Process Expert (m/f/d), Schaftenau, Tyrol - temporary

Job ID
REQ-10061560

9月 11, 2025

Austria

摘要

Shift Lead II
Der Shift Leader ist verantwortlich f ü r die Verwaltung seines Teams, um die Fertigung nach Zeitplan in Ü bereinstimmung mit den HSE- und GMP-Regeln durchzuf ü hren.

Prozessexperte
Bereitstellung technischer und wissenschaftlicher Expertenunterst ü tzung an vorderster Front f ü r alle prozessspezifischen Fragen, um die terminliche Durchf ü hrung von Prozessen sicherzustellen (Business Continuity); einhaltung von cGMPs, SOPs und anwendbaren Richtlinien und funktionalen Standards (z. B. HSE, NOSSCE) und eine kontinuierliche Verbesserung der Qualit ä t und Produktivit ä tseffizienz zu erm ö glichen.

Operational Scheduler
Der Operational Scheduler erstellt und f ü hrt einen aktuellen Plan f ü r die Aktivit ä ten im Zusammenhang mit der Fertigungseinheit. Der Operational Scheduler ist verantwortlich f ü r die Entwicklung verschiedener Produktionsszenarien (Laufzeiterh ö hung, Prozess ä nderungen, Hochlauf, Schichtmodellanpassung etc.) sowie die Verbesserung des Planungstools.

Manufacturing Systems Expert
MES Expert bietet technisches Fachwissen zur Unterstützung aller Fragen im Zusammenhang mit elektronischen Chargendatensätzen (eBRs). MES Expert unterstützt den MES-Einsatz, die Implementierung und kontinuierliche Verbesserung der Fertigungseinheiten und bietet routinemäßigen technischen Support in der Werkstatt.

Technischer Trainer
Liefert technische Schulungen und bewertete Lernergebnisse für den zugewiesenen Bereich. Kann auch Lerninterventionen entwerfen und entwickeln. Kann auch die Schulung Audit-Antwort für den Standort leiten.

About the Role

Key Responsibilities:

- Provide support in the coordination and production of biopharmaceuticals in Cell Culture Process as well as in GMP processes.
- Handling of deviations, implementation / coordination of defined actions as well as implementation of approved change requests.
- Collaborating with cross-functional teams
- Support during regulatory inspections and in responding to regulatory inquiries.
- Creation, approval, and distribution of production-relevant documents.
- Ensure real-time shop floor support and the completion of production operations on time, in accordance with the documentation and in compliance with GMP rules.
- Collaboration and management of improvement projects in the production environment.
- Preparation / execution of training for production personnel regarding industrial hygiene, GMP and safety.
- Demonstrates a strong awareness of quality and a proactive, self-directed work approach.

Essential Requirements:

- Completed studies in the field of Process Engineering, Biotechnology, Biology, Pharmaceutical Technology, Technical Chemistry, Pharmacy (or comparable studies), min. Master's degree or equivalent.
- Team player, good communication skills
- Good technical and automation understanding
- Proficiency in English (spoken & written); German native speaker

Desirable Requirements:

- First experience in the pharmaceutical industry, preferably manufacturing of large molecules or development.

You ' ll receive:

You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook. https://www.novartis.com/careers/benefits-rewards
In addition to a market-competitive base salary, we offer an attractive incentive program, a modern company pension scheme, childcare facilities, learning and development opportunities as well as worldwide career possibilities within the Novartis group. In accordance with Austrian law, we are obliged to disclose the minimum salary as stated in the collective bargaining agreement. For this position the minimum salary is € 59.781,96 year (on a full-time basis). In most cases, the actual salary will be higher, as we strive to maintain a competitive position in the market and consider your previous experience, qualifications and individual competencies. We are open for part-time and job-sharing models and support flexible and remote working where possible.

This is a temporary role (1.5 years) with no relocation package offered.

Commitment to Diversity & Inclusion:*We are committed to building an outstanding, inclusive work environment and diverse teams representative of the patients and communities we serve*.

Adjustments for Applicants with Disabilities:
If because of a medical condition, physical disability or a neurodiverse condition you require an adjustment during the recruitment process, please reach out to disabilities.austria@novartis.com and let us know the nature of your request as well as your contact information. The support which we can provide will include advice on suitable positions as well as guidance at all stages of the application process. Austrian law provides candidates the opportunity to involve the local disability representative, Behindertenvertrauensperson (BVP), in the application process. If you would like to request this, please let us know in advance as a note on your CV.

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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
Operations

Business Unit
Innovative Medicines

地点
Austria

站点
Schaftenau

Company / Legal Entity
AT33 (FCRS = AT033) Novartis Pharmaceutical Manufacturing GmbH

Functional Area
Technical Operations

Job Type
Full time

Employment Type
Befristet (Innendienst) (Befristet)

Shift Work
No

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