NOVARTIS

Strokovni sodelavec za oskrbo zdravil (m/ž/d) / Associate Expert Drug Supply (m/f/d)

Job ID
REQ-10077800
май 20, 2026
Словения
Available in: English

Сводка

Location: Menges, Slovenia #onsite
*please note that this is a 1 year temp role
*Prosimo, upoštevajte, da gre za začasno delovno mesto za 1 leto

Z veseljem napovedujemo ustanovitev novega obrata klinične proizvodnje v Sloveniji, namenjenega hitrejšemu odkrivanju inovativnih zdravil za bolnike po vsem svetu. Najsodobnejši objekt, ki se nahaja v Biocampusu v Mengšu, nudi izjemno priložnost za sodelovanje na področju razvoja in proizvodnje inovativnih zdravil.

Iščemo navdušene in usposobljene strokovnjake za proizvodni tim. Kot strokovni sodelavec za oskrbo zdravil boste del tima Proizvodnih operacij.

Odgovorni boste za izvajanje proizvodnega procesa bioloških učinkovin, vključno s pripravo raztopin, upravljanjem bioreaktorjev, uporabo opreme za izolacijo učinkovine (kromatografski sistemi) in druga procesna oprema. Aktivnosti se izvajajo skladno z internimi postopki in zahtevami GxP.

Postanite del dinamičnega tima, ki na novo opredeljuje zdravljenje in prinaša upanje tistim, ki ga najbolj potrebujejo. Pridružite se nam pri oblikovanju prihodnosti varovanja zdravja in pri ustvarjanju pomembnih razlik v življenju bolnikov po vsem svetu. Veselimo se vašega prihoda v naš tim!

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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 Mengeš, offers an exceptional opportunity for collaboration in the development and production of innovative medicines.

We are currently looking to hire passionate and skilled specialists in the Manufacturing Operations team.

As part of our team, you will be responsible for executing the production process of biological active substances, including the preparation of solutions, operation of bioreactors, use of equipment for active substance isolation such as chromatography systems, and other process equipment. Activities are performed in accordance with internal procedures and GxP requirements.

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

Opis delovnega mesta

Vaše ključne odgovornosti:

- Izvedba procesa proizvodnje bioloških učinkovin, vključno z izvajanjem in-procesne analitike.
- Planiranje, izvajanje in dokumentiranje GxP aktivnosti.
- Izpolnjevanje GxP dokumentacije (dnevniki, obrazci, proizvodna poročila itd.), vključno s pregledom izvornih podatkov. Pisanje enostavnih postopkov, protokolov in poročil (delovni postopki, poročila o trendih itd.)
- Sodelovanje pri reševanju izzivov in odpravljanju težav. Prepoznavanje, sporočanje in prispevanje k reševanju odstopanj ter izvajanje korektivnih in preventivnih ukrepov. Uporaba pridobljenih izkušenj.
- Delovanje v skladu s standardi za kakovost, etiko, varnost, zdravje, okolje in informacijsko varnost ter zagotavljanje upoštevanja predpisov GxP.
- Odgovornost za osebni in strokovni razvoj.
- Izkazovanje pozitivne delovne etike in pozitivno vplivanje na druge.

Vaš doprinos k delovnem mestu:

- Srednješolska izobrazba.
- Tekoče znanje slovenščine. Tehnično znanje angleščine.
- Dobre organizacijske sposobnosti in sposobnosti upravljanja z dokumentacijo, ki zagotavljajo vodenje evidenc v skladu s pravili podjetja.

- Sposobnost natančnega upoštevanja navodil in postopkov.
- Ustrezno poznavanje programske opreme in računalniških orodij.

Zaželene izkušnje:

- Izkušenje s primerljivega delovnega mesta.
- Ustrezno strokovno ali tehnično znanje s področja naravoslovnih ved.
- Dobro poznavanje dobre proizvodne prakse (GMP) in izkušnje z delom v reguliranem proizvodnem okolju.

Z izbranim kandidatom bomo sklenili delovno razmerje za določen čas s poskusno dobo 6 mesecev in možnostjo podaljšanja. Prijavo oddajte z življenjepisom v slovenskem jeziku.

Kaj nudimo:

Konkurenčen plačni paket, letni bonus, fleksibilen način dela z možnostjo prilagajanja urnika, zaposlitev v podjetju s certifikatom TOP Employer, pokojninsko shemo, shemo nagrajevanja in priznanja dosežkov, razširjeni program promocije zdravja na področju telesnega, duševnega in družbenega počutja (Polni življenja) ter dogodke, neomejene priložnosti za učenje in razvoj.

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.

Job description

Your key responsibilities:

- Executing the production process of biological active substances, including performing in-process analytics.
- Planning, performing, and documenting GxP activities.
- Completing GxP documentation, such as logs, forms, production reports, etc., including the review of source data. Writing simple procedures, protocols, and reports, such as work procedures, trend reports, etc.
- Participating in resolving challenges and troubleshooting. Identifying, reporting, and contributing to the resolution of deviations, as well as implementing corrective and preventive actions. Applying lessons learned.
- Acting in accordance with standards for quality, ethics, safety, health, environment, and information security, and ensuring compliance with GxP regulations.
- Taking responsibility for personal and professional development.
- Demonstrating a positive work ethic and positively influencing others.

Your contribution to the role:

- High school education.
- Fluent knowledge of Slovenian lenguage. Technical knowledge of English.
- Good organizational and documentation management skills, ensuring record-keeping in accordance with company rules.
- Ability to follow instructions and procedures accurately.
- Appropriate knowledge of software and computer tools.

Desired experience:

- Experience in a comparable role.
- Relevant professional or technical knowledge in the field of natural sciences.
- Good knowledge of Good Manufacturing Practice, GMP, and experience working in a regulated manufacturing environment.

An employment contract will be concluded with the selected candidate for a fixed term, with a 6-month probationary period and the possibility of extension. Please submit your application with a CV in Slovenian lenguage.

What we offer:

A competitive salary package, annual bonus, flexible working arrangements with the possibility of schedule adjustments, employment in a company certified as a TOP Employer, a pension scheme, a rewards and recognition scheme, an expanded health promotion program focused on physical, mental, and social well-being, “Full of Life,” as well as events and unlimited opportunities for learning and development.

We are committed to diversity and inclusion

Novartis is committed to creating an outstanding, inclusive work environment and building diverse teams, as these represent 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

Benefits and Rewards: Learn about all the ways we’ll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

Primary location salary range
€20,300.00 - €37,700.00
Дивизион
Development
Business Unit
Development
Место
Словения
Сайт
Mengeš
Company / Legal Entity
SI19 (FCRS = SI019) Novartis farmacevtska proizvodnja d.o.o.
Functional Area
Research & Development
Job Type
Full time
Employment Type
Temporary (Fixed Term)
Shift Work
No

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.inclusion_slo@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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