

QC Microbiology Technician

Job ID
REQ-10068443

12月 16, 2025

USA

摘要

The QC Microbiology Technician is responsible for assisting routine and batch-related Environmental Monitoring in controlled environments, following current Good Manufacturing Practices.

About the Role

LOCATION: This opportunity is located in Morris Plains, NJ and will not have the ability to be located remotely.

#LI-Onsite

Number of positions available: 1

Shifts available:

Tuesday - Saturday (1:00pm - 9:00pm)

Essential Duties & Responsibilities

- Completes all necessary documentation on GMP documents in real-time.
- Responsible for the accurate recording, review, and storage of laboratory data.
- Strictly follow Data Integrity principles for reliable, accurate, and complete laboratory data.
- Assist with the setup and daily running of the EM laboratory.
- Participate in EM lab operations on a daily, weekly, monthly, and quarterly schedule.
- Incubate and enumerate organisms on cultured media.
- Prepare for shipping to a contract lab, Out-of-Specification (OOS) media plates, and finished product.
- Provide EM support including review, tracking of EM of Microbiology results in support of product release. In conjunction with Quality Assurance, ensure timely completion and data review of routine and batch-related microbiological data.
- Contributes to the gowning qualification program by taking samples and analyzing data.
- Identify process improvements.
- Ensure proper equipment function, calibration, maintenance, and troubleshooting of laboratory equipment.
- Participate in hazardous waste training. Transfer of hazardous waste between lab and trash accumulation area/storage.
- Seeks continuous improvement within EM related activities.
- Supports the QC Micro Management in projects relating to EM technician tasks.
- Follow best practices and regulatory requirements.
- Follows Standard Operating Procedures designed to ensure quality in EM tasks.
- Perform monthly review of laboratory equipment logbooks and perform monthly laboratory cleaning.
- Perform quarterly equipment cleaning.
- Perform additional duties as required and assigned by the Quality Control Management, such as but not limited to the following:
- Receive and inspect incoming shipments of GMP materials and equipment following established procedures and Certificate of Analysis.
- Ensure QC Micro department supplies are inventoried and ordered to avoid stock-out.

Knowledge, Skills & Abilities

- Must be goal-oriented, quality-conscientious, and customer-focused.
- Knowledge of laboratory science and aseptic techniques and principles.
- Effective oral and written communication skills.
- Ability to read, understand, and follow SOPs, work instructions and laboratory test methods.
- Ability to work independently and cooperatively on a team.

Core Values

- Consistently operate with the highest standards of ethics and compliance.

- Take ownership of your actions, success and setbacks.

Ideal Background:

Education & Experience

- Associates degree or Bachelor ' s in Microbiology or closely related field is strongly preferred, or equivalent combination of education and experience.
- A minimum experience of 1 year in the pharmaceutical and biopharmaceutical industry is preferred.
- Knowledge of LIMS preferred.
- Knowledge and understanding of cGMPs and understanding of GLPs used in the industry preferred.
- Detail oriented with expertise in problem solving and solid decision-making abilities.
- Strong interpersonal skills which include a professional demeanor when interacting with Novartis associates.

Languages:

Fluent in English.

Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between \$45,300 and \$84,100 per year. 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: Read our handbook to learn about all the ways we ' ll help you thrive personally

and professionally: <https://www.novartis.com/careers/benefits-rewards>

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.

部门

Operations

Business Unit

Quality

地点

USA

状态

New Jersey

站点

Morris Plains

Company / Legal Entity

U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area
Quality

Job Type
Full time

Employment Type
Regular

Shift Work
No

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