QA Operations Specialist

Job ID REQ-10029818 Nov 18, 2024 China

About the Role

Key Responsibilities

- Provide shop-floor QA support and oversight of production and testing activities, ensures compliance with cGxP, ISO9001, incl. data integrity and eCompliance
- Batch document review, support product release
- Support deviation and compliant investigations within MU, QC, Engineering and warehouse and other relevant areas.
- Performs routine documentation activities, Review and approval of production and QC documents and records, MBR review
- Supports change control activities
- IPC oversight
- Support OpEx improvement projects
- Support internal and external audit or inspection activities
- Comply with all HSE guidelines. Detect and report potential accident, risks and propose solutions. Participate in HSE risk assessments. Preparation and participation to internal HSE audits
- · Responsible for participating in initial training and retraining

Essential Requirements:

- University degree with a scientific / technological background (e.g. Chemistry, Pharmacy, Biology, Biochemistry, or equivalent)
- Qualified Person (depending on local requirements)
- Good (oral and written) in English; fluent in local language (oral and written)
- More than 3 years of experience in pharmaceutical quality control, quality assurance or production
- Operations Management and Execution; Functional Breadth; Collaborating across boundaries; Applied Practice

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Role Requirements

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Operations
Business Unit
Innovative Medicines
Location
China
Site
Changping County (Beijing)
Company / Legal Entity
CN06 (FCRS = CN006) Beijing Novartis Pharma Co., Ltd
Functional Area
Quality
Job Type
Full time
Employment Type

Job ID

Nο

Regular Shift Work

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QA Operations Specialist

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