

Analyst, Quality Control - Raw Materials

Job ID

REQ-10073795

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.
- Perform sampling of incoming materials as required for use in manufacturing and production.
- Initiate all paperwork (LIMS, SOPs, assay forms) for testing and transfers of methods for projects.
- Responsible for limited range of laboratory support functions and procedures as assigned, developing capability in basic technical skills, disciplines, and procedures within assigned discipline area(s).
- May be assigned to specific disciplines, but will support all necessary laboratory and assay functions, including housekeeping, safety, logbook/equipment use and maintenance, and updates to existing operating procedures.
- Capable of delivering to assigned work schedule with attention to detail and accuracy.
- Notifies management and initiates events (such as Laboratory Investigations) in the quality systems, with guidance from senior analysts or management.
- Assist in special projects on analytical and instrument problem solving by execution of assay.
- Gain familiarity with basic process improvement methodologies, learning and applying concepts of lean lab and six sigma that are applicable to the QC lab environment.
- Other related job duties as assigned.

Essential Requirements:

- Bachelor's degree in scientific disciplines such as Biochemistry, Biology, Microbiology or related field.
- Learns to use professional concepts.
- Applies company policies and procedures to resolve routine issues.
- Ability to communicate and work in a team environment.
- Normally receives detailed instructions on all work.

Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between \$32.12 and \$59.62 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

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