NOVARTIS

Specialist, QA Operations (Wed-Sat, 2nd Shift)

Job ID
REQ-10093377
Ноя. 22, 2026
США

Сводка

Location: Millburn, NJ

Relocation Support: This role is based in Millburn, New Jersey. Novartis is unable to offer relocation support: please only apply if accessible.

Please note that Novartis is unable to provide visa or immigration support for this position. Applicants must have authorization to work in the United States without requiring current or future sponsorship.

Bring your Quality expertise to our Millburn Radioligand Therapy site, where your work directly supports product quality, compliance, and timely delivery to patients. As QA Operations Specialist, you will provide hands-on Quality Assurance oversight of production and testing activities, with a strong presence on the shop floor. You will review and approve critical production and Quality records, support investigations and change controls, and help maintain cGMP, data integrity, and electronic compliance while collaborating across teams to strengthen Quality performance and drive continuous improvement.

#LI-Onsite

About the Role

Key Responsibilities

- Provide shop-floor Quality Assurance oversight for production and testing activities, ensuring compliant execution.
- Review and approve production records for completeness, accuracy, compliance, and timely disposition.
- Review Master Batch Records and associated documentation to support compliant manufacturing operations.
- Investigate non-critical deviations, assess potential impact, and support appropriate corrective and preventive actions.
- Initiate and manage non-complex change controls, ensuring Quality requirements and implementation activities are addressed.
- Partner across functions to resolve Quality issues and advance operational excellence and continuous improvement.

Essential Requirements

- University degree with a scientific or technological background, such as Chemistry, Pharmacy, Biology, Biochemistry, or equivalent.
- 3+ years of experience in pharmaceutical Quality Control, Quality Assurance, or Production.
- Good knowledge of cGMP and applicable Quality standards.
- Experience collaborating across functions, with a results-oriented approach and focus on operational excellence.
- Proficiency with Microsoft Office and standard applications supporting Quality activities.

The salary for this position is expected to range between $43.08 and $80.00 per hour. 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.

To learn more about the culture, rewards and benefits we offer our people clickhere.

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
Operations
Business Unit
Quality
Место
США
Состояние
New Jersey
Сайт
Millburn
Company / Legal Entity
U473 (FCRS = US473) Novartis Innovative Technologies Inc
Functional Area
Quality
Job Type
Full time
Employment Type
Regular
Shift Work
No

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