Regulatory Affairs Specialist

Job ID
REQ-10023226

9月 20, 2024

Serbia/Monten.

摘要

贡献和支持提交产品注册，进度报告，补充，修订和/或定期经验报告的发展。支持该部门的所有注册活动，以确保符合当地药品监管环境的要求。

About the Role

Major Accountabilities

·根据注册计划，通过具有商业吸引力的标签实现最佳产品注册
·根据当地法规/法律/准则、公司战略和全球合规性，维护并保护
CMC/CDS/安全更新方面的产品许可证
·确保遵守 NP4、KRPIA 行为准则、相关 CPO
活动的相关法规和法律发更新、RMP、包装材料、促销材料/活动、PMS/药物安全报告等）
·促进和保持与内部和外部利益相关者之间的良好关系
·在收到诺华产品后24小时内报告与诺华产品相关的技术投诉/不良事件/特殊情况

~营销样本的分发如适用）

Key Performance Indicators

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Work Experience

运营管理和执行
项目管理
职能广度
跨文化经历

Skills

分析能力
项目规划
临床试验
协作
生命科学
注重细节
法规遵从性

Language

英语

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
Innovative Medicines

地点
Serbia/Monten.

站点
Serbia

Company / Legal Entity
RSP0 (FCRS = CH024) NPHS RO Serbia

Functional Area
Research & Development

Job Type
Full time

Employment Type
正式

Shift Work
No

Apply to Job

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