

# Pilot Plant Quality Control Associate Expert

Job ID REQ-10068760

12月 16, 2025

Italy

### 摘要

Location: Ivrea, Italy #onsite

#### Role Purpose:

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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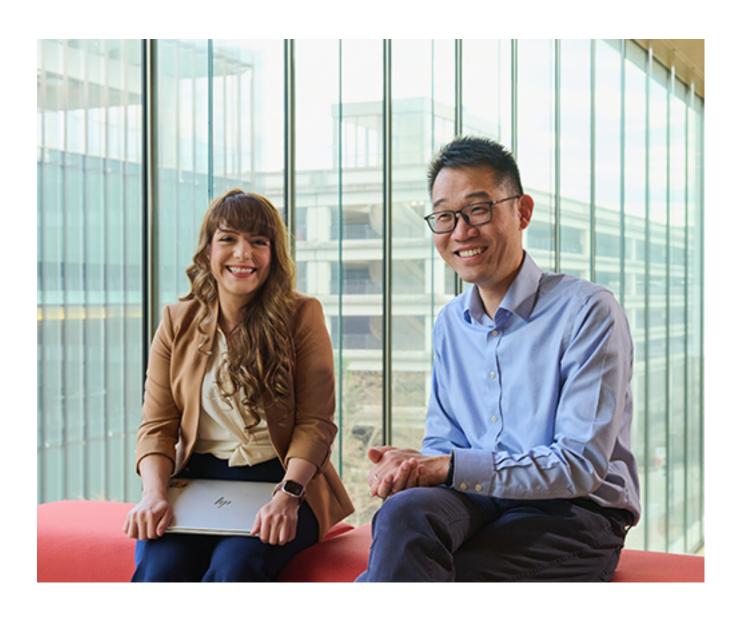
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部门 Development
Business Unit Development
地点 Italy
站点 Ivrea
Company / Legal Entity IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl
Functional Area Research & Development
Job Type Full time
Employment Type Regolare
Shift Work No

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