

Qualified Person

Job ID

REQ-10073536

Apr 07, 2026

LOC_IT

About the Role

Major accountabilities:

- Guarantee and certify that each batch of medicines is produced and checked in compliance with the law and the conditions imposed in the marketing authorization.
- Assessment and release of manufactured medicinal products, in accordance with national legislation.
- Guarantee that the documentation attesting the suitability of each product lot is available and can be shown at the request of the health authority.
- Collaborate in the approval of deviation investigations.
- Make sure that the batch record of the released batch is stored correctly and can be exhibited at the request of the health authority.
- Communicate immediately to the national Health Authority (AIFA) and to the Management any substantial irregularity detected in the product that has already been placed on the market.
- Work in collaboration with Quality Control and Production departments in the activities related to the manufactured batches.
- Identify and propose technological and organizational interventions aimed at improving manufacturing processes in terms of quality, productivity and costs and the optimization of resources.
- Collaborate with the Function Managers in order to guarantee the correctness of the Quality Management System.
- Management of deviations, complaints, change control and CAPA.

Essential requirements:

- Degree in Pharmacy, CTF or Chemistry.
- Previous experience in the role within a pharmaceutical sterile manufacturing environment (Authorized Qualified Person certificate according to Legislative Decree n. 219 of April 24th, 2006).
- Strong affinity with quality and awareness of quality issues.
- Open and clear collaboration and communication to make sure the daily production operation runs smoothly and safely.
- Fluent in Italian and English.

This role is 100% site based in Ivrea.

Role Requirements

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>

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Division

DIV_TO

Business Unit

Quality

Location
LOC_IT
Site
Ivrea
Company / Legal Entity
IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl
Functional Area
FCT_QA
Job Type
Full time
Employment Type
Regolare
Shift Work
No
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