOCC/Sr-Scientific-Writer-II_REQ-10036662)
5. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Dublin-NOCC/Sr-Scientific-Writer-II\\_REQ-10036662](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Dublin-NOCC/Sr-Scientific-Writer-II_REQ-10036662)

