Vi š ji AS&T specialist (m/ ž /d) / Senior AS&T Specialist (m/f/d)

Job ID
REQ-10046554

4月 01, 2025

Slovenia

摘要

#LI-Hybrid

Kot Vi š ji AS&T specialist boste odgovorni za vodenje analitskih projektov in boste sodelovali s š irokim timom strokovnjakov za vzpostavitev analitskih metod za podporo proizvodnji inovativnih biolo š kih zdravil. Sodelovali boste v strokovni in tehni č ni podpori na podro č ju analitike, pri odpravljanju kompleksnih te ž av ter pri upravljanju sprememb.
Č e vas veseli delo v dinami č nem okolju, se vidite v projektnem vodenju in ž elite vseeno ostati vpeti v strokovno delo, se pridru ž ite skupini strokovnjakov v AS&T.

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As a Senior AS&T Specialist, you will be responsible for leading the analytical projects and collaborating with a broad team of experts to establish analytical methods to support the production of innovative biological medicines. You will participate in expert and technical support in the field of analytics, in resolving of complex problems and in change management.
If you enjoy working in a dynamic environment, see yourself in project management, and still want to stay involved in an expert role, then join our group of experts in AS&T.

About the Role

Va š **e klju** č **ne odgovornosti:**

- Zagotavljanje podpore transfernim projektom, povezanih s proizvodnjo.
- Odgovornost za strate š ko planiranje in vodenje analitskih dejavnosti (razvoj, primerjave, validacije in prenosi metod ...) v sodelovanju z drugimi oddelki / funkcijami in projektnimi timi.
- Obvladovanje ž ivljenskega cikla analitskih metod za testiranje izdelkov ter nudenje strokovne in tehni č ne podpore pri odpravljanju kompleksnih te ž av, obvladovanju odstopov in sprememb.
- Odgovornost za interpretacijo, statisti č no vrednotenje in poro č anje rezultatov, sprejemanje zaklju č kov ter priprava poro č il.
- Nadgradnja in prenos strokovnih znanj, izobra ž evanje in razvoj sodelavcev ter odgovornost za osebni in strokovni razvoj.
- Zagotavljanje skladnosti vseh dejavnosti z dobrimi praksami (cGxP), celovitostjo podatkov ter doma č o in evropsko zakonodajo, ter sodelovanje pri internih in zunanjih presojah.
- Sodelovanje pri razvoju, implementaciji in nadzoru sistema kakovosti v skladu s slovensko in evropsko zakonodajo, FDA, mednarodnim svetom za usklajevanje tehni č nih zahtev glede zdravil, Konvencijo o farmacevtski in š pekciji in Novartisovimi standardi.
- Odgovornost za kvalifikacijo in kalibracijo opreme ter priprava na č rta kvalifikacij opreme.

Va š **doprinos k delovnem mestu:**

- Visoko š olska stopnja izobrazbe farmacevtske, biolo š ke, kemijske, mikorbiolo š ke ali druge naravoslovne smeri.
- Minimalno 5 let delovnih izku š enj na podro č ju kakovosti, razvoja ali proizvodnje ali drugega ustreznega podro č ja.
- Aktivno znanje angle š kega jezika.
- Proaktivnost, samoiniciativnost in hitro dojemanje informacij.
- Izku š nje z vodenjem projektne skupine ter delom z ve č dele ž niki.
- Aktiven, dinami č en in komunikativen pristop k delu.
- Napredna uporaba Officeovih orodij.

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.

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.

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Key Responsibilities:

- Providing support to transfer projects related to production.
- Responsibility for strategic planning and leading of the analytical activities (development, comparabilities, validations and method transfers...) in collaboration with other departments / functions and project teams.
- Managing of lifecycle of analytical methods for product testing and providing expert and technical support in the resolution of complex problems, control of deviations and changes.
- Responsibility for interpretation, statistical evaluation and reporting of results, drawing conclusions and preparing reports.
- Upgrading and transferring professional knowledge, education and development of colleagues and responsibility for personal and professional development.
- Ensuring compliance of all activities with good practices (cGxP), data integrity and domestic and European law, and participation in internal and external audits.
- Contribution in the development, implementation and assurance of the quality system in accordance with Slovenian and European legislation, the FDA, ICH guidelines, the Inspection Convention and Novartis standards.
- Responsibility for equipment qualification and calibration and preparing an equipment qualification plan.

Essential Requirements:

- Bachelor's degree in pharmaceutical, biological, chemical, microbiological or other natural science direction.
- Minimum of 5 years of work experience in the field of quality, development or production, or other relevant field.
- Active knowledge of the English language.
- Proactivity, self-initiative, and quick comprehension of information.
- Experience in leading project groups and working with multiple stakeholders.
- Active, dynamic and communicative approach to work.
- Advanced use of Office tools.

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 (Wellbeing), 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
Operations

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
Quality

Job Type
Full time

Employment Type
Regular

Shift Work
No

Apply to Job

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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