Country Risk Management Plan Manager , Patient Safety the Netherlands

Job ID
REQ-10055565

7月 22, 2025

Netherlands

摘要

#LI-Hybrid
Location: Amsterdam, Netherlands
Internal job title: CO RMP Manager PS

Step into a pivotal role where your expertise will directly shape patient safety across the Netherlands. As CO RMP Manager PS, you ' ll be the driving force behind the local implementation of global Risk Management Plan (RMP) commitments–ensuring compliance, cross-functional alignment, and timely delivery. You ' ll collaborate with diverse teams, support audits and inspections, and play a key role in safeguarding public health. If you're passionate about making a meaningful impact through defined processes and strategic oversight, this is your opportunity to lead with purpose.

About the Role

Key Responsibilities

- Lead local implementation of Risk Management Plan (RMP) commitments across all Novartis entities.
- Align cross-functional teams to ensure consistent execution of RMP processes and deliverables.
- Monitor and verify timely rollout of RMP educational materials by relevant departments.
- Maintain local archiving processes for RMP documentation and training records.
- Ensure local preparation and updates of RMP annexes or full plans as required.
- Deliver and document RMP training for all relevant associates, including third parties.
- Act as country liaison for global RMP systems, ensuring accurate tracking and compliance.
- Provide local input to global RMP teams on implementation metrics and regulatory updates.
- Support audits and inspections by ensuring readiness and documentation of RMP activities.
- Escalate non-compliance issues and contribute to continuous improvement of RMP practices.

Essential Requirements

- Degree in Health Care Sciences or equivalent higher education, training, and experience
- Proficient in both English and Dutch, with strong written and verbal communication, interpersonal and negotiation skills
- Solid understanding of safety and Risk Management Plan (RMP) processes
- Strong knowledge of pharmacovigilance and RMP regulatory requirements
- Proven ability to work cross-functionally and influence within a matrix organization
- Highly organized, results-driven, and committed to compliance and quality standards

Desirable Requirements

- Experience delivering training programs related to safety or compliance
- Familiarity with pharmacovigilance agreements and case processing activities

Commitment to Diversity & Inclusion:

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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

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部门
Development

Business Unit
Universal Hierarchy Node

地点
Netherlands

站点
Amsterdam

Company / Legal Entity
NL08 (FCRS = NL008) Novartis Pharma NL

Functional Area
Research & Development

Job Type
Part time

Employment Type
Tijdelijk (bepaalde tijd)

Shift Work

No

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