

# **Quality Assurance Specialist**

Job ID REQ-10016971 Oct 23, 2024 Italia

#### Resumen

-Manages Quality aspects and projects within area of responsibility. -Ensures and supports overall GxP conformity and Compliance with the Novartis Quality Management Systems. Prepare Documentation for Batch Release of commercial Pharmaceuticals

#### **About the Role**

# Major accountabilities:

Oversight of all production and testing activities, ensures compliance with cGxP, incl. data integrity and
eCompliance -Support exception investigations -Review and approval of production, QC, and AS and T
records -MBR review -Support OpEx improvement projects Qualified Person – Executes batch release in
compliance with registration -Reporting of technical complaints / adverse events / special case scenarios
related to Novartis products within 24 hours of receipt -Distribution of marketing samples (where
applicable)

# Key performance indicators:

 On-time and GMP-compliant release of dosage forms -No complaints about inspections by authorities in your own area of responsibility without these being noticed and communicated beforehand -Successfully support continuous improvement projects -Executes batch release in compliance with registration

#### **Minimum Requirements:**

# Work Experience:

- Functional Breadth.
- QC/ QA in pharmaceutical ind./ biotech with environmental monitoring &.
- Collaborating across boundaries.
- cleanliness zones.

#### Skills:

- Continuous Learning.
- Dealing With Ambiguity.
- Gmp Procedures.
- Qa (Quality Assurance).
- Quality Control (Qc) Testing.
- · Quality Standards.
- Self Awareness.

- Technological Expertise.
- Technological Intelligence.

#### Languages:

• English.

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División

Operations

**Business Unit** 

Innovative Medicines

Ubicación

Italia

Sitio

Milano

Company / Legal Entity

IT08 (FCRS = IT008) Novartis Farma S.p.A.

Alternative Location 1

Remote, Suiza

**Functional Area** 

Quality

Job Type

Full time

**Employment Type** 

Temporary (Fixed Term)

Shift Work

No

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