NOVARTIS

Technical Manager

Job ID
REQ-10087234
Sep 07, 2026
LOC_CN

About the Role

Major accountabilities:

- Responsible for technology transfer activities at site level (within, inbound and outbound), including any scale-up or other process adaptations.

- Leads technical transfer project team at site and liaises efficiently with involved functions (e.g. Technical Development, Supply Chain, Production Unit, Quality Control, HSE, other sites.).

- Owns the process knowledge of the product(s) assigned throughout the commercial lifecycle, maintains the oversight on process capability, through data trending and statistical analysis of critical variables, ensuring process(es) are robust, in continued state of validation and continuously improving.

- Ensures seamless flow of knowledge and information across functions, and with other Sites when applicable, with focus on the assigned product(s).

- Oversees processes and standards to maintain and improve existing and to implement new innovative manufacturing technologies when applicable.

- Responsible for developing, implementing and managing the site process validation, primary packaging validation, cleaning validation and revalidation strategies to meet cGMP and quality requirements on time and on budget to ensure that programs are aligned with Regulatory Authorities' expectations and related SOPs.

- Design, plan, perform, interpret and report scientific experiments under the lead of the department head to contribute to overall MS&T strategies and objectives.

Scientist MS&T

- Complex projects with deep understanding of development process requirements
- Difficult to solve topics (eg. new by-product investigation, complex deviations)
- Cross product alignment in support approach (larger sites)
- Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt
- Distribution of marketing samples (where applicable)

Education:

- Higher scientific degree in Pharmacy, Pharmaceutical Technology, Chemistry or equivalent scientific field.

Languages:

- Fluency in English languages.
- undefined

Experience:

- Minimum 5 years of experience in pharmaceutical (chemical) manufacturing and in sterile drug product manufacturing , with a comprehensive know-how in pharmaceutical technology

Compentency

- Applied Business Insights
- Operational Excellence
- Project Excellence
- Manufacturing Process/Product Expertise

Key performance indicators:

- Cost, C-Sat and productivity targets
- Achievement of project plans & milestones
- Internal customer satisfaction with quality of services provided

Role Requirements

Division
DIV_TO
Business Unit
Product Supply Chain
Location
LOC_CN
Site
Changshu (Jiangsu Province)
Company / Legal Entity
CN06 (FCRS = CN006) Beijing Novartis Pharma Co., Ltd
Functional Area
FCT_TO
Job Type
Full time
Employment Type
Regular
Shift Work
No
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Technical Manager

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