NOVARTIS

Production Technician I (weekdays)

Job ID
REQ-10078421
май 27, 2026
США

Сводка

#LI-Onsite

Location: Indianapolis, IN, United States

Relocation Support: This role is based in Indianapolis, IN, United States. Novartis is unable to offer relocation support: please only apply if accessible.

Join a team at the forefront of radioligand therapy manufacturing and play a vital role in delivering innovative treatments to patients who need them most. As a Production Technician I, you will be hands-on in a highly controlled, cutting-edge environment, ensuring every product is manufactured safely, accurately, and on time. This is an opportunity to build your expertise in aseptic techniques, advanced manufacturing systems, and regulatory compliance, while contributing directly to life-changing therapies. If you are energized by fast-paced production environments and take pride in precision, quality, and teamwork, this role offers the chance to make a meaningful impact every day.

Shift: Monday- Thursday 6am- 6pm and then Monday- Wednesday 6 am-6pm
mandatory overtime may be required

About the Role

Key Responsibilities:

· Manufacture radioligand therapy products by following batch instructions and maintaining strict aseptic discipline.

· Operate and maintain Grade A isolators to meet safety, quality, and key performance indicator targets.

· Follow radiation safety requirements and comply with all state, federal, and Novartis guidelines.

· Complete required training in Standard Operating Procedures, aseptic technique, gowning qualification, and Health, Safety, and Environment.

· Clean production cells manually and sterilize isolators to ensure production readiness.

· Perform routine and dynamic environmental monitoring per procedure and document results accurately.

· Prepare materials, maintain identity and traceability, and update the batch monitoring system as required.

· Execute work in alignment with current Good Manufacturing Practice and site quality standards.

· Support qualification and validation activities; contribute to deviation investigations and inspection readiness.

· Prepare and maintain batch records, shipping documentation, and training materials to ensure compliant operations.

Essential Requirements:

· Bachelor’s degree in relevant Engineering or Scientific discipline is highly preferred; If the applicant does not have a degree, a minimum of 1+ year’ of experience in cGMP or aseptic environment is required.

· Working knowledge of current Good Manufacturing Practice for sterile or aseptic manufacturing environments.

· Ability to follow United States Food and Drug Administration guidance relevant to aseptic manufacturing.

· Ability to gown aseptically and work extended periods in a Grade C cleanroom environment.

· Strong attention to detail and ability to complete batch records and controlled documentation accurately.

· Near vision equivalent to 20/20 with no color vision impairment; corrective lenses permitted.

· Ability to lift or carry up to 35 pounds and perform hands-on production tasks.

· Reliable teamwork and communication skills to succeed in a fast-paced, shift-based schedule.

Desirable Requirements:

· Radiopharmaceutical manufacturing experience preferred, especially in aseptic or cleanroom operations.

· English proficiency for reading, writing, and speaking in a regulated manufacturing environment.

Novartis Compensation and Benefits:

The salary for this position is expected to range between $25.19/hr. and $46.83/hr.

The final salary offered is determined based on factors like, but not limited to, relevant skills and experience, and upon joining Novartis will be reviewed periodically. Novartis may change the published salary range based on company and market factors.

Your compensation will include a performance-based cash incentive and, depending on the level of the role, eligibility to be considered for annual equity awards.

US-based eligible employees will receive a comprehensive benefits package that includes health, life and disability benefits, a 401(k) with company contribution and match, and a variety of other benefits. In addition, employees are eligible for a generous time off package including vacation, personal days, holidays and other leaves

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

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

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.

Дивизион
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Production / Manufacturing
Место
США
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Indiana
Сайт
Indianapolis
Company / Legal Entity
U469 (FCRS = US469) AAA USA Inc.
Functional Area
Technical Operations
Job Type
Full time
Employment Type
Regular
Shift Work
No

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