# **U** NOVARTIS

# **QC** Chemist

Job ID REQ-10031005 Nov 29, 2024 USA

## Sommario

Location: Indianapolis, IN #LI-Onsite

About this role:

The QC 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 is also required.

# About the Role

#### Major accountabilities:

- Provide support to peers within the Quality Assurance, Quality Control and AS&T teams.
- On-time and GMP-compliant release of dosage forms
- 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
- 3+ years of experience in a GMP quality control environment
- Strong HPLC experience a must
- General HSE Knowledge
- Knowledge of GMP Manufacturing Process Execution
- Quality Control (QC) Testing
- Quality Control Sampling
- Dealing With Ambiguity.
- GMP Procedures

- Quality Control (Qc) Testing.
- Quality Standards.
- Self Awareness.
- Technological Expertise.
- Technological Intelligence.

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# **QC Chemist**

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