Expert Science & Technology (Oral Solid Dosage forms)

Job ID
REQ-10052839

5月 27, 2025

India

摘要

Perform and document scientific experiments in the laboratory for drug substances (DS) and drug products (DP) in collaboration with multifunctional project teams. Contribute to maintenance of lab instruments/day-to-day operations. Timely execution of project related activities to support TRD-NCE strategies and goals.

About the Role

Major accountabilities:

- Plan, organize, execute, and document scientific experiments (e.g., analytical method developments/ validations/ transfers/ stability/ release testing, formulation development analytics etc.) according to the agreed timelines and appropriate quality standards.
- Accountable for documentation and submission of raw data in appropriate data

system (for e.g., LIMS test activation and results entry).
- Responsible for good documentation practices (GDP) and good laboratory practices (GLP) during execution of laboratory activities.
- Support in evaluation and interpretation of results including investigations on SST failures, OOX/Deviations/Change controls as needed.
- Responsible for assigned laboratory related area/activities (e.g., chemical/reagents/consumables/samples/column/ glassware management etc.).
- Responsible for implementation and maintenance of lean/efficient/environmentally sustainable practices in the laboratory.
- Proactively communicate key issues and any other critical topics in a timely manner to the manager and/or to any other relevant project team member(s).
- Responsible to meet KQI (Key quality indicators) and KPI (Key performance indicators) for all assigned activities.
- Support internal and external audits and ensure no critical findings within the assigned scope.
- Actively contribute to team and organization goals.
- Work according to appropriate SOPs, GMP, GLP, QM, HSE, ISRM & Novartis Guidelines.
- Additional specific roles/tasks: See Up4Growth training assignments for the business roles for the associate as per the team matrix and completion of trainings in transcript of learning system (e.g., Up4Growth).

Minimum Requirements:

- Masters in Life Science (e.g., analytical / organic chemistry /pharmacy / pharmaceutical development) or equivalent.
- 5+ years of relevant work experience in OSD forms- hands on in chromatography, multimedia dissolutions, In-vivo & Invitro dissolutions, quality investigations, QBD etc.
- Fluent in English (oral and written).Knowledge of site language, if required.
- Knowledge in quality principles driving drug development such as GMP.
- Understanding of general regulatory and quality expectations.
- Good scientific background, communication skills including presentation and scientific/technical writing.

Work Experience:

- Functional Breadth.
- Operations Management and Execution.
- Collaborating across boundaries.

Skills:

- Environment.
- Experiments Design.
- Health And Safety (EHS).
- Laboratory Equipment.
- Manufacturing Process.
- Materials Science.
- Process Simulation.

- Project Management.
- Sop (Standard Operating Procedure).
- Technical Writing.

Languages :

- English.

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部门
Development

Business Unit
Innovative Medicines

地点
India

站点
Hyderabad (Office)

Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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