NOVARTIS

Manufacturing Science & Technology experts

Job ID
REQ-10067047
Dec 05, 2025
LOC_RO

About the Role

Key Responsibilities

- Maintain end-to-end process knowledge and lifecycle oversight for assigned products.
- Lead tech transfers and launch readiness activities, ensuring seamless scale-up and compliance.
- Develop and execute validation strategies (process, cleaning, packaging) aligned with cGMP and regulatory standards.
- Provide technical support, risk mitigation, and cross-functional collaboration to ensure business continuity and quality.

Essential Requirements:

- BSc in Pharmacy, Pharmaceutical Technology, Chemistry, or an equivalent scientific degree; MSc desirable or equivalent experience.
- Fluent in English (oral and written).
- 2–3 years of experience in manufacturing, manufacturing science and technology (MS&T), technical development, or quality—ideally within GMP/GxP-regulated environments.
- Experience in GMP-regulated pharmaceutical manufacturing and MS&T.
- Strong skills in process validation, CPV, and statistical data analysis.
- Proven leadership in tech transfer and cross-functional project management.
- Excellent documentation and compliance mindset.

Commitment to Diversity and Inclusion / EEO paragraph:

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

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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Role Requirements

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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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Targu Mures
Functional Area
FCT_TO
Employment Type
Regular
Shift Work
No
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