

# Global Labelling Associate Director, Content

Job ID  
REQ-10077914  
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 Associate Director, Content 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.

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

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