

Global Labeling Compliance & Project Lead

Job ID
REQ-10011598
ספט' 30, 2024
United Kingdom

תקציר

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

About the Role

:Major Accountabilities

This role offers hybrid working, requiring 3 days per week/ 12 days per month in our White City, London office.

:As Global Labelling Compliance Project Lead, you will be responsible for

- The tracking, data mining, monitoring and reporting on the global implementation of safety related product information changes e.g. Core Data Sheet amendments and updates. This includes routine interaction and follow-up with Country (CO) and Regional Regulatory Associates and other cross-functional teams, including Quality Assurance (QA), Process Improvement & Excellence (PIE) and Technical Operations
- Routine compliance reporting for global and local regulatory milestones of the Novartis safety label change process; support root cause analysis for any delays or non-compliance
- Supporting internal Pharmacovigilance (PV) CO audits, interacting with the COs to ensure timely and current safety label change (SLC) compliance data and reports are provided
- Supporting local and global PV inspections through preparation of requested SLC data and reports as required
- Supporting the evolution and maintenance of the Novartis safety label change tracking tool (SALTO) and

.related projects as a SALTO Super User

Providing input for process improvements, training manuals and materials, IT labelling projects and user •
.testing

.Supporting development of Novartis systems and company projects •

:Your Experience

Bachelor's degree required with relevant regulatory and/or technical experience gained in the •
.pharmaceutical industry

Knowledge of Regulatory Affairs, specifically labelling regulations globally, labelling submission business •
.processes and related tools

Proven knowledge of data analysis and reporting, excellent verbal and written communication skills, •
technical/IT skills and proven ability to be innovative and a creative problem solver with quality and
.compliance approach

.Fluency in English as a business language required •

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>

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up:
<https://talentnetwork.novartis.com/network>

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>

מלוקה

Development

Business Unit

Innovative Medicines

מיקום

United Kingdom

אתר

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

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