NOVARTIS

Quality Assurance Specialist

Job ID
REQ-10080312
Jun 10, 2026
LOC_IT

About the Role

Key responsibilities:

- Drive technological and organizational improvements to enhance manufacturing quality, productivity, cost efficiency, and resource optimization.
- Ensure full GMP compliance across operations, aligning site practices with company policies, standards, and regulatory requirements.
- Support development and lifecycle management of new products, process improvements, and existing product enhancements.
- Review and approve GMP documentation (e.g., batch records, specifications, protocols) ensuring accuracy, traceability, and compliance.
- Oversee document lifecycle management (paper and electronic), ensuring proper control, data integrity, and archival.
- Manage quality systems activities including deviations, CAPAs, Change Controls, complaints, and laboratory investigations (OOS/OOT/OOE).
- Lead or support audits, inspections, self-inspections, training compliance, supplier qualification, and Site Master File/APQR preparation.
- Contribute to continuous GMP governance and regulatory expertise development, supporting readiness for future QP responsibilities.

Essential requirements:

- Scientific degree.
- 2/3+ experience in a similar role within a sterile production environment.
- Good knowledge of GMP.
- Fluent in Italian. Good knowledge of English.

Why Novartis? Our purpose is to reimagine medicine to improve and extend people’s lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: https://www.novartis.com/about/strategy/people-and-culture

At Novartis we’re committed to reimagining medicine together – and rewarding the people who make it happen.

Expected annual Salary Base Range for the role: EUR 29.800,00 - 55.300,00

Benefits and rewards: Read our handbook to learn about all the ways we’ll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

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.

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

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

Division
DIV_TO
Business Unit
Quality
Location
LOC_IT
Site
Saluggia
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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List of links present in page

1. https://prod1.jobapi.novartis.com.cn/req-10080312-quality-assurance-specialist-it-it
2. https://www.novartis.com/about/strategy/people-and-culture
3. https://www.novartis.com/careers/benefits-rewards
4. https://www.novartis.com/about/strategy/people-and-culture
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7. https://novartis.wd3.myworkdayjobs.com/it-IT/Novartis_Careers/job/Saluggia/Quality-Assurance-Specialist_REQ-10080312-1