

Head of Quality Control

Job ID
REQ-10039124
Feb 06, 2025
USA

Summary

The Head of Quality Control is responsible for designing and implementing innovative and robust processes for testing activities associated with a viral gene therapy manufacturing site which includes bulk product and sterile fill operations.

Location- Durham, NC #LI- onsite

About the Role

Key Responsibilities:

- Develops and executes corporate quality policies, practices, procedures, standards, and systems necessary to ensure adherence to aseptic production and product management in accordance with the cGXP compliance to US and EU regulations.
- Oversees the testing and validation strategy, with a concentrated focus on method verification and validation.
- Maintains a robust Environmental Monitoring program that meet US/EU and other applicable regulatory requirements.
- Organizes and directs cross functional relationships with Manufacturing, Engineering, MS&T, Quality Assurance, and Regulatory.
- Hires staff and manages contract vendors for programs related to focus area.
- Directs laboratory staff, set goals and expectations, and maintain efficient utilization of resources.
- Authors and approve documents required for regulatory submissions.
- Develops, implements, and ensures laboratory procedures and policies are followed.
- Provides presentations, explain laboratory qualification and operations, and defend testing results during FDA and other inspections.
- Other duties for which QC is responsible, as assigned.

Requirements:

- BS/MS in Microbiology discipline and 12 years of related experience in cGMP laboratory environment, with strong knowledge of regulatory, USP and Eur. Phr guidelines (Ph.D. in life sciences is preferred).
- 8 years of laboratory management experience.
- Experience starting up lab facilities supporting clinical and/or commercial manufacturing.
- Experience with regulatory agency inspections.

Novartis Compensation and Benefit Summary: The pay range for this position at commencement of employment is expected to be between \$138,600 and \$257,400 /year; *however, while salary ranges are*

effective from 1/1/25 through 12/31/25, fluctuations in the job market may necessitate adjustments to pay ranges during this period. Further, final pay determinations will depend on various factors, including, but not limited to geographical location, experience level, knowledge, skills, and abilities. The total compensation package for this position may also include other elements, including a sign-on bonus, restricted stock units, and discretionary awards in addition to a full range of medical, financial, and/or other benefits (including 401(k) eligibility and various paid time off benefits, such as vacation, sick time, and parental leave), dependent on the position offered. Details of participation in these benefit plans will be provided if an employee receives an offer of employment. If hired, employee will be in an “at-will position” and the Company reserves the right to modify base salary (as well as any other discretionary payment or compensation program) at any time, including for reasons related to individual performance, Company or individual department/team performance, and market factors.

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>

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Division
Operations
Business Unit
Innovative Medicines
Standort
USA
State
North Carolina
Site
Durham
Company / Legal Entity

U473 (FCRS = US473) Novartis Gene Therapies

Functional Area

Quality

Job Type

Full time

Employment Type

Regular

Shift Work

No

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