Vodja upravljanja kakovosti - operacije (m/ ž /d) / QA Operations Lead (m/f/d)

Job ID
REQ-10069041

12月 19, 2025

Slovenia

摘要

#LI-Hybrid
Location: Ljubljana, Slovenia

Predstavljajte si, da igrate klju č no vlogo pri tem, da ž ivljenjsko pomembna zdravila varno in skladno pridejo do bolnikov. Kot QA Operations Lead boste usmerjali ekipe, optimizirali procese in skrbeli za najvi š je standarde kakovosti skozi razvoj, prenos in komercializacijo. Va š e vodenje bo oblikovalo operativno odli č nost ter spodbujalo inovacije v dinami č nem, globalnem okolju.

Imagine playing a pivotal role in ensuring life - -changing medicines reach patients safely and compliantly. As QA Operations Lead, you ’ ll guide teams, optimize processes, and uphold the highest quality standards across development, transfer, and commercialization. Your leadership will shape operational excellence and drive innovation in a dynamic, global environment.

Relocation Support: This role is based in Ljubljana, Slovenia. Novartis is unable to offer relocation support: please only apply if accessible.

About the Role

Va š e klju č ne odgovornosti:

- Zagotavljanje skladnosti vseh aktivnosti z veljavnimi standardi dobrih praks (cGxP).
- Vodenje in podpora pri GxP presojah ter in š pekcijskih pregledih regulatornih organov.
- Nadzor kakovostnih operacij na lokaciji, vklju č no z vhodnim in izhodnim QA.
- Pregled in odobritev glavnih proizvodnih zapisov (Master Batch Records, MBR).
- Upravljanje spro š č anja izdelkov v skladu z globalnimi in lokalnimi predpisi.
- Vodenje QA aktivnosti za kontrolo kakovosti ter Analiti č ne znanosti in tehnologijo (AS&T).
- Spodbujanje operativne odli č nosti in kulture nenehnih izbolj š av.
- Optimizacija procesov in u č inkovito upravljanje operativnih stro š kov QA enote.

Va š doprinos k delovnem mestu:

- Univerzitetna izobrazba iz farmacije, biologije, kemije, mikrobiologije ali druge ustrezne naravoslovne oziroma tehni š ke smeri.
- Najmanj 5 let izku š enj na podro č ju kakovosti, razvoja, proizvodnje ali primerljivih delovnih mest.
- Funkcionalno znanje angle š kega jezika.
- Odli č no poznavanje cGxP, EU/FDA regulative in mednarodnih standardov.
- Dokazane vodstvene sposobnosti ter izku š nje z razvojem ekip.
- Sposobnost vodenja presoj, in š pekcijskih pregledov in procesov spro š č anja izdelkov.

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, mo ž nost vklju č itve v kolektivno zdravstveno zavarovanje, shema nagrajevanja in priznanja dose ž kov, raz š irjeni program promocije zdravja na podro č ju fizi č nega in du š evnega dobrega po č utja ter delovne obremenitve (Polni ž ivljenja), š tevilne 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.

Zakaj Novartis: Pomagati bolnikom in njihovim dru ž inamzahteva ve čkot 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 ivljenjapacientov.
Ste pripravljeni ustvariti svetlej š oprihodnost skupaj z nami?
https://www.novartis.com/about/strategy/people-and-culture

Pridru ž itese Novartisu: Ni pravo delovno mesto za vas? Prijavite se v na š obazo talentov, da ostanete v kontaktu z nami in se seznanite z ustreznimi kariernimi prilo ž nostmtakoj, ko se pojavijo:

Key Responsibilities:

- Ensure all activities comply with current good practices (cGxP) standards.
- Lead and support GxP audits and regulatory inspections.
- Oversee Quality Operations across the site, including inbound and outbound QA.
- Review and approve Master Batch Records (MBR).
- Manage product release in line with global and local regulations.
- Drive QA for Quality Control and Analytical Science & Technology (AS&T).
- Champion operational excellence and continuous improvement initiatives.
- Optimize processes and manage QA operational unit costs effectively.

Essential Requirements:

- University degree in Pharmacy, Biology, Chemistry, Microbiology, or equivalent natural or engineering science.
- Minimum 5 years of experience in Quality, Development, Manufacturing, or comparable positions.
- Functional knowledge of English.
- Strong understanding of cGxP, EU/FDA regulations, and international standards.
- Demonstrated leadership and team development skills.
- Ability to manage audits, inspections, and product release 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, possibility of joining collective health insurance scheme, Employee Recognition Scheme, Expanded program for the promotion of health in the field of physical and mental well-being and managing workload (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.

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: 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
Quality

地点
Slovenia

站点

Ljubljana

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

Functional Area
Quality

Job Type
Full time

Employment Type
Regular

Shift Work
No

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.

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