Vodja tehnolo š kih prenosov (m/ ž /d)/Technical Transfer Lead (m/f/d)

Job ID
REQ-10046522

6月 09, 2025

Slovenia

摘要

V Biolo š kih u č inkovinah Menge š i š č emo vodjo tehnolo š kih prenosov (ž /m/d), ki bo odgovorena za vodenje prenosov tehnolo š kih postopkov pridobivanja biolo š kih u č inkovin. Delo vklju č uje vodenje in koordinacijo projektnih aktivnosti, ocenjevanje tehnolo š ke zmogljivosti in izvedljivosti prenosov, pripravo dokumentacije za prenose, ter zagotavljanje u č inkovitosti in kvalitete znotraj predvidenih č asovnic. Kot vodja tehnolo š kih prenosov na lokaciji boste odgovorni za vodenje in usmerjanje ekipe strokovnjakov z razli č nih podro č ij ter sodelovanje z drugimi proizvodnimi lokacijami in razvojnimi organizacijami pri prenosu tehnologij v proizvodnjo. Pridru ž ite se nam v dinami č nem in inovativnem okolju!

English:

At Biolo š ke u č inkovine Menge š , we are looking for a Technical Transfer Lead (m/f/d) who will be responsible for leading the transfer of technological processes for the production of biological substances. The job includes leading and coordinating project activities, assessing technological

capacity and feasibility of transfers, preparing documentation for transfers, and ensuring efficiency and quality within the planned timelines. In this role you will be responsible for leading and guiding a team of experts from various fields and collaborating with other production sites and development organizations in transferring technologies into production. Join us in a dynamic and innovative environment!

About the Role

Va š e klju č ne odgovornosti:

- Odgovorni ste za prenos tehnologije proizvodnje biolo š kih u č inkovin na lokaciji.
- V sodelovanju z drugimi dele ž niki izdelate oceno tehni č ne izvedljivosti za odlo č itev o lokaciji oskrbe, dolo č a obseg in strategijo tehni č nih serij za prenos, pripravlja splo š no projektno strategijo in projektni na č rt, vklju č no z viri in č asovnim okvirom. Spro ž ite in nadzirate postopek za lokalni nadzor sprememb povezanih s prenosom tehnolo š kih procesov.
- V primeru pogodbenega sodelovanja zagotavljate skladnost procesov tehnolo š kih prenosov s pogodbami o kakovosti in poslovnem sodelovanju.
- Oblikujete in vodite projektno ekipo na lokaciji, koordinirate dejavnosti projektne ekipe na lokaciji, povezujete strokovnjake obeh lokacij in poro č ate projektnemu vodstvu in vodstvu enote.
- Zagotovite pravo č asno razpolo ž ljivost tehni č ne dokumentacije. Izdelate in uskladite dokumente o prenosu proizvodnega procesa (protokol, poro č ilo).
- V sodelovanju z drugimi strokovnjaki pregledate in posodobite oceno tveganj kakovost (QRA) pred prenosom in pred validacijo, po potrebi prilagodite strategijo nadzora procesa. Pregledate prvi APQR po prenosu, da zagotovite ustrezno delovanje procesa in kvaliteto izdelka.
- Prispevajte k izbolj š anju in optimizaciji procesov za prenose izdelkov.
- Zagotovite, da so na voljo vse ustrezne tehni č ne informacije in dokumentacija za validacijo.
- Koordinirate izdelavo tehni č nih, regulatornih in validacijskih serij na lokaciji v skladu z opredeljeno strategijo validacije. Nudite podporo ekspertu validacij pri ustvarjanju validacijskega protokola in poro č ila.

Va š doprinos k delovnem mestu:

- Univerzitetna diploma iz farmacije, farmacevtske tehnologije, kemije, biologije, biotehnologije, in ž enirskih ved ali drugega ustreznega znanstvenega podro č ja. Za ž elen je magisterij ali doktorat.
- Izku š nje pri vodenju tehnolo š kih projektov v proizvodnji ali razvoju, poznavanje GMP proizvodnje, izku š nje v biotehnolo š ki proizvodnji ali drugih primerljivih industrijah ali institucijah (farmacevtska, ž ivilska, kemijska industrija).
- Projektne vodstvene sposobnosti.
- Mo č ne komunikacijske sposobnosti.
- Znanje angle š kega jezika.

Z izbranim kandidatom bomo sklenili delovno razmerje za **nedolo** č **en** č **as** poskusno dobo **6 mesecev.**

Zakaj Novartis?

