

Global Program Safety Lead - CRM

Job ID
REQ-10065643

11月 07, 2025

Spain

摘要

Are you ready to make a significant impact in medical safety?

Novartis is seeking passionate individuals to join our innovative Cardio Renal Metabolism team. You will work with cutting-edge technology, collaborate with experts, and drive excellence in patient safety.

If you're eager to advance your career and be part of a forward-thinking company, we want you on our team!

About the Role

#LI HYBRID

Location: Barcelona, Spain or London, United Kingdom

Key Responsibilities:

- **Safety Input and Team Participation:** Provide expert safety input to the clinical development program for assigned projects/products and actively participate in the Global Program Team (GPT), Global Clinical Team (GCT), and Clinical Trial Team (CTT). Lead the Safety Management Team (SMT). Responsible for managing safety issues from the formation of the GPT through Life Cycle Management.
- **Safety Strategy Preparation:** Own the projects/products safety strategy and lead the production of medical safety deliverables.
- **Core Global Labelling:** Responsible for initial development and ongoing maintenance of safety information in Core Data Sheet, addressing safety issues in all project/product indications.
- **Signal Detection and Safety Management:** Oversee overall signal detection, monitoring, evaluation, interpretation, and appropriate management of safety information, based on data from all relevant sources.
- **Regulatory and Professional Inquiries:** Lead the preparation of the safety strategy for health authority responses and collaborate with other project team members. Respond to inquiries from regulatory authorities or healthcare professionals regarding safety issues.
- **Departmental and Functional Goals:** Contribute to and often lead the development of departmental and functional/business unit goals and objectives. Proactively engages in the development of competencies across the Medical Safety Function

Role Requirements:

- Medical Degree or equivalent is preferred; PhD, PharmD or equivalent graduate level health care professional degree required
- 3 years clinical experience postdoctoral
- Minimum 5 years ' experience in drug development in a major pharmaceutical company (of which 2 years in a global position), including 2 years in safety at an operational or medical position.
- Experience in clinical trial methodology, regulatory requirements, scientific methodology, statistics and writing of publications
- Expertise in preparing or contributing to preparation of clinical safety assessments and regulatory reports/submissions involving safety information
- Experience with (safety or others) issue management
- Experience in leading cross-functional, multicultural teams

Languages:

- Fluent English (both spoken and written) is mandatory.

Additional desirable skills:

- Additional languages are an advantage.
- CRM experience

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>

部门

Development

Business Unit

Universal Hierarchy Node

地点

Spain

站点

Barcelona Gran V í a

Company / Legal Entity

ES06 (FCRS = ES006) Novartis Farmac é utica, S.A.

Alternative Location 1

London (The Westworks), United Kingdom

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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