NOVARTIS

Global Labelling Manager, Content

Job ID
REQ-10077916
май 14, 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 Manager, Content to support the development and delivery of high-quality global labelling for our products.

In this role, you will contribute to ensuring that global product information is accurate, consistent, and compliant across markets. You will work closely with cross-functional partners to develop and maintain core labelling documents and support effective implementation across the product lifecycle.

About the Role

Major Accountabilities

- Act as labelling lead for assigned products, developing and maintaining global core labelling documents and major market labels (e.g. CDS, USPI, EU SmPC/PIL).
- Organise and facilitate cross-functional labelling discussions (e.g. ELTF meetings) to support alignment on content.
- Collaborate with Global Labelling colleagues and cross-functional teams to ensure consistent, compliant, and competitive labelling content.
- Conduct research across competitor labels, regulatory guidance, and clinical data to inform labelling updates.
- Prepare documentation to support labelling changes and contribute to responses to Health Authority queries.
- Support timely implementation of labelling updates across countries, ensuring alignment with global standards.
- Maintain high-quality documentation, including version control, references, and rationale for changes.
- Support audit and inspection readiness, mentoring colleagues where appropriate, and contributing to continuous improvement initiatives.

Essential Requirements

- Fluency in English (written and spoken).
- Demonstrated capability in Global Labelling, Regulatory Affairs, or a related pharmaceutical discipline.
- Working knowledge of core labelling concepts and major market formats (e.g. CDS, USPI, EU SmPC/PIL).
- Ability to interpret clinical and safety data and translate it into clear, consistent labelling content.
- Strong attention to detail, with the ability to maintain accurate documentation and traceability.
- Ability to collaborate effectively across cross-functional teams and manage competing priorities.

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.
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No

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List of links present in page

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