NOVARTIS

QC Analyst

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REQ-10091367
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Италия

Сводка

The QC Analyst performs analytical testing and related laboratory activities on raw materials, drug substances, drug products, water and primary packaging, in compliance with GMP requirements, approved procedures and Data Integrity principles. The role supports routine Quality Control activities as well as stability, validation, qualification and technology transfer activities within the assigned area of expertise.

About the Role

The QC Analyst is responsible for performing analytical testing and laboratory activities according to approved methods, internal procedures and GMP requirements.

Main responsibilities include:

- Perform analytical testing on excipients, drug substances, drug products, water, stability samples and primary packaging.
- Execute analytical activities related to stability studies, process validation, method validation/verification and technology transfer.
- Perform radionuclidic purity testing on products from AAA sites, as applicable.
- Perform instrument performance qualification activities, as required.
- Record and document analytical activities accurately, ensuring compliance with ALCOA+ principles, Data Integrity requirements and applicable SOPs.
- Know and apply the procedures and GMP requirements relevant to the assigned activities.
- Contribute to the achievement of Quality Control department objectives and agreed priorities.
- Support the maintenance and update of Quality Control SOPs and laboratory documentation.
- Contribute to maintaining laboratories, instruments and the working environment in appropriate operating and housekeeping conditions.

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Дивизион
Operations
Business Unit
Quality
Место
Италия
Сайт
Saluggia
Company / Legal Entity
IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl
Functional Area
Quality
Job Type
Full time
Employment Type
Temporaneo (tempo determinato)
Shift Work
No

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