

## QC Analyst - tempo determinato

Job ID  
REQ-10050694

May 15, 2025

Italy

### Summary

Questo ruolo utilizza le competenze di laboratorio di chimica per testare e misurare prodotti o materiali, garantendo al contempo che l'analisi venga eseguita secondo le procedure operative standard (SOP), i metodi analitici e gli attuali compendi.

### About the Role

#### Key Responsibilities:

- Execute analyses on excipients, raw materials, pharmaceutical products, water, and primary and secondary packaging according to internal SOPs
- Perform stability studies, process validation protocols, method validation protocols and studies related to pharmaceutical products involved in technological transfer for the relevant sector

- Conduct qualifying tests for instrument performances • Properly record analyses in compliance with ALCOA+ principles
- Collaborate to achieve departmental goals
- Know and apply the relevant SOPs for the responsible sector
- Contribute to maintaining good conditions of laboratories, instruments and working environment
- Assist in keeping the Quality Control SOPs updated
- Review of GMP documentation

#### Essential Requirements:

- Diploma/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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部門

Operations

部門

Innovative Medicines

国

Italy

勤務地

Saluggia

Company / Legal Entity

IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl

Functional Area

Quality

職種

Full time

雇用形態

Temporaneo (tempo determinato)

Shift Work

No

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