

Specialist upravljanja kakovosti za področje skladnosti računalniških sistemov / QA eCompliance Specialist (m/f/d)

Job ID
REQ-10071882
mar 12, 2026
Словения

Сводка

#LI-Hybrid
Location: Ljubljana, Slovenia

Vas zanima upravljanje kakovosti v farmacevtski industriji v povezavi s skladnostjo računalniških sistemov? Ste proaktivna in visoko motivirana oseba, ki si želi izzivov in tesnega timskega dela na globalnem nivoju? Imate izobrazbo tehnične ali druge primerljive smeri? Vas je vedno zanimalo kako izgledajo procesi v farmacevtski industriji? Ste natančna, zanesljiva in komunikativna oseba, ki si prizadeva za stalno profesionalno rast in išče priložnosti za napredovanje na naslednjo stopnjo v svoji karieri?

Če se najdete v zgornjem opisu vas vabimo, da se pridružite ekipi QA eCompliance na Novartisovi lokaciji v Ljubljani!

Are you interested in quality management in the pharmaceutical industry in connection with computer system compliance? Are you a proactive and highly motivated person who seeks challenges and close teamwork at a global level? Do you have a technical education? Have you always been interested in how processes work in the pharmaceutical industry? Are you a precise, reliable, and communicative person who strives for continuous professional growth and seeks opportunities for advancement to the next level in your career?

If you identify with the above description, we invite you to join the QA eCompliance team at Novartis' location in Ljubljana!

About the Role

Vaše ključne odgovornosti:

- Nudjenje podpore pri dejavnostih za kvalifikacijo in validacijo računalniško podprtih sistemov (načrtovanje, svetovanje, pregled).
- Zagotavljanje implementacije veljavnih Novartisovih in regulatornih zahtev za področje GxP računalniško podprtih sistemov.
- Pregled / odobritev nadziranja sprememb v sistemu.
- Zagotavljanje kakovosti procesa v skladu s predpisi.
- Zagotavljanje strokovnega znanja oz. usmeritev za zagotavljanje kakovosti in ustreznosti GxP relevantnih računalniško podprtih sistemov, ocenjevanje dobaviteljev, nadzor nad spremembami, obvladovanje odstopov in povezanih aktivnosti, s čimer se zagotovi skladnost z regulatornimi predpisi in uresničijo pričakovanja podjetja.
- Pregledovanje in potrjevanje ocen opreme/sistemov glede GxP relevantnosti.
- Implementiranje in razvijanje novih zmogljivosti v skladu s poslovnimi potrebami.
- Priprava in podpora pri revizijah in inšpekcijskih pregledih.

Vaš doprinos k delovnem mestu:

- Visokošolska/univerzitetna izobrazba tehnične, računalniške ali druge naravoslovne smeri.
- Aktivno znanje angleškega jezika.
- Poznavanje orodja Microsoft Office.
- Minimalno 3 let delovnih izkušenj s področja avtomatizacije/CSV in /ali avtomatizacije procesov in sistemov ter standardov s področja računalniških sistemov v farmacevtski industriji ali drugi ustrezni industriji.

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

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 dogodki in 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.

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.inclusion_slo@novartis.com in navedite, kakšne prilagoditve potrebujete ter vaše kontaktne podatke. Prosimo, vključite tudi podatek o številki razpisa, na katerega se prijavljate.

- Support site qualification and validation activities (planning, advising, review).
- Audit and inspection preparation and support.
- Change control review/approval.
- Ensure process quality assurance acc. to regulations.
- Ensure implementation of the applicable Novartis and regulatory requirements for GxP regulated computerized systems.
- Provide quality assurance expertise / guidance for GxP computerized systems classification, qualification, supplier assessment, change control, deviation management and associated activities that ensure compliance to regulatory and company expectations.
- Review and approve determination of computerized system for GxP applicability.
- Adopts & develops new capabilities in alignment with Business needs.

Essential Requirements:

- University-level education in a technical, computer science, or other natural science field.
- Active knowledge of the English language.
- Familiarity with Microsoft Office tools.
- At least 3 years of work experience in the field of automation/CSV and/or process and system automation, as well as standards related to computer systems in the pharmaceutical industry or another relevant industry.
- We offer temporary employment with 6 months of probation period

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.

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.

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 \(PDF 30 MB\)](#)

Дивизион

Operations

Business Unit

Quality

Место

Словения

Сайт

Ljubljana

Company / Legal Entity

SI19 (FCRS = SI019) Novartis farmacevtska proizvodnja d.o.o.

Functional Area

Quality

Job Type

Full time

Employment Type

Začasni sodelavec (za določen čas)

Shift Work

No

Dostop in prilagoditve

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3. mailto:diversity.inclusion_slo@novartis.com
4. https://novartis.wd3.myworkdayjobs.com/sl-SI/Novartis_Careers/job/Ljubljana/Specialist-upravljanja-kakovosti-za-podroje-skladnosti-raunalnikih-sistemov--QA-eCompliance-Specialist--m-f-d-_REQ-10071882
5. https://novartis.wd3.myworkdayjobs.com/sl-SI/Novartis_Careers/job/Ljubljana/Specialist-upravljanja-kakovosti-za-podroje-skladnosti-raunalnikih-sistemov--QA-eCompliance-Specialist--m-f-d-_REQ-10071882