

生产技术员II

Job ID
REQ-10053841

7月 11, 2025

China

摘要

生产技术员在放射配体疗法Radioligand Therapies, 简称 RLT的日常生产中发挥积极作用, 同时负责仪器和设备的设置与准备。生产技术员在履行职责时遵守监管要求, 按照批记录和标准操作规程SOP执行生产任务。相关职责在团队中完成, 并根据指定的生产轮班时间表进行。生产技术员与经理和主管密切合作, 确保生产过程安全、及时地执行。

About the Role

主要职责：

- 执行与放射配体疗法RLT产品制造相关的所有活动。职责包括操作和维护A级隔离器, 专注于关键绩效指标 (KPI 目标, 并确保遵守所有州级、联邦及诺华的辐射安全指南。
- 负责按时成功完成所需的培训

- 课程,包括必要的标准操作程序(SOP
,无菌技术,更衣资格认证以及与该岗位相关的其他培训包括健康、安全与环境 HSE。)
- 支持与生产准备相关的所有技术方面,包括手动清洁工作单元和使用过氧化氢蒸汽对隔离器进行灭菌。
 - 根据要求进行常规和动态的环境监测。
 - 推动“勇于发声”的文化,确保所有cGMP合规活动得到遵守。
 - 准备相关文件和记录,如批记录、运输文件和培训材料。
 - 鉴于产品的时间敏感性,该职位可能需要临时加班以确保流程的连续性和完成。
 - 灵活穿戴洁净室服装和个人防护装备(RPE)
 - 近距离视力相当于20/20,且无色觉障碍。允许使用矫正镜片以达到所需的视力水平。
 - 在受限区域内禁止使用化妆品、佩戴首饰、涂指甲油、使用香水/古龙水及其他可能成为微生物来源的物品。
 - 搬运或提举最多达35磅约16公斤的物品。
 - 根据需要,可能会分配其他职责。

关键绩效指标:

- 批记录执行“第一次即正确”,无重复偏差。
- 遵守出勤指南及所有与安全相关的操作程序。
- 与A级隔离器或B/C级区域相关的审计中无重大或关键性发现。
- 生产过程符合所有良好文件规范(GDP和良好生产规范(GMP原则。
- 无菌/洁净室行为符合GMP指南要求。

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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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
Operations

Business Unit
Innovative Medicines

地点
China

站点
Haiyan (Zhejiang Province)

Company / Legal Entity
CN27 (FCRS = CN027) Novartis Pharmaceutical Technology Zhejiang Co., Ltd.

Functional Area
Technical Operations

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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