NOVARTIS

Expert, Science & Technology

Job ID
REQ-10090578
Oct 27, 2026
LOC_IN

About the Role

Key Responsibilities

- Responsible for performing method feasibilities and validation of robust analytical methodologies applied to innovative oral solid dosage forms. Strong experience in chromatography techniques and experience in handling techniques like Dissolution, Titrator etc., used for OSD testing is a pre-requisite.
- 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 an 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.
- Generate and 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.

Essential Requirements:

- Strong expertise in Chromatography and Dissolution Testing, including method selection and handling
- Proven experience in Analytical Method Development, Validation, and Transfer and multimedia dissolution
- Expertise in method validation, method transfer, IcH guideline, HPLC and UPLC.
- Solid understanding of Regulatory Compliance and GLP standards
- M.Pharm./M.Sc. with 4 -8 years of relevant experience, with demonstrated experience in a related domain.

Desirable Requirements:

- Understanding digital tools
- Hands on experience with GLIMS, Chromeleon and ELN

Role Requirements

Division
DIV_GD
Business Unit
Development
Location
LOC_IN
Site

Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Functional Area
FCT_RD
Job Type
Full time
Employment Type
Regular
Shift Work
No
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