NOVARTIS

# Senior Expert Science & Technology I

Job ID
REQ-10081103
Jun 22, 2026
Китай
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### Сводка

-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.

### About the Role

**Major accountabilities:**

- 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)

**Key performance indicators:**

- Adherence to costs, quality, quantity, and timelines for all assigned tasks.
- Adherence to Novartis standards, in particular, quality, ethical, health, safety, and environment (HSE), and information security (ISEC) standards.
- Feedback from other team leaders and advisory boards.
- Measurable contributions to the success, efficiency and productivity of the department and new programs/initiatives started and implemented.
- Refer to annual individual and team objective setting.
- Internal and external publications/presentations, invited lectures.
- Successful and effective execution of assigned tasks within given timelines at expected quality; right the first time and on time; demonstrate initiative and strive for high level of quality2.
- Adherence to appropriate standards as defined in Quality Manual, SOPs, ethical, health, safety, environment (HSE), and information security (ISEC) guidelines3.
- Refer to annual individual and team objective setting4.
- Measurable contributions to efficiency increase and productivityScientist: -Adherence to costs, quality, quantity, and timelines for all assigned tasks.
- Adherence to Novartis standards, in particular, quality, ethical, health, safety, and environment (HSE), and information security (ISEC) standards.
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- Refer to annual individual and team objective setting.
- Internal and external publications/presentations, invited lectures.

**Minimum Requirements:**

- Coaching Skills.
- Data Science.
- Environment.
- Experiments Design.
- Health And Safety (Ehs).
- Laboratory Equipment.
- Manufacturing Process.
- Materials Science.
- Process Simulation.
- Project Management.
- 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

**Benefits and Rewards:** Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

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CN23 (FCRS = CN023) Suzhou Novartis Technical Development Co., Ltd.
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No

### 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.

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