

Ekspert upravljanja življenjske krivulje izdelka (m/ ž /d) / Lifecycle Implementation Expert (m/f/d)

Job ID
REQ-10048412

4月 30, 2025

Slovenia

摘要

The Life Cycle Implementation Expert (LCI) leads the implementation of Lifecycle (LC) projects, aligning and coordinating all stakeholders and functions involved in Lifecycle projects. The LCI ensures the timely and compliant implementation of Lifecycle projects (launches/ changes/ transfers/ divestment and pruning) in accordance to project plan and implementation strategy.

Join us and become our next talent.

About the Role

Key Responsibilities:

- Leads the implementation of LC projects at SKU material level, ensuring on-time and compliant first deliveries.
- Creates, maintains and communicates a detailed Change Over Plan (COP) for LC projects regarding implementation dates, according to the overall strategic project/ program plan, in order to allow for early local Master Data set-up and planning at site and in countries.
- Works with a broad variety of stakeholders: Actively seek alignment with Project Lead and with the experts from different functions (e.g. project team members, CPO DRA, CPO demand planning, site tactical/ operational schedulers, Master Data Governance, other Supply Chain functions).
- Is responsible for the up-to-date assortment at SKU level of the assigned projects. Initiates the Material creation in absence of Master Data Steward and provides candidates and evaluation to clean obsolete SKU.
- Tracks regulatory approvals and trigger implementation of changes in the Change Control system in accordance to regulatory status and implementation strategy.
- Ensures compliance with GMP, regulatory requirements, HSE (including record management) and continuous improvement of quality relevant processes within area of responsibility.
- Plans and tracks changeover activities and project implementation milestones with all involved functions and proactively addresses/resolves or escalates potential issues through the Project Lead or project team members (e.g. incomplete Master Data causing missