

# Global Labelling Manager

Job ID  
REQ-10020301  
Set 17, 2024  
Regno Unito

## Sommario

Our Development Team is guided by our purpose: to reimagine medicine to improve and extend people's lives.

To do this, we are optimizing and strengthening our processes and ways of working. We are investing in new technologies and building specific therapeutic area and platform depth and capabilities – all to bring our medicines to patients even faster.

We are seeking key talent, like you, to join us and help give people with disease and their families a brighter future to look forward to.

Apply today and welcome to where we thrive together!

The role:

This role offers hybrid working, requiring 3 days per week in our London office.

As Global Labelling Manager (GLM) you will be responsible for the creation and maintenance of regulatory compliant, competitive, and up-to-date core labelling documents for assigned developmental programs and Novartis Innovative Medicines products.

The assigned products will be for complex products and may include developmental programs.

The RA GLM also provides strategic and operational regulatory labeling input, working in close collaboration with Expert Labeling Task Force (ELTF) members in creating or maintaining core labeling documents. You will also handle Health Authority or Country Operations labeling queries for assigned products.

## About the Role

### Major accountabilities:

- You will maintain regulatory compliant, competitive and up-to-date global labelling documents for assigned products, presenting to the Global Labelling Committee on core data sheet changes.
- You will coordinate and lead the ELTF to agree labelling strategy, labelling course of action and text.
- You will also represent global labelling in relevant sub-teams, research and understand labelling topics across different markets, the competition and regulations.

- You will contribute to the creation of high-quality documents for Health Authorities and interact with Country Organisations to ensure the timely implementation of labelling changes in local product information.

**Your experience:**

- Bachelors or advanced life science degree.
- Experience in global labelling, or in related areas of the pharmaceutical industry or Health Authorities.
- Strong interpersonal, project management, communication and negotiation skills.
- Ability to lead cross functional teams in a complex work environment.
- Fluency in English

**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>

**Commitment to Diversity & Inclusion:**

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

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**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>

Divisione

Development

Business Unit

Innovative Medicines

Posizione

Regno Unito

Sito

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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