Production Lead (Nights)

Job ID
REQ-10050887

May 05, 2025

USA

Summary

Production Leads play an active role in daily production of isotope manufacturing as well as setup and preparation of instruments and equipment. The Production Lead adheres to regulatory requirements while performing job functions, executing production as per batch records and SOPs. Responsibilities are performed within a team and according to an assigned production shift schedule. The Production Lead works closely with the Production Manager and Shift Supervisor to ensure production is executed in a safe and timely manner. This role will transition to night shifts in the future.

About the Role

Major accountabilities:

- Executes all activities related to the manufacturing of RLT isotope products. Responsibilities include operating and maintaining grade C isolators, focusing on KPI goals as well as

ensuring all state, federal and Novartis radiation safety guidelines are adhered to.
- Responsible for successful on time completion of required training curriculum comprising of the necessary Standard Operating Procedures (SOPs) and Techniques, Gowning Qualifications and other relevant training including HSE for the specific role.
- Supports all technical aspects related to production readiness including manually cleaning the cell and performing sterilization of the isolators.
- Conducts routine and dynamic environmental monitoring as required.
- Prepares all materials while maintaining material identity in accordance with the batch monitoring system as defined by procedure.
- Participation in assigned qualification/validation activities, as necessary.
- Facilitates a culture of "speaking up" and ensuring all cGMP compliance activities are followed.
- Prepares applicable documents and records such as batch records, shipping documents, and training materials.
- Participates in periodic mandatory overtime to ensure process continuity and completion.
- Ensures technicians complete all required training in accordance with published curriculum.
- Participate in technician professional development counselling to foster a growth culture.
- Other duties may be assigned, as necessary.

Essential Requirements:

- Training in radiochemistry or radio pharmacy is preferred.
- 4+ years of experience in pharmaceutical manufacturing, with low bioburden manufacturing preferred.
- Good understanding of manufacturing and validation requirements and activities.
- Exploitation of new technology and techniques to eliminate non-value adding activities and improve productivity / performance through new processes.
- Knowledge of cGMP regulations and FDA guidance applicable to isotope manufacturing.
- Proficient in MS Office applications.
- Flexibility to don clean room garments and personal protective equipment (PPE).
- Near vision performance should be the equivalent of 20/20 with no impairment of color vision. The use of corrective lenses to achieve the desired visual acuity is permitted.
- Makeup, jewelry, nail polish, perfume/cologne and other potential microbial sources are prohibited in restricted areas.
- Ability to lift or carry up to 35 pounds.
- Bachelor of Science strongly preferred. If the applicant does not have a degree, a minimum of 2 years of experience in a cGMP or aseptic environment can be substituted

#LI-Onsite

Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: novartis-life-handbook.pdf.

The pay range for this position at commencement of employment is expected to be between $77,0000 and $ 143,000 per year; however, while salary ranges are effective from 1/1/25 through 12/31/25 fluctuations in the job market may necessitate adjustments to pay ranges during this period.

Further, final pay determinations will depend on various factors, including, but not limited to geographical location, experience level, knowledge, skills and abilities. The total compensation package for this position may also include other elements, including a sign-on bonus, restricted stock units, and discretionary awards in addition to a full range of medical, financial, and/or other benefits (including 401(k) eligibility and various paid time off benefits, such as vacation, sick time, and parental leave), dependent on the position offered. Details of participation in these benefit plans will be provided if an employee receives an offer of employment. If hired, employee will be in an "at-will position" and the Company reserves the right to modify base salary (as well as any other discretionary payment or compensation program) at any time, including for reasons related to individual performance, Company or individual department/team performance, and market factors.

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

EEO Statement:

The Novartis Group of Companies are Equal Opportunity Employers. We do not discriminate in recruitment, hiring, training, promotion or other employment practices for reasons of race, color, religion, sex, national origin, age, sexual orientation, gender identity or expression, marital or veteran status, disability, or any other legally protected status.

Accessibility & Reasonable Accommodations

The Novartis Group of Companies are committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the application process, or to perform the essential functions of a position, please send an e-mail to us.reasonableaccommodations@novartis.com or call +1(877)395-2339 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

部門
Operations

部門
Innovative Medicines

国
USA

State
Indiana

勤務地
Indianapolis

Company / Legal Entity
U469 (FCRS = US469) AAA USA Inc.

Functional Area
Technical Operations

職種
Full time

雇用形態
Regular

Shift Work
No

Apply to Job.

Job ID
REQ-10050887

Production Lead (Nights)

Apply to Job.

Source URL:
https://prod1.novartis.com/jp-ja/careers/career-search/job/details/req-10050887-production-lead-nights

List of links present in page

1. https://www.novartis.com/sites/novartiscom/files/novartis-life-handbook.pdf
2. https://www.novartis.com/about/strategy/people-and-culture
3. https://talentnetwork.novartis.com/network
4. https://www.novartis.com/careers/benefits-rewards
5. mailto:us.reasonableaccommodations@novartis.com
6. https://novartis.wd3.myworkdayjobs.com/en-US/NovartisCareers/job/Indianapolis/Production-Lead--Nights-REQ-10050887

7. https://novartis.wd3.myworkdayjobs.com/en-US/NovartisCareers/job/Indianapolis/Production-Lead--Nights-REQ-10050887