

# **Senior QC Analytical Chemist**

Job ID REQ-10053058 Jun 03, 2025 USA

# **Summary**

The Senior QC Analytical Chemist is responsible for performing tasks associated with release testing and reviewing laboratory data. Communicating with and supporting internal & external partners of the Quality Control organization. Supports site as technical expert in related field.

Location: Indianapolis, IN #LI-Onsite

Shift: Friday-Tuesday, 2nd (4 PM to 12:30 AM)

- This position may involve shift work which will be defined through site commercialization needs.
- This position may involve on-call shifts, if required, when scheduled.

#### **About the Role**

#### **Key Responsibilities:**

- Provide support to peers within the Quality Assurance, Quality Control and AS&T teams.
- On-time and GMP-compliant release of patient batches
- Support Quality Control and AS&T as a valued business partner, with a culture of safety, quality, delivery to patients, cost, compliance, and data integrity.
- Author, review and support procedures, investigations, corrective and preventive actions, change controls, complaints, and training as it relates to quality control testing.
- Ensure that QC testing is properly conducted and documented for all performed activities, with emphasis on Data Integrity. Evaluate and approve QC records as required.
- Provide oversight and monitoring of quality control KPIs and programs.
- Perform QC related validations, transfers, improvements, investigations related activities (deviations, OOS, OOE, OOT, CAPAs, trending), and Change Control systems.
- Prepare and participate in health authorities' inspections and internal audits of QC. Ensure quality control area is inspection ready.

## **Essential Requirements:**

- BSc in Chemistry or relevant scientific discipline
- 5+ years of experience in a GMP quality control environment
- General HSE Knowledge
- Knowledge of GMP Manufacturing Process Execution
- Quality Control (QC) Testing
- · Quality Control Sampling

The salary for this position is expected to range between \$85,400 and \$158,600 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? <a href="https://www.novartis.com/about/strategy/people-and-culture">https://www.novartis.com/about/strategy/people-and-culture</a>

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**Benefits and Rewards:** Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <a href="https://www.novartis.com/careers/benefits-rewards">https://www.novartis.com/careers/benefits-rewards</a>

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Division
Operations
Business Unit
Innovative Medicines
Location
USA
State

Indiana

Site

Indianapolis

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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REQ-10053058

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