Na š namen je soustvarjati medicino za izbolj š anje in podalj š evanje ž ivljenja ljudi, na š a vizija pa je postati najbolj cenjeno in zaupanja vredno farmacevtsko podjetje na svetu. Kako lahko to dose ž emo? S pomo č jo na š ih ljudi. Prav na š i sodelavci nas vsak dan spodbujajo, da dose ž emo svoje ambicije. Postanite del te misije in se nam pridru ž ite! Ve č na spodnji povezavi: https://www.novartis.com/about/strategy/people-and-culture

Kaj nudimo:

Konkuren č en pla č ni paket, letni bonus, fleksibilen na č in dela, z mo ž nostjo prilagajanja urnika in delom od doma, zaposlitev v podjetju s certifikatom TOP Employer, pokojninsko shemo, shemo nagrajevanja in priznanja dose ž kov, raz š irjeni program promocije zdravja na podro č ju telesnega, du š evnega in dru ž benega po č utja (Polni ž ivljenja) ter dogodke, neomejene prilo ž nosti za u č enje in razvoj.

Predani smo raznolikosti in vklju č **enosti:**

Novartis si prizadeva ustvariti izjemno, vklju č ujo č e delovno okolje in oblikovanje raznolikih timov, saj ti predstavljajo na š e bolnike in skupnosti, ki jih oskrbujemo.

Pridru ž **ite se na** š **i mre** ž **i Novartis:**

V kolikor se ne prepoznate v zgornjem opisu delovnega mesta, vas vabimo, da se vpi š ete na spodnji povezavi v Novartisovo bazo talentov saj lahko tako va š o vlogo upo š tevamo za podobne pozicije v prihodnosti: https://talentnetwork.novartis.com/network

English version:

Key Responsibilities:

- You are responsible for the transfer of technology for the production of biological substances at the location.
- In collaboration with other stakeholders, you create a technical feasibility assessment for the decision on the supply location, determine the scope and strategy of technical batches for the transfer, prepare the overall project strategy and project plan, including resources and timeline. You initiate and oversee the process for local change control related to the transfer of technological processes.

- In case of contractual cooperation, you ensure compliance of technology transfer processes with quality and business cooperation agreements.
- Form and lead the project team at the location, coordinate the activities of the project team at the location, connect experts from both locations, and report to the project management and unit management.
- Ensure the timely availability of technical documentation. You create and harmonize documents for the transfer of the production process (protocol, report).
- In collaboration with other experts, you review and update the quality risk assessment (QRA) before the transfer and before validation, adjust the process control strategy if necessary. You review the first APQR after the transfer to ensure proper process operation and product quality.
- Contribute to the improvement and optimization of processes for product transfers.
- Ensure that all relevant technical information and documentation for validation are available.
- Coordinate the production of technical, regulatory, and validation batches at the location in accordance with the defined validation strategy. You provide support to the validation expert in creating the validation protocol and report.

What you will bring to the role:

- A university degree in pharmacy, pharmaceutical technology, chemistry, biology, biotechnology, engineering, or another relevant scientific field is required. A master's or doctoral degree is preferred.
- Experience in leading technological projects in production or development, knowledge of GMP production, and experience in biotechnological production or other comparable industries or institutions (pharmaceutical, food, chemical industry).
- Project management skills.
- Strong communication abilities.
- Proficient in English language.

We offer **permanent employment** with **6 months** of probation period.

Why Novartis?

Our purpose is to reimagine medicine to improve and extend people ' s lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: https://www.novartis.com/about/strategy/people-and-culture

You ' ll receive:

Competitive salary, Annual bonus, Flexible working schedule, tailored to your needs, possibility to work from home, Pension scheme, Employee Recognition Scheme, Expanded program for the promotion of health in the field of physical, mental and social well-being, Unlimited learning and

development opportunities.

Commitment to Diversity and Inclusion:

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

Join our Novartis Network:
If this role is not suitable to your experience or career goals but you wish to stay connected to hear more about Novartis and our career opportunities, join the Novartis Network here: https://talentnetwork.novartis.com/network

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
Operations

Business Unit
Innovative Medicines

地点
Slovenia

站点
Menge š

Company / Legal Entity
SI19 (FCRS = SI019) Novartis farmacevtska proizvodnja d.o.o.

Functional Area
Technical Operations

Job Type
Full time

Employment Type
Regular

Shift Work
No

Apply to Job

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 diversity.inclusionslo@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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List of links present in page

1. http://www.novartis.com/about/strategy/people-and-culture
2. http://talentnetwork.novartis.com/network
3. http://talentnetwork.novartis.com/network
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