NOVARTIS

Quality Control, Specialist

Job ID
REQ-10073806
мар 18, 2026
США

Сводка

At Novartis, we are reimagining medicine to improve and extend people's lives. The Specialist, Quality Control, assists and supports the organization with compliance and ongoing preparation, testing and monitoring conformance to established quality processes and standards for manufacturing and production.

#LI-Onsite
Shift: 1st

Location: Durham, NC
This role is based in Durham, NC. Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

About the Role

Key Responsibilities

- Responsible for directing and executing method transfers/verifications and assisting in execution of bioanalytical (BAL) analysis for cGMP release and characterization testing using techniques including but not limited to CE-SDS, SDS-PAGE and CZE methods.
- Supports BAL assay onboarding/execution, including drafts and maintenance of QC SOPS, forms, protocols, reports, Analytical Master Plans and change controls.
- Reviews and approves data for reports and specification compliance; supports and leads resolution of nonconformances (SSFs, deviations, OOS/OOT/OOE investigations)
- Assist with enrollment of vendors, ordering reagents and consumables for new assays.
- Assists in the evaluation of internal controls, communications, risk assessments and maintenance of documentation as related to compliance with internal and external safety, quality, and regulatory standards.
- Performs qualifications, maintenance and trend analysis of bioanalytical methods assays/critical controls & standards and draws conclusions.
- Capable of delivering to assigned work schedule with attention to detail and accuracy.
- Support department risk assessments and participate in audit walkthroughs.
- Oversee special projects on analytical and instrument problem solving. May develop testing and analysis methods and procedures in accordance with established guidelines.
- Supports training of departmental personnel in appropriate techniques and related topics.
- Other related job duties as assigned.

Essential Requirements

- Bachelor's degree in scientific disciplines such as Biochemistry, Biology or related field required.
- 5 years of experience in GMP environment
- Excellent interpersonal, verbal and written communication skills with strong technical writing experience required. Previous investigation experience a plus.
- Proven ability to work effectively in a team environment. Collaborates cross functionally with other departments to achieve site goals.
- Works on problems of moderate scope where analysis of situations or data requires a review of a variety of factors.
- Exercises judgment within defined procedures and practices to determine appropriate action including critical thinking, troubleshooting and problem-solving skills.
- Normally receives general instructions on routine work, detailed instructions on new projects or assignments.
- Self-motivated, detail-oriented, and willing to accept temporary responsibilities outside of core duties.

The salary for this position is expected to range between $41.06 and $76.25 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.

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
North Carolina
Сайт
Durham
Company / Legal Entity
U473 (FCRS = US473) Novartis Gene Therapies
Functional Area
Quality
Job Type
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Shift Work
No

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Quality Control, Specialist

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