Vodja podro č ja klini č ne proizvodnje (m/ ž /d) / Associate Director Clinical Manufacturing - Sterility Assurance (m/f/d)

Job ID
REQ-10046513

4月 03, 2025

Slovenia

摘要

Z veseljem napovedujemo ustanovitev novega obrata klini č ne proizvodnje v Sloveniji, namenjenega hitrej š emu odkrivanju inovativnih zdravil za bolnike po vsem svetu. Najsodobnej š i objekt, ki se nahaja v Biocampusu v Meng š u, nudi izjemno prilo ž nost za sodelovanje, inoviranje in vpliv.

I š č emo navdu š ene in usposobljene strokovnjake za tim Klini č ne proizvodnje zdravilnih u č inkovin na š lokaciji TRD v Meng š u, Slovenija.

Kot Vodja podro č ja klini č ne proizvodnje boste odgovorni predvsem za vodenje timov v klini č ni proizvodnji za zagotavljanje klini č nega materiala visoke kakovosti ter za vodenje in razvijanje strategije klini č ne proizvodnje.

Postanite del dinami č nega tima, ki na novo opredeljuje zdravljenje in prina š a upanje tistim, ki ga najbolj potrebujejo. Pridru ž ite se nam pri oblikovanju prihodnosti varovanja zdravja in pri ustvarjanju pomembnih razlik v ž ivljenju bolnikov po vsem svetu. Veselimo se va š ega prihoda v na š 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 new 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 Sterile Drug Product Clinical Manufacturing Team at our TRD site in Menge š , Slovenia.

As Associate Director Clinical Manufacturing - Sterility Assurance, you will be primarily responsible for leading and managing teams responsible for ensuring sterility assurance and contamination control at the Clinical Manufacturing Plant.

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:**

- Odgovornost za vodenje timov v klini č ni proizvodnji za zagotavljanje klini č nega materiala visoke kakovosti ter za vodenje in razvijanje strategije klini č ne proizvodnje.
- Usmerjanje talentov na dodeljenem podro č ju: Vodenje procesov, povezanih z ljudmi, vklju č no z zaposlovanjem, usposabljanjem, izmenjavo talentov, coachingom in ocenjevanjem uspe š nosti, ter s tem izpolnjevanje operativnih zahtev ob ohranjanju konkuren č nosti in raznolikosti.
- Coaching, vodenje, razvoj in motivacija sodelavcev. Podpora gradnje tima, opolnomo č enje in vodenje tima v organizacijo, usmerjeno v procese.
- Vzpostavljanje in vzdr ž evanje trdnih odnosov s poslovnimi partnerji ter zagotavljanje u č inkovite komunikacije vizije in vrednot podjetja.
- Vodenje in nadziranje programov č i š č enja, sanitacije in okoljskega monitoringa.

Va š **doprinos k delovnem mestu:**

- Univerzitetna izobrazba s podro č ja biotehnologije, farmacije ali sorodnih ved.
- Minimalno 3 leta relevantnih vodstvenih izku š enj.
- Minimalno 5 let relevantnih izku š enj v proizvodnji / razvoju, npr. Asepti č na proizvodnja, Mikrobiolo š ko testiranje, Kontrola kakovosti, Tehni č ni prenos procesov.
- Poglobljeno znanje zahtev GMP in njihove uporabe v asepti č ni proizvodnji.
- Poglobljene komunikacijske spretnosti, sposobnost u č inkovite predstavitve kompleksnih idej in re š itev za vodstvo, tehni č ne strokovnjake in regionalne / lokalne ekipe.
- Dobro razumevanje proizvodnega okolja; sposobnost ocenjevanja poslovnih prilo ž nosti in/ali odpravljanja tveganj skupaj z relevantnimi ekipami.
- Teko č e znanje angle š č ine.

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

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, 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.

Key Responsibilities:

- Responsible for leading and managing a team responsible for Sterility assurance and contamination control within the Clinical Manufacturing Plant to deliver on high-quality clinical material and actively driving and developing the Clinical Manufacturing strategy.
- Driving the talent agenda for area of responsibility: Leading people processes through recruitment, training, talent exchange, coaching and performance to meet all operation requirements sustaining both competitiveness and diversity.
- Coaching, leadership, development and motivation of associates. Supporting team-building, empowerment and conducting the team to a process oriented organization.
- Establishing and maintaining strong relationships with business partners and ensure effective communication of company vision and values.
- Leading and controlling cleaning, sanitation, and environmental monitoring programs.

Essential Requirements:

- University degree in biotechnology, pharmacy or related sciences.
  Minimum 3 years of relevant leadership experience.
- Minimum 5 years of relevant production / development experience e.g. Aseptic Manufacturing, Microbiological Testing, Quality Control, Technical Process Transfer.
- Profound knowledge of GMP requirements and its applications in aseptic Manufacturing.
- Profound communication skills, ability to effectively present complex ideas and solutions to management, technical experts and regional / local teams.
- Good understanding of production environment; ability to evaluate business opportunities and/or remediate risks jointly with relevant teams.
- Fluent English.

We offer **permanent employment** with **6 months** of probation period. Submit your application with the

CV in Slovenian and English language.

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 (Energized for Life), 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.

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
Regul ä r

Shift Work
No

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