Expert- Science and Technology

Job ID
REQ-10057517

7月 14, 2025

India

摘要

Design, plan, perform, interpret and report results of scientific experiments for the method development of drug substances (DS) and drug products (DP) within global ARD. Lead and manage all project/local network activities, support/coach team members, participate in sub-teams and contribute to overall TRD strategies and goals.

About the Role

Major activities

· Develop and qualify various analytical methods (e.g., fast LC, titration, dissolution).

· Provide analytical and technical support to PHAD/project team at various stages of product development (e.g., CSF, FMI and LCM).

· Design and author analytical documents (e.g., Analytical methods, Stability protocols/reports, Excipient compatibility (EC) protocol/reports; APS protocols/reports, etc.)

· Support Analytical project leader for setting analytical development strategy.

· Support in data interpretation, results compilations and sharing the information with critical observations and proposals to project team.

· Responsible for project related sample handling (e.g., sampling plans, issuance, storage, distribution, reconciliation/destruction of the samples).

· Support planning for assigned project activities. Accountable to meet KQI (Key quality indicators) and KPI (Key performance indicators) for all assigned project activities.

· Provide requests for lab activities to the associates and stakeholders.

· Manage project activities including logistics at third parties and external testing laboratories.

· Proactively communicate key issues and any other critical topics in a timely manner to the appropriate management level and/or to any other relevant project team member(s).

· Work according to appropriate SOPs, GMP, GLP, QM, HSE, ISEC & Novartis Guidelines.

· Subway: Author (EC/APS protocol and reports), review of test methods and compatibility study plan.

· ESOPS: Read SOP access and Review of SOPs.

Ideal background

Education and experience:

M.Sc. /M. Pharm/ Ph.D. with relevant experience.

Languages:

Good knowledge of English and site language (oral and written).

Professional requirement:

- Recognized expertise in analytical methodology and broad scientific as well as technical and strategic background.
- Demonstrated successful experience with working in interdisciplinary and cross-cultural teams.
- Demonstrated leadership and advanced coaching and mentoring skills.

- Thorough knowledge of relevant SOP, GMP and Novartis regulations and policies if applicable.
- Excellent communication/presentation skills and scientific.

Why Novartis: Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients ' lives. Ready to create a brighter future together? https://www.novartis.com/about/strategy/people-and-culture

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部门
Development

Business Unit
Innovative Medicines

地点
India

站点
Telangana

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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Accessibility and accommodation

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