

Quality Assurance Specialist

Job ID
REQ-10016971
Ott 23, 2024
Italia

Sommario

-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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Divisione

Operations

Business Unit

Innovative Medicines

Posizione

Italia

Sito

Milano

Company / Legal Entity

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

Alternative Location 1

Remote, Svizzera

Functional Area

Quality

Job Type

Full time

Employment Type

Temporary (Fixed Term)

Shift Work

No

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