Pilot Plant Quality Control Associate Expert

Job ID
REQ-10063927

10月 06, 2025

Italy

摘要

This role utilizes chemistry laboratory skills to test and measure product or materials while ensuring that analysis is performed according to established Standard Operating Procedures (SOPs), Analytical Methods & current Compendia.

About the Role

Key responsibilities:

- Executes analyses on excipients, raw materials, finished products, and packaging per internal SOPs and applicable pharmacopeias.
- Conducts stability studies, process and method validations, and activities related to tech transfer of pharmaceutical products.
- Performs instrument qualification and re-qualification (IQ/OQ/PQ) according to internal procedures.

- Records analytical data in compliance with ALCOA+ principles and data integrity requirements.
- Reviews GMP documentation, batch records, analytical reports, and certificates for completeness and compliance.
- Collaborates with QA, Production, and R&D; supports method transfers, troubleshooting, and safe lab/5S practices.
- Participates in audits and inspections by providing data and responses; contributes to CAPA, deviations, change control, and SOP updates.
- Tracks KPIs: on-time completion and right-first-time rates, data integrity compliance, audit outcomes, documentation quality, and continuous improvement impact.

Essential Requirements:

- Degree in Chemistry
- Solid technical-scientific knowledge of pharmaceutical/chemical QC or equivalent analysis
- Previous experience working in a laboratory environment in the pharmaceutical industry
- Experience in GMP environment
- Fluent in Italian and English

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部门
Development

Business Unit
Innovative Medicines

地点
Italy

站点
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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