Vi š ji ekspert za oskrbo zdravil (m/ ž /d) / Senior Expert Drug Supply (m/f/d)

Job ID
REQ-10057640

7月 16, 2025

Slovenia

摘要

S ponosom sporo č amo, da smo ustanovili novo proizvodno kliniko v Sloveniji, namenjeno pospe š evanju ustvarjanja inovativnih zdravil za bolnike po vsem svetu. Najsodobnej š a klinika, ki se nahaja v Biocampusu Menge š , ponuja neprimerljive prilo ž nosti za sodelovanje, inovacije in vpliv.

Trenutno i š č emo zainteresirane in usposobljene strokovnjake za ekipo za klini č no proizvodnjo zdravilnih u č inkovin na na š i lokaciji TRD v Meng š u, Slovenija.

Kot Vi š ji ekspert za oskrbo zdravil bo va š a vloga vklju č evala delo kot tehnolog v klini č ni proizvodnji zdravilnih u č inkovin. Klju č ne odgovornosti vklju č ujejo zagotavljanje tehni č ne podpore za vzpostavitev proizvodnih procesov, razvoj konceptov, kot je validacija č i š č enja, pripravo dokumentacije, npr. glavni proizvodni zapisi, in podporo rutinskim proizvodnim dejavnostim.

Postanite del dinami č ne ekipe, ki si prizadeva za preoblikovanje medicine in prina š anje upanja tistim, ki ga najbolj potrebujejo. Pridru ž ite se nam pri oblikovanju prihodnosti zdravstvenega varstva in ustvarjanju pomembne razlike v ž ivljenju bolnikov po vsem svetu. Veselimo se va š ega prihoda v

na š o ekipo!

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We are thrilled to announce the establishing of new clinical manufacturing facility in Slovenia, dedicated to accelerating the creation of innovative medicines for patients around the globe. This cutting-edge facility, located at Biocampus Menge š , offers unparalleled opportunities for collaboration, innovation, and impact.

We are currently looking to hire passionate and skilled specialists in the Drug Substance Clinical Manufacturing Team at our TRD site in Menge š , Slovenia.

As Senior Expert Drug Supply, your role will involve working as a technologist within drug substance clinical production. Key responsibilities include providing technical support for establishing production processes, developing concepts such as cleaning validation, preparing documentation like master batch records, and supporting routine production activities.

Be part of a dynamic team that is reimagining medicine and delivering hope to those who need it most. Join us in shaping the future of healthcare and making a meaningful difference in the lives of patients worldwide. We look forward to welcoming you to our team!

About the Role

Va š **e klju** č **ne odgovornosti:**

- Razumevanje praks, konceptov in procesov na podro č ju klini č ne proizvodnje zdravilnih u č inkovin in sodelovanje pri vzpostavitvi in vzdr ž evanju proizvodnega objekta za klini č ni material zgodnjega tehni č nega razvoja.
- Na č rtovanje, razporejanje in koordiniranje serij, kvalifikacij, validacij in drugih dejavnosti ekipe; kot vi š ji strokovnjak podpira dejavnosti prenosa tehnologije.
- Vodenje razvoja in uvajanje novih tehnologij na tem podro č ju.
- Samostojno na č rtovanje, izvajanje, nadziranje in spremljanje vseh dejavnosti dodeljenih ekip/projektov v dolo č enih č asovnih okvirih.
- Prispevanje k notranjim (npr. GGA) in zunanjim presojam (npr. JAZMP), sodelovanje z zdravstvenimi organi in delovanje kot tehni č ni strokovnjak pri revizijah.
- Grajenje in vzdr ž evanje mre ž e znotraj in zunaj organizacije za pridobivanje znanj in najbolj š ih praks.
- Biti vzor kulture odgovornosti, raznolikosti in vklju č enosti, zaupanja, visoke u č inkovitosti ter nenehnih izbolj š av.
- Prevzemanje odgovornosti za osebni in strokovni razvoj.

Va š **doprinos k delovnem mestu:**

- Vsaj diploma ali enakovredna izobrazba iz farmacije, biokemije, biotehnologije, kemije, mikrobiologije ali enakovrednega podro č ja. Za ž elena je magistrska diploma iz tehni č ne, znanstvene ali druge ustrezne discipline.
- Najmanj 3 leta sorodnih izku š enj kot strokovnjak na podro č ju tehni č nega razvoja zdravil.

- Tehni č no znanje angle š č ine.
- Miselnost re š evanja problemov s proaktivnim pristopom k iskanju re š itev.
- Dobre komunikacijske ve š č ine. Sposobnost u č inkovitega komuniciranja in sodelovanja z razli č nimi medfunkcijskimi ekipami.

