

Ekspert za oskrbo zdravil (m/ ž /d) / Expert Drug Supply (m/f/d)

Job ID
REQ-10060721

8月 26, 2025

Slovenia

摘要

#LI-Hybrid

Z veseljem sporočamo, da smo v Sloveniji odprli nov klinični proizvodni obrat, namenjen pospeševanju ustvarjanja inovativnih zdravil za paciente po vsem svetu. Ta najsodobnejši obrat, ki se nahaja v Biocampusu Menges, ponuja neprimerljive priložnosti za sodelovanje, inovacije in vpliv.

Trenutno iščemo motivirane in usposobljene strokovnjake za ekipo za klinično proizvodnjo zdravilnih učinkovin na naši lokaciji TRD v Mengesu v Sloveniji.

Kot Ekspert za oskrbo z zdravili boste procesni specialist v klinični proizvodnji zdravilnih učinkovin. Vaša ključna odgovornost vključuje mikro načrtovanje, podporo izvajanju in nadzor proizvodnega procesa ali dejavnosti kvalifikacije proizvodne opreme, ter podporo drugim dejavnostim (npr. change over, upravljanje odstopanj v proizvodnji). Postanite del dinamičnega tima, ki na novo opredeljuje zdravljenje in prinaša upanja tistim, ki ga najbolj potrebujejo.

Pridružite se nam pri oblikovanju prihodnosti zdravstva in ustvarjanju pomembnih sprememb v

življenju pacientov po vsem svetu. Veselimo se, da vas bomo pozdravili v naši ekipi!

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 Mengeš, 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 Mengeš, Slovenia.

As Expert Drug Supply, your role will be a process specialist within the Drug substance clinical production, with key responsibilities including microplanning, supporting the execution and overseeing the production process or production equipment qualification activities, as well as supporting other activities (e.g. change over, management of deviations within the production).

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

Vaš ključna odgovornosti:

- Načrtovanje, razporejanje in usklajevanje proizvodnega procesa, kvalifikacij, validacij in drugih aktivnosti ekipe.
- Samostojno ocenjevanje in interpretiranje rezultatov ter sprejemanje ustreznih zaključkov.
- Zagotavljanje znanstvenega in tehničnega vodenja; iskanje informacij, aktivno spodbujanje izmenjave znanja znotraj skupine.
- Sodelovanje v multidisciplinarni ekipi in zagotavljanje podpore, mentorstva in nadzora mlajšim članom. Prenos znanja znotraj ekip in na druge dele ženske.
- Samostojno pisanje protokolov, zapisov glavnih serij, standardnih operativnih postopkov, proizvodnih receptov in druge dokumentacije. Aktivno sodelovanje pri revizijah in zagotavljanje skladnosti.
- Komuniciranje, obravnavanje in reševanje odstopanj. Prispevanje k odpravljanju težav, preiskavam temeljnih vzrokov in predlaganje korektivnih ukrepov.
- Prezemanje odgovornosti, odprte komunikacije in kulture reševanja problemov/osredotočenosti na rezultate, pri čemer je kakovost glavna prednostna naloga.
- Prezemanje odgovornosti za osebni in poklicni razvoj.

Vaš doprinos k delovnem mestu:

- Univerzitetna izobrazba s področja farmacije, biokemije, biotehnologije, kemije, mikrobiologije ali druge enakovredne smeri.
- Tehnično znanje angleščine.

- Več let sorodnih izkušenj kot strokovnjak za proizvodnjo bioloških zdravil.

Zaželene izkušnje:

- Poznavanje dobre proizvodne prakse (GMP) in izkušnje z delom v reguliranem proizvodnem okolju.
- Dobro poznavanje instrumentov ali opreme, pomembne za proizvodnjo bioloških zdravilnih učinkovin, in dobro poznavanje biotehnoloških proizvodnih procesov.

Z izbranim kandidatom bomo sklenili delovno razmerje za nedoločeno časovno dobo 6 mesecev. Prijavo oddajte z življenjepisom v slovenskem in angleškem jeziku.

Ugodnosti in nagrajevanje: Konkurenčni plačni paket, letni bonus, fleksibilen način dela z možnostjo prilagajanja urnika in delom od doma, pokojninska shema, shema nagrajevanja in priznanja dosežkov, razširjeni program promocije zdravja na področju telesnega, duševnega in fizičnega počutja (iniciativa Polni življenja), številne priložnosti za učenje in razvoj.

Preberite naš priročnik, da spoznate načine, s katerimi bomo spodbujali vaš osebni in profesionalni razvoj: <https://www.novartis.com/careers/benefits-rewards>

Zakaj Novartis: Pomagati bolnikom in njihovim družinam zahteva več 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 življenja pacientov. Ste pripravljeni ustvariti svetlejšo prihodnost skupaj z nami? <https://www.novartis.com/about/strategy/people-and-culture>

Pridružite se Novartisu! Ni pravo delovno mesto za vas? Prijavite se v našo bazo talentov, da ostanete v kontaktu z nami in se seznanite z ustreznimi kariernimi priložnostmi takoj, ko se pojavijo: <https://talentnetwork.novartis.com/network>

Predani smo raznolikosti in vključenosti! Novartis si prizadeva ustvariti izjemno, vključujoče delovno okolje in oblikovanje raznolikih timov, saj ti predstavljajo naše bolnike in skupnosti, ki jih oskrbujemo.

Dostop in prilagoditve: V Novartisu si prizadevamo k vključenosti 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 pišite na naslov diversity.inclusionslo@novartis.com in navedite, kakšne prilagoditve potrebujete ter vaše kontaktne podatke. Prosimo, vključite tudi podatek o številki razpisa, na katerega se prijavljate.

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Key Responsibilities:

- Plans, schedules, and coordinates batches, qualification, validation, and other team activities.
- Evaluates and interprets results, draws relevant conclusions independently. Provides scientific and technical guidance; performs information searches; actively fosters knowledge exchange within peer population.
- Works collaboratively within a multidisciplinary team, providing support, coaching, mentoring, and supervision to junior members. Transfers process know-how within teams and other stakeholders.
- Writes protocols, master batch records, SOPs, working procedures, production recipes, and other documents independently. Actively contributes to audits and ensures compliance.
- Communicates, addresses, and solves deviations. Contributes to troubleshooting, root cause investigations, and proposes corrective actions.
- Role models accountability, open communication and problem solving/ result driven culture, with quality as a top priority.
- Takes Responsibility for personal and professional development.

Essential Requirements:

- Bachelor's degree or equivalent in Pharmacy, Biochemistry, Biotechnology, Chemistry, Microbiology or equivalent.
- Technical knowledge of English.
- Several years of related experience as expert in biologics manufacturing.

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>

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

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