QC Chemist

Job ID
REQ-10060923

8月 29, 2025

USA

摘要

The QC Chemist will support all technical aspects related to quality control testing readiness, including QC reagents and materials management, equipment preparation and daily cleaning and maintenance activities, sample management and QC testing, and documentation completion and review in full compliance with GMP regulation, procedures, and product specifications.

Location: Carlsbad, CA
#LI-Onsite

Shift: This position involves shift work which will be defined through site start up and commercialization readiness, and likely will include weekend and on-call shifts. Apply only if you are available for weekend work.

About the Role

Key Responsibilities:

- Finished Product testing, Environmental Monitoring and Sterility QC testing, and reporting of the QC results.
- Escalation in case of non-conformances and deviations and manage these quality incidents as per procedures.
- Support deviation investigations, OOS/OOT/OOE investigations, CAPA follow up and implementation, and Change Control management, including procedure and form revisions.
- Participation in assigned qualification/validation activities, as necessary.
- Responsible for successful on time completion of required training curricula comprising of the necessary Standard Operating Procedures (SOPs) and Aseptic Techniques, Gowning Qualifications, Testing and specifications, and other relevant training including HSE for the specific role.
- Prepares applicable documents, forms, and records such as analytical batch records and follows Good Documentation Practices.
- Support internal and external Audits and Inspections, as required.

Essential Requirements:

- Education: Bachelors' degree required in relevant Scientific discipline (e.g Chemistry, Microbiology).
- Minimum of 3-year experience in cGMP or aseptic environment required.
- Knowledge of cGMP regulations and FDA guidance applicable to Quality Control for product and Environmental Monitoring testing, as well as Aseptic techniques.
- Practical experience with Microbiology method verification and routine testing practices, EM Monitoring and basic knowledge of method/equipment validation principle and methodologies.
- Ability to interpret analytical data and convert into technical documentation.
- Basic knowledge and understanding of aseptic principles and techniques

Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between $89,600 and $166,400/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

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: https://talentnetwork.novartis.com/network

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
Innovative Medicines

地点
USA

状态

California

站点
Carlsbad

Company / Legal Entity
U469 (FCRS = US469) AAA USA Inc.

Functional Area
Quality

Job Type
Full time

Employment Type
Regular

Shift Work
No

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