

Global Labelling Associate Director, Content (Oncology)

Job ID
REQ-10079655
Июн. 02, 2026
Великобритания

Сводка

#LI-Hybrid (12 days per month on-site if living within 50 miles of our London office)
#LI-Remote (Homeworker if living further than 50 miles of our London office)
Office Location: London (The Westworks), United Kingdom

We are looking for a Global Labelling Associate Director, Content (Oncology) to contribute to the development and delivery of global labelling strategies across a portfolio of products.

In this role, you will support the creation of clear, consistent, and scientifically robust labelling content across development and lifecycle stages. You will work closely with cross-functional partners to ensure alignment on labelling strategy, enabling compliant and competitive product information for global markets.

About the Role

Major Accountabilities

- Develop and maintain global labelling strategies and core labelling documents for assigned products (e.g. CDS, USPI, EU SmPC/PIL).
- Lead and facilitate cross-functional discussions to support alignment on labelling content and strategy.
- Present labelling proposals and updates to governance bodies and project teams
- Identify emerging labelling considerations and contribute to planning, risk assessment, and mitigation strategies.
- Analyse competitor labelling, regulatory guidance, and scientific data to inform content development.
- Support and contribute to responses to Health Authority queries and interactions, including preparation of supporting documentation.
- Collaborate with global and regional partners to ensure consistency and alignment across markets.
- Mentor colleagues and contribute to audit readiness, inspections, and continuous improvement initiatives.

Essential Requirements

- Fluency in English (written and spoken).
- Demonstrated capability in Global Labelling and/or Global Regulatory Affairs, with a focus on labelling across development and lifecycle activities.
- Ability to develop and maintain core labelling documents (e.g. CDS) and support major market labelling (e.g. USPI, EU SmPC/PIL) with scientific accuracy and compliance.
- Ability to interpret clinical efficacy and safety data and translate it into clear, consistent labelling content and supporting documentation.
- Working knowledge of global labelling standards and expectations, including major Health Authority requirements.
- Strong collaboration and communication skills, with the ability to facilitate cross-functional discussions and support alignment.
- Strong planning, prioritisation, and attention to detail to deliver high-quality work within timelines.

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 \(PDF 30 MB\)](#)

Дивизион
Development
Business Unit
Development
Место
Великобритания
Сайт
London (The Westworks)
Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.
Alternative Location 1
Home Worker, Великобритания
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work

No

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