Za ž **elene izku** š **nje:**

- Znanje GMP in izku š nje z delom v reguliranem proizvodnem okolju.
- Dobro poznavanje instrumentacije ali opreme, ki je pomembna za proizvodnjo biolo š kih zdravilnih u č inkovin, in dobro poznavanje biotehnolo š kih proizvodnih procesov.

Z izbranim kandidatom bomo sklenili delovno razmerje za **nedolo** č **en** č **as** s poskusno dobo **6 mesecev.** Prijavo oddajte z ž ivljenjepisom v slovenskem in angle š kem jeziku.

Ugodnosti in nagrajevanje: Konkuren č en pla č ni paket, letni bonus, fleksibilen na č in dela z mo ž nostjo prilagajanja urnika in delom od doma, pokojninska shema, shema nagrajevanja in priznanja dose ž kov, raz š irjeni program promocije zdravja na podro č ju telesnega, du š evnega in fizi č nega po č utja (iniciativa Polni ž ivljenja), š tevilne prilo ž nosti za u č enje in razvoj.

Preberite na š priro č nik, da spoznate na č ine, s katerimi bomo spodbujali va š osebni in profesionalni razvoj: https://www.novartis.com/careers/benefits-rewards

Zakaj Novartis: Pomagati bolnikom in njihovim dru ž inam zahteva ve č kot le inovativno znanost. Potrebna je skupnost zavzetih ljudi, kot ste vi. V Novartisu cenimo sodelovanje, podporo in navdihovanje drug drugega za razvoj prebojnih terapij, ki spreminjajo ž ivljenja pacientov. Ste pripravljeni ustvariti svetlej š o prihodnost skupaj z nami?
https://www.novartis.com/about/strategy/people-and-culture

Pridru ž **ite se Novartisu:** Ni pravo delovno mesto za vas? Prijavite se v na š o bazo talentov, da ostanete v kontaktu z nami in se seznanite z ustreznimi kariernimi prilo ž nostmi takoj, ko se pojavijo: https://talentnetwork.novartis.com/network

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.

Dostop in prilagoditve: V Novartisu si prizadevamo k vklju č enosti oseb z invalidnostjo in zagotavljanju

ustreznih prilagoditev delovnega okolja posameznikom z omejitvami. V kolikor zaradi bolezni ali invalidnosti potrebujete ustrezne prilagoditve v kateremkoli delu selekcijskega procesa oziroma potrebujete prilagoditve pri izvajanju osnovnih nalog na delovnem mestu, nam pi š ite na naslov diversity.inclusionslo@novartis.com in navedite, kak š ne prilagoditve potrebujete ter va š e kontaktne podatke. Prosimo, vklju č ite tudi podatek o š tevilki razpisa, na katerega se prijavljate.

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Key Responsibilities:

- Understands applied practice, concepts and processes of Drug substance clinical manufacturing field and engages in setting-up and maintaining the FiH facility.
- Plans, schedules and coordinates batches, qualification, validation and other team activities; supports tech transfer activities as a senior expert.
- Drives development and implementation of new technologies in the field.
- Independently designs, plans, performs, supervises and monitors all activities of assigned teams/ projects within timelines.
- Significantly contributes to internal (e.g. GGA) and external audits (e.g. JAZMP), interacts with health authorities and acts as technical expert in audits.
- Builds and sustains strong network in- and out-side the organization for knowledge gain, best practice, lessons learned, and synergies.
- Role models a culture of accountability, diversity and inclusion, trust, high performance as well as continuous improvement
- Takes Responsibility for personal and professional development.

Essential Requirements:

- At least Bachelor ' s degree or equivalent in Pharmacy, Biochemistry, Biotechnology, Chemistry, Microbiology or equivalent. Master ' s degree in technical, scientific or relevant discipline is desirable.
- Minimum 3 years of related experience as an expert in biologics drug product technical research and development.
- Technical knowledge of English.
- Problem-solving mindset with a proactive approach to finding solutions.
- Good communication skills. Ability to effectively communicate and collaborate with diverse cross-functional teams.

Desirable Requirements:

- Knowledge of GMP and experience working in a regulated manufacturing environment.
- Good knowledge of instrumentation or equipment relevant to biologic drug substance production and good knowledge of biotechnological manufacturing processes.

We offer **permanent employment** with **6 months** of probation period. Submit your application with the CV in Slovenian and English language.

Benefits and Rewards: 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 (Well-being), Unlimited learning and development opportunities.

Read our handbook to learn about all the ways we ' ll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

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.

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.

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
Development

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
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

Apply to Job

Accessibility and accommodation

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