NOVARTIS

# Quality Control Analyst

Job ID
REQ-10092903
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Италия

### Сводка

The TRD QC Analyst supports the start-up of new laboratories by implementing analytical capabilities, qualifying equipment and systems, and supporting method validation and transfer activities. After start-up, the role performs analytical testing for method validation and routine QC, ensuring compliance with cGMP, GLP, and Novartis quality standards. The position also supports stability studies, investigations, equipment qualification, and continuous improvement initiatives. During routine operations, the role may require shift-based work according to manufacturing and testing needs.

#LI-Onsite
Location: Ivrea, Italy

### About the Role

The TRD QC Analyst plays a key role in establishing and operating Quality Control laboratories that support pharmaceutical development and manufacturing activities. During the laboratory start-up phase, the position supports the implementation of analytical capabilities, laboratory systems, equipment qualification, and method transfer activities. Once operations are established, the role is responsible for executing analytical testing, supporting method validation and verification, and ensuring compliance with cGMP, GLP, and internal quality standards. The TRD QC Analyst also serves as a Super User for laboratory software platforms, providing technical expertise, troubleshooting support, and contributing to system optimization and continuous improvement initiatives.

**Main Activities**:

- Collaborate on the validation / verification / transfer / troubleshooting.
- Support the qualification of new equipment according to Novartis and regulatory standards.
- Execute the required analysis in the QC department based on the plans and needs.
- Contribute on investigations of deviations and out of specifications.
- Keep the laboratory cleanliness and order and actively collaborate to maintain the laboratory and related instruments and auxiliaries in the best possible status.
- Support the safety objective of laboratory personnel and propose improvements to avoid accidents.
- Keep the analytical material stock with the aim of guaranteeing business continuity.
- Ensure proper compliance with Good Laboratory Practices and documentation in the inspection tests used.
- Identify problems, needs and areas for improvement of the laboratory. In accordance with this, design, develop and implement projects that allow improvements in quality, safety and performance of the analysis process in the laboratory.
- Ensure that the work is carried out in accordance with the requirements of the Novartis QMS and in compliance with GMP regulations.
- Oversight of external laboratories activities
- Support for inspections preparation.
- Review of SOPs

**Education**:

- Degree in Pharmacy, Biology, Chemistry, or equivalent

**Languages:**

- Proficiency in Italian
- Good understanding (mandatory), fluent in verbal and written desirable.

**Experience/Professional requirement:**

- Open and clear collaboration and communication
- Shows the appropriate sense of urgency around given tasks
- Scientific, technical and regulatory knowledge in a specific area.
- Knowledge of cGMP, working knowledge of safety and environmental regulations and guidelines.
- At least 2 years of experience in pharmaceutical companies (or analogue) in equivalent roles.
- Proficiency in the use of laboratory equipment such as pipettes, balances, pH meters, and HPLC systems.
- Preferred knowledge of Chromeleon and GLIMS software.

**Benefits & Rewards**

At Novartis, we’re committed to reimagining medicine together - and rewarding the people who make it happen.

**Expected Annual Base Salary Range for role: € 30,200 to € 56,000**

The base salary offered is determined based on gender-neutral objectives, such as relevant skills, competencies and experience in accordance with the Novartis pay setting policy and upon joining Novartis will be reviewed periodically.

In addition to your base salary, you may be eligible for a performance-based bonus depending on certain performance parameters.

The rewards of being part of our team go far beyond base pay and incentives. We also offer a variety of competitive benefits in kind to help you thrive personally and professionally, such as insurance plans, retirement plans, wellbeing resources and global recognition programs. In addition, we provide flexible and hybrid working options, where possible, and minimum 14 weeks paid parental leave.

**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:** Learn about all the ways we’ll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

Primary location salary range
€30,200.00 - €56,000.00
Дивизион
Development
Business Unit
Development
Место
Италия
Сайт
Ivrea
Company / Legal Entity
IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regolare
Shift Work
No

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