

## Senior Expert Science & Technology

Job ID  
REQ-10036379

8月 03, 2025

India

### 摘要

400! This is the number of associates in Global Analytical R&D, across 4 countries, working tirelessly on innovative and patient centric medicines. As part of this group, you design, plan and/or perform scientific/technical studies. By bridging the analytical science to the clinical performance you will drive the transformation of our molecules into medicines that improve and extend patient 's lives.

### About the Role

Major Accountabilities:

- Design, plan and perform scientific experiments for projects at different clinical phases of drug substance and drug product with minimal guidance.

- Monitors degradation pathways and shelf-life of products.

Well versed with regulatory guidelines, scientific literature, technology transfer and interpretation of the results to draw conclusions in reports.

- Adhere to Quality metrics, Compliance and Good Documentation Practices following ALCOA+ principles, GLP, OQM, HSE, ISEC and Novartis guidelines.

- Should be a Team player by adding value in collaborating with other teams to support project deliverables within agreed timelines, mentoring new joiners, active participation in project meetings / networks / meetings and contributing to Team goals while meeting individual objectives.

- Responsible for Qualification of instruments / equipment ' s (URS to Report) and periodic calibrations as per applicable site procedures.

Responsible for implementation and maintenance of lean/efficient/environmentally sustainable practices in the laboratory

- Ability to perform investigations, guide team members, communicate proactively and clearly to global stakeholders.

Cross-functional collaboration

What you bring to the role:

- Broad scientific or technical knowledge in a specific area.

- Adequate understanding of development processes in own function.

- Advanced knowledge of laboratory and/or technical tools.

- Good knowledge of software and computer tools.

- Proficient in literature searches

Good understanding about Regulatory Guidelines

Strong problem-solving and critical thinking

Knowledge on ICH guidelines (Q2(R1)), QbD principles etc

Minimum requirements:

- Minimum: M. Pharm./M.Sc. Good knowledge of English (oral and written).

- PhD on technical subject with 4+yrs of relevant experience. or Master of Science with 12+ years of relevant experience in testing of Solid oral dosage forms.

•Good presentation skills and scientific/technical writing skills.

•Good communication skills

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

Universal Hierarchy Node

地点

India

站点

Hyderabad (Office)

Company / Legal Entity

IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Alternative Location 1

Telangana, India

Functional Area  
Research & Development

Job Type  
Full time

Employment Type  
Regular

Shift Work  
No

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Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.



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