NOVARTIS

Quality Control Analyst (Temporary Position)

Job ID
REQ-10078372
май 20, 2026
Испания

Сводка

#LI-Onsite
Location: Zaragoza – La Almunia, Spain

Relocation Support: This role is based in Zaragoza – La Almunia, Spain. Novartis is unable to offer relocation support: please only apply if accessible.

Bring quality to life in a role where your expertise safeguards patient health. As a Quality Control Analyst, you will ensure products meet the highest standards of safety and compliance while working in a dynamic laboratory environment.

This is a temporary opportunity where your analytical skills will directly support the release and stability of pharmaceutical products, contributing to Novartis' mission to reimagine medicine, with availability required to support a 24/7 operation.

This role requires availability to work rotating shifts, including mornings, afternoons, and nights.

About the Role

Key Responsibilities

- Perform analytical testing of drug products, finished goods, and materials following approved procedures and standards
- Document laboratory activities and results accurately in line with Good Manufacturing Practice requirements
- Manage sample storage, tracking, and handling to ensure integrity and compliance
- Conduct stability testing and maintain proper documentation of stability studies
- Investigate and report technical complaints, adverse events, and quality issues within 24 hours
- Ensure full adherence to Standard Operating Procedures and regulatory guidelines at all times
- Support audit and inspection readiness through consistent compliance and documentation practices
- Monitor deadlines and ensure timely completion of all testing and reporting activities
- Identify and implement process improvements to enhance efficiency and reduce operational costs
- Collaborate with cross-functional teams to maintain quality standards and resolve issues effectively

Essential Requirements

- Degree in Chemistry, Pharmacy, or a related scientific field
- Experience in quality control within a pharmaceutical or regulated manufacturing environment
- Knowledge of Good Manufacturing Practice and quality standards
- Hands-on experience with analytical laboratory techniques and equipment
- Ability to follow Standard Operating Procedures with high attention to detail
- Strong problem-solving skills and ability to manage competing priorities
- Intermediate level of English, with good reading and writing skills

Desirable Requirements

- Experience with stability testing and lifecycle management of pharmaceutical products
- Familiarity with audit and inspection processes within regulated environments

Commitment to Diversity and Inclusion:
Novartis is committed to building an outstanding, inclusive work environment and diverse teams representative of the patients and communities we serve

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)

Дивизион
Operations
Business Unit
Quality
Место
Испания
Сайт
Zaragoza
Company / Legal Entity
ES45 (FCRS = ES045) Advanced Accelerator Applications Iberica S.L.U.
Functional Area
Quality

Job Type
Full time
Employment Type
Temporary (Fixed Term)
Shift Work
No

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