NOVARTIS

Specialist, Quality Control - Raw Materials

Job ID
REQ-10073796
Mar 12, 2026
LOC_US

About the Role

Location:

- This position will be located in Durham, NC and will be an On-Site role.
- Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

Key Responsibilities:

- Execute routine and non-routine analysis for cGMP release and characterization testing using techniques including but not limited to chromatography (HPLC, UPLC), AUC, SEC, compendial assays (pH, Conductivity, Osmolality), electrophoresis (CE, Western Blot) and assist with data review.
- Execute method verifications/transfers as required for various projects.
- Assist with enrollment of vendors, ordering reagents and consumables for new assays.
- Initiate all paperwork (LIMS, SOPs, assay forms) for testing and transfers of methods for projects.
- 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.
- Reviews data obtained for compliance with specifications and reports abnormalities. Performs trend analysis of methods /environmental data / assay 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.
- Participates in the preparation of investigations, summaries and reports. Reviews data obtained for compliance with specifications and reports. Investigates and resolves non-conforming test results by completing thorough Deviation, OOS/OOT/OOE and Investigation.
- Authors new/revise Standard Operating Procedures, Protocols / Summary Reports / Analytical Master Plans for QC.
- 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, Microbiology or related field with 5 years’ experience in GMP environment or 4 years’ at GTx.
- 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.

Novartis Compensation and Benefit Summary:

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.

#LI-Onsite

Role Requirements

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)

Division
DIV_TO
Business Unit
Quality
Location
LOC_US
Site
Durham
Company / Legal Entity
U473 (FCRS = US473) Novartis Gene Therapies
Functional Area
FCT_QA
Job Type
Full time
Employment Type
Regular
Shift Work
No
Apply to Job

Job ID
REQ-10073796

Specialist, Quality Control - Raw Materials

Apply to Job

Source URL: *https://prod1.jobapi.novartis.com.cn/req-10073796-specialist-quality-control-raw-materials*

List of links present in page

1. https://prod1.jobapi.novartis.com.cn/req-10073796-specialist-quality-control-raw-materials
2. https://www.novartis.com/about/strategy/people-and-culture
3. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Durham/Specialist--Quality-Control---Raw-

Materials_REQ-10073796-1
5. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Durham/Specialist--Quality-Control---Raw-Materials_REQ-10073796-1