

Senior Expert Science & Technology - Process Research & Development

Job ID REQ-10067102

11月 23, 2025

China

摘要

-Design, plan, perform, interpret and report results of scientific experiments for the preparation and timely delivery of drug substances (DS), drug products (DP), processes and procedures. Lead and manage all project/local network activities, support/coach team members, participate in sub-teams and contribute to overall TRD strategies and goals. -Management Track -Lead a team for the development of pharmaceutical/biological/cell-gene therapies working in a multidisciplinary environment. Execute and support developing the functional strategy and drive operational excellence in line with TRD vision and strategy. Ensure full portfolio support in line with GDD, NTO and NIBR plans.

About the Role

Major accountabilities:

- Oversee and lead all activities of assigned teams /projects; meet customer needs.
- Work according to appropriate standards for quality, ethics, health, safety, environment, protection and information security; lead initiatives to ensure continuous improvement; all activities have to be aligned with organizational workflows and procedures.
- Evaluate and interpret results, draw relevant conclusions; supervise project related activities; perform complex tasks without having established procedures.
- Oversess and may also write protocols, scientific reports, lab procedures or process. related SOPs; write scientific documents intended for external partners or for generation of registration documents; interact with authorities -Communicate, address and solve problems within own and broader area of responsibility; communicate effectively across organizational interfaces; lead the transfer of know how to other departments or external contractors, including troubleshooting and on-site training.
- For technical development units: Develop complex methods (lab or plant); lead the
 optimization of project related scientific /technical activities or processes, co-ordinate local
 team(s); guide development and implementation of new technologies. For GMP units: ensure
 compliance to cGMP. For technology focused role: Provide scientific and technical guidance;
 actively foster knowledge exchange.
- Develop, mentor and coach other scientific associates; present scientific /technical results
 internally and contribute to publications, presentations and patents. For project-focused role:
 Lead assigned teams; represent own technical function in teams and fulfill all project tasks
 and responsibilities related to the own discipline -Broadly uses professional concepts in
 accordance with company objectives to solve complex problems in creative and effective
 ways -Contributes to many cost center goals and objectives; may contribute to service line
 goals.
- Develop detailed plans and timelines with the manager, develop formulation strategies and plans for designated projects from development to cGMP manufacture. Ensure accurate, speedy reports are produced to enable reg. Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt. Distribution of marketing samples (where applicable)

Minimum Requirements:

- People Challenges. Managing Crises. Functional Breadth.
- Project Management. Operations Management and Execution.
- Collaborating across boundaries. Coaching Skills.
- Data Science. Environment. Experiments Design.
- Health And Safety (Ehs). Laboratory Equipment. Manufacturing Process. Materials Science.
 Process Simulation.
- Sop (Standard Operating Procedure). Technical Writing.
- English.

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

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: https://talentnetwork.novartis.com/network

Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

部门 Development
Business Unit Development
地点 China
站点 Changshu (Jiangsu Province)
Company / Legal Entity CN23 (FCRS = CN023) Suzhou Novartis Technical Development Co., Ltd.
Functional Area Research & Development
Job Type Full time
Employment Type Regular

Shift Work

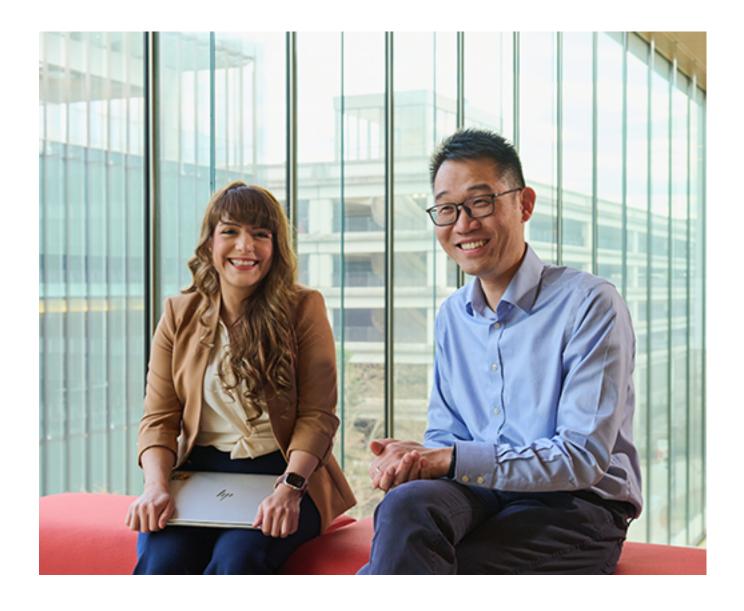
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Accessibility and accommodation

Novartis is committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to perform the essential functions of a position, please send an e-mail to diversityandincl.china@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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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