U NOVARTIS

Global Clinical Operations- Supply Chain Expert

Job ID REQ-10046370

Apr 25, 2025

China

Summary

在适当的监督下,为全球临床试验的各个方面做出贡献,在时间表,预算,质量/合规性和绩效标准范围内提供研究成果。可能引领全球临床试验的具体方面。临床试验团队的核心成员,通过流程改进和知识共享,为卓越运营做出贡献

About the Role

Key responsibilities:

Customs and Trade Expertise

- Review, assign and monitor correct customs attributes in NVS trade system for clinical shipments (clinical/ancillary supplies)
- Ensure valid import and export licenses are available. Execution of import/export customs clearances and coordination of logistics to ensure activities in accordance with trade

compliance policy and SOP.

- Ensure and validate the correctness of customs declarations and liaise with the relevant internal and external partners where post-declaration amendments are required.
- Act as point of contact of SSO China in case of queries from Customs or related authorities.
- Liaise and communicate with Customs and Suppliers/Agencies related to the Customs and Trade Topics.
- Cross-function collaboration with Country Customs and Trade Compliance, Global Funds Flow to ensure the Global Customs Valuation is up-to-date.
- Be responsible for Customs agency/supplier management and supervision, ensure they strictly follow NVS standard.
- Support the feasibility from clinical supply chain perspective at early stage of trial.
- Maintain accurate records of trade activities, including documentation, license, agreements, in accordance with legal and regulatory requirements.
- Stay current with changes in international trade laws and China regulations.
- Responsible for the tariff supplementary with Customs and NVS commercial/finance team after product is launched.

Essential requirements :

- Minimum of 3-5 years of experience in Customs and international trade function of MNC
- Minimum 2-4 years ' experience in Customs clearance, trade compliance and clinical supply chain management or equivalent
- Must keeping up to date with changes in Customs/NMPA regulations

Desirable requirements :

- Good understanding of Customs/NMPA related regulations and trade compliance requirements
- Ability to proactively recognize and solve problems
- Excellent communication skills: ability to effectively communicate with stakeholders at various levels, including internal cross-functional teams, external agencies and HAs.
- Strong negotiation skills to ensure alignment and collaboration in the supply chain process.
- Knowledge of GCP is highly desirable
- Ability to build strong working relationships with supply chain internal and external stakeholders
- Strong attention to detail in managing complex supply chain activities
- Must be proactive, team player. Ability to multitask, prioritize tasks
- Project Management skills required to manage and/or support project supply chain to meet tight deadlines
- Well-structured, good analytical and conceptual thinking
- Strong process-orientation including ability to define and document business processes
- Able to adjust to the changing and different job requirements

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? <u>https://www.novartis.com/about/strategy/people-and-culture</u>

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部門 Development

部門 Innovative Medicines

国 China

勤務地 Beijing (Beijing)

Company / Legal Entity CN14 (FCRS = CN014) China Novartis Institutes for BioMedical Research Co., Ltd.

Alternative Location 1 Shanghai (Shanghai), China

Functional Area Research & Development

職種 Full time

雇用形態

正式

Shift Work No

Apply to Job.

无障碍及便利 设 施

诺华 承 诺 与残障人士共事并 为 他 们 提供合理的便利 设 施。如果您由于健康状况或残障 在招聘 过程的任何 环 节 需要合理便利 设 施 或者 为 了履行 职 位的基本 职 能 请发 送 电 子 邮 件至 <u>diversityandincl.china@novartis.com</u> 告知您的需求和 联 系方式,并在 邮 件中附上您的 职 位申 请编 号。



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