

Change Control Specialist

Job ID
REQ-10066615

11月 13, 2025

Mexico

摘要

-Manages Quality aspects and projects within area of responsibility. -Ensures and supports overall GxP conformity and Compliance with the Novartis Quality Management Systems.

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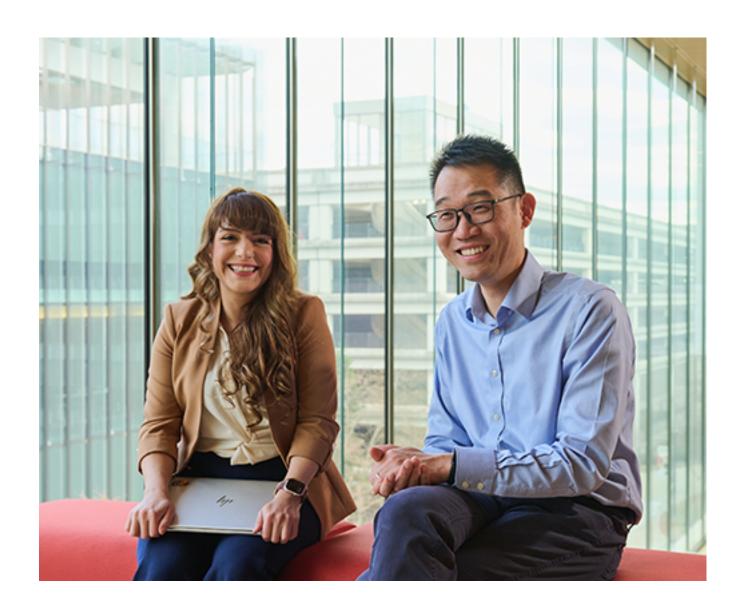
Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

部门 Operations **Business Unit** Quality 地点 Mexico 站点 **INSURGENTES** Company / Legal Entity MX06 (FCRS = MX006) Novartis Farmac é utica S.A. de C.V. **Functional Area** Quality Job Type Full time **Employment Type** Regular Shift Work No Apply to Job

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