Patient Safety Manager

Job ID
REQ-10003344

9月 04, 2024

France

摘要

Suit et audite le programme de contr ô le des m é dicaments, produits biologiques ou m é dicaux de l'entreprise, dont la r é ception, l' é valuation, le traitement et le suivi des rapports pr é judiciables. Participe à la r é solution de toute obligation l é gale et au respect des r è gles gouvernementales. Assure une r é ception, maintenance et é valuation pr é cises face à l' é tiquetage des produits. Signale les incidents ou r é actions selon les exigences des r é gulateurs, y compris les donn é es des effets n é gatifs des essais, les sources spontan é es ou sollicit é es, les rapports p é riodiques et d'exp é rience. Peut fournir tendances et d é tection et é valuation des signaux de s é curit é . Soutient toutes les activit é s d'essais cliniques et apr è s vente

About the Role

Key Responsibilities:

- To be the accountable for specific operational vigilance process(es) at the Country

Organization
- To mentor less experienced staff, maintaining a professional network of key contacts and role model Novartis values and behaviors.
- Ensure oversight and compliance in terms of reporting/submission/distribution of safety reports/updates/information (e.g., SAE, SR, IN, SUSAR, PSUR, DSUR, changes in risk-benefit profile) to Local Health Authorities (LHA) according to regulatory requirements and Novartis procedures.
- Work in close collaboration other local and global medical safety functions to ensure accurate evaluation of safety data.
- Interact and exchange relevant safety information with Health Authorities, other functional groups, third-party contractors, and PS associates, as applicable.
- Monitor national pharmacovigilance regulations and provide update to global PS organization.
- Set up, update, and implement local procedures to ensure compliance with PS global procedures and national requirements.
- Ensure local PS-related RMP commitments are executed and properly documented.

Essential Requirements:

- Education : Health Care Sciences Professional (e.g. Medical Doctor, Nurse, Pharmacist)
- Excellent communications and negotiation (networking) skills.
- Quality focused and results oriented.
- 2 years ' experience in pharmacovigilance or equivalent field
- Knowledge of national and international regulations for pharmacovigilance

Desirable Requirements:

- Fluent in both written and spoken English and French
- Project management skills

Why Novartis? Our purpose is to reimagine medicine to improve and extend people ' s lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: https://www.novartis.com/about/strategy/people-and-culture

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Commitment to Diversity & Inclusion: 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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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
Innovative Medicines

地点
France

站点
Paris Headquarter (Novartis Pharma S.A.S.)

Company / Legal Entity
FR12 (FCRS = FR012) Novartis Pharma S.A.S.

Functional Area
Research & Development

Job Type
Full time

Employment Type
CDI

Shift Work
No

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