NOVARTIS

TRD QC Senior Expert (m/f/d)

Job ID
REQ-10090995
Ноя. 09, 2026
Италия

Сводка

Location: Ivrea, Italy #onsite

Role Purpose:
The TRD QC Senior Expert is a key technical and scientific role responsible for supporting the start-up and operation of the Pilot Plant QC laboratories. During the build-up phase, the role focuses on laboratory setup, instrument qualification, system implementation, SOP development, and validation activities to ensure GMP compliance and inspection readiness. Once operational, the position leads analytical activities related to technology transfer, method validation, and QC testing, while providing technical leadership, coordinating stakeholders, supporting investigations (OOS, deviations, CAPAs), and ensuring reliable, compliant, and timely analytical results. The role also contributes to continuous improvement, audit readiness, and the development of laboratory expertise.

About the Role

Main activities

- Support activities during the laboratory build-up phase for the start-up of the new Pilot Plant QC laboratories.
- Support and coordinate laboratory activities associated with technology transfer projects, including implementation of analytical methods, specifications, reference standards, and testing strategies.
- Act as the primary technical point of contact for cross-functional teams.
- Provide scientific and technical expertise for analytical methods, supporting method qualification, verification, troubleshooting, and lifecycle management.
- Evaluate analytical data, method performance, and technical risks, ensuring reliable, reproducible, and scientifically sound analytical results.
- Oversee routine Quality Control laboratory operations, ensuring effective planning, prioritization, and allocation of resources to meet business and project requirements.
- Support the management of deviations, OOX investigations, and CAPAs, performing technical assessments and contributing to root cause analysis and implementation of corrective and preventive actions.
- The role is responsible for the review and approval of analytical results supporting the release of finished products and the testing of raw materials
- Ensure compliance with cGMP, GLP, Data Integrity principles, pharmacopeial requirements, internal procedures, and regulatory expectations.
- Contribute to the qualification and lifecycle management of laboratory equipment, computerized systems, analytical standards, and reagents.
- Support audit and inspection readiness through accurate documentation, effective data review, and maintenance of a compliant laboratory environment.
- Provide technical coaching and mentoring to laboratory personnel.

Additional specific roles/tasks

- Act as point of contact for escalations from analysts and for cross-functional interfaces (QA, Manufacturing, TRD).
- Support change control activities impacting QC.
- Contribute to updates of SOPs/WIs and to the implementation of new tools or digitalization initiatives.
- Track and report relevant KPIs.
- Participate in internal audits, self-inspections, and regulatory inspection preparation activities

Ideal Background

Education (minimum/desirable):

Master's Degree in Pharmacy, Biology, Chemistry, or equivalent

Languages:

- Proficiency in Italian (Local Language)
- Fluency in spoken and written English

Experience/Professional requirement:

- Open and clear collaboration and communication

- Shows the appropriate sense of urgency around given tasks
- Scientific, technical and regulatory knowledge in the specific area.
- Strong knowledge of GMP regulations, analytical technologies, laboratory equipment qualification, and data integrity principles
- At least 5 years of experience in pharmaceutical companies in equivalent roles.
- Previous experience in radiopharmaceutical companies is preferred.
- Good organizational skills.
- Knowledge of computerized systems such as Chromeleon, GLIMS, and SAP.
- Good knowledge of analytical instrumentation, including HPLC, FT-IR, TLC, and UV spectroscopy.

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: €45,300 to € 84,100

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
€45,300.00 - €84,100.00
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
Regular
Shift Work
No

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