Vodja kontrole kakovosti (m/ž/d) / QC Lead (m/f/d), Mengeš

Job ID
REQ-10032642
Ene 08, 2025
Eslovenia

Resumen

Želiš sodelovati pri postavitvi strategije za učinkovito podporo proizvodnjih procesov s state of the art analitskimi metodami in novimi tehnologijami v analitiki? Voditi skupino visoko motiviranih sodelavcev za testiranje inovativnih in biološko podobnih učinkovin?

Pridruži se nam kot Vodja kontrole kakovosti (QC Lead) na naši lokaciji v Mengšu in postani del super ekipe.

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Do you want to be part of developing a strategy for effective support of production processes with state-of-the-art analytical methods and new technologies in analytics? Lead a team of highly motivated colleagues in testing innovative and biosimilar active ingredients?

Join us as a Quality Control (QC) Lead at our location in Mengeš and become part of a great team.

About the Role

Vaše ključne odgovornosti:

- Zagotavljanje izvajanja aktivnosti KK v skladu s standardi cGxP.
- Nadzor nad pravočasnim testiranjem vzorcev za zagotavljanje nemotenih proizvodnjih procesov
- Zagotavljanje skladnosti operacij analitičnih laboratorijev in razvoja z veljavnimi regulativnimi zahtevami in smernicami.
- Iskanje priložnosti za izboljšanje produktivnosti in usklajevanje njihove izvedbe.
- Doseganje ključnih kazalnikov uspešnosti (KPI) v zvezi s kakovostjo, stroški in ljudmi
- Omogočanje pravočasnega sproščanja izdelkov v skladu z GMP (Good Manufacturing Practice) in regulatornimi zahtevami.
- Upoštevanje in izvajanje pravil ter smernic za področje zdravja, varnosti in okolja (ZVO).

Vaš doprinos k delovnem mestu:

- Spretnost prilagajanja spreminjajočim se prioritetam in okoljem.
- Vodenje presoj
- Močno razumevanje poslovne strategije in njene uporabe v operacijah QC.
- Zavezanost odličnosti v laboratoriju in spoštovanju najboljših praks.
- Obvladovanje principov in metod nadzora kakovosti.

- Spretnost v upravljanju deležnikov za usklajevanje ciljev različnih funkcij.
- Aktivno znanje angleškega jezika (pisno in ustno).

Zaželene izkušnje:

- Izkušnje v vodenju ljudi, z zmožnostjo motiviranja in usmerjanja ekipe.
- Minimalno 5 let delovnih izkušenj na področju kakovosti, razvoja, proizvodnje v farmacevtski industriji ali na primerljivih delovnih mestih
- Visokošolska stopnja izobrazbe farmacevtske, biološke, kemijske, mikrobiološke ali druge naravoslovne smeri.

Z izbranim kandidatom bomo sklenili delovno razmerje za **nedoločen čas** s poskusno dobo **6 mesecev.**

Prijavo oddajte z življenjepisom v slovenskem in angleškem jeziku.

Kaj nudimo:

Konkurenčen plačni paket, letni bonus, fleksibilen način dela, z možnostjo prilagajanja urnika in delom od doma, 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.

Key Responsibilities:

- Ensure QC activities are executed in accordance with cGxP standards.
- Oversee the on-time testing of samples to maintain workflow efficiency.
- Ensure compliance of analytical laboratory operations and development with applicable regulatory requirements and guidelines.
- Identify opportunities for productivity improvements and coordinate their implementation.
- Achieve KPIs related to quality, cost, people, and delivery.
- Facilitate timely product release in line with GMP (Good Manufacturing Practice) and license requirements.
- Maintain a safe work environment with zero HS&E (Health, Safety & Environment) incidents.

Essential Requirements:

- Agility in adapting to changing priorities and environments.
- Experience with audit management and ensuring compliance readiness.
- Strong understanding of business strategy and its application to QC operations.
- Commitment to laboratory excellence and adherence to best practices.
- Proficiency in quality control principles and methods.
- Skilled in stakeholder management to align cross-functional goals.
- Active knowledge of English (written and spoken).

Desirable Requirements:

- Proven experience in people leadership, with the ability to motivate and guide teams.
- Over 5 years of experience in QA/QC/Production or related fields.
- University degree in pharmacy, biology, chemistry, microbiology or other natural science degree

We are offering a **permanent** employment contract, including a **6-month** 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 (Wellbeing), employment at Top SI Employer, 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

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Operations
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Innovative Medicines
Ubicación
Eslovenia
Sitio
Mengeš
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
Apply to Job

Accessibility and accommodation

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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