

## Expert Science & Technology (Oral Solid Dosage forms)

Job ID  
REQ-10052839

8月 29, 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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Universal Hierarchy Node

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India

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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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