NOVARTIS

Global Labelling Associate Director, Content (Cardiovascular, Renal & Metabolism)

Job ID
REQ-10082742
июл 09, 2026
Ирландия
Available in: English

Сводка

#LI-Hybrid

Location:
Dublin, Ireland

Relocation Support:
This role is based in Dublin, Ireland. Novartis is unable to offer relocation support: please only apply if accessible.

Great medicines deserve clear, compelling storytelling. As a Global Labelling Associate Director, Content (Cardiovascular, Renal & Metabolism), you will play a pivotal role in shaping how critical product information is communicated across global markets. Working at the intersection of science, strategy, and regulatory excellence, you will lead the development of high-quality global labelling content that supports patient safety, regulatory compliance, and commercial success. This is an opportunity to influence important healthcare decisions, collaborate with diverse international experts, and make a meaningful impact on established medicines that continue to improve patients' lives worldwide.

About the Role

Key Responsibilities

- Develop and maintain global labelling strategies and content for established products across development and lifecycle activities.
- Lead cross-functional discussions to achieve alignment on global labelling strategy, content, and key product claims.
- Drive the creation and maintenance of core and major market labelling documents.
- Present labelling recommendations, updates, and strategic proposals to governance bodies and project teams.
- Anticipate emerging labelling risks and develop mitigation, escalation, and contingency plans.
- Support health authority interactions by preparing evidence-based responses and negotiation strategies.
- Mentor colleagues and contribute to continuous improvement, inspection readiness, and knowledge sharing initiatives.

Essential Requirements

- Degree in a scientific discipline with strong understanding of pharmaceutical development and regulatory requirements.
- Experience developing and maintaining global labelling content for medicinal products across the product lifecycle.
- Ability to interpret clinical efficacy and safety data and translate findings into clear labelling content.
- Strong knowledge of global labelling regulations and requirements across major health authorities.
- Proven ability to collaborate effectively across multidisciplinary teams and influence diverse stakeholders.
- Excellent planning, prioritisation, communication, and attention to detail with a focus on quality.

Benefits & Rewards

At Novartis, we’re committed to reimagining medicine together - and rewarding the people who make it happen.

Expected Annual Base Salary Range for role: €75,880 to €140,920

The base salary offered is determined based on gender-neutral objectives, such as relevant skills, competencies and experience in accordance with the Novartis pay setting policy and upon joining Novartis will be reviewed periodically.

In addition to your base salary, you may be eligible for a performance-based bonus depending on certain performance parameters.

The rewards of being part of our team go far beyond base pay and incentives. We also offer a variety of competitive benefits in kind to help you thrive personally and professionally, such as insurance plans, retirement plans, wellbeing resources and global recognition programs. In addition, we provide flexible and hybrid working options, where possible, and minimum 14 weeks paid parental leave.

You may be eligible for a company vehicle or a car allowance in accordance with the applicable local Novartis policies and guidelines.

Pay equity is a fundamental principle of our employment policy and reflects our commitment to create a diverse, equitable and inclusive environment that treats all employees with dignity and respect, as outlined in our Code of Ethics.

Read our brochure to learn more about our global total rewards offering:

https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf

Note: Benefits and compensation may vary by country and are subject to local legal requirements, including provisions of collective bargaining agreements where applicable. A full overview of your compensation package, including any relevant collective bargaining agreement details applicable to your role based on your employment location and Novartis employer entity, will be communicated separately to you during the application process.

Commitment to Diversity and Inclusion / EEO paragraph

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)

Primary location salary range
€75,880.00 - €140,920.00
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

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List of links present in page

1. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
2. https://www.novartis.com/about/strategy/people-and-culture
3. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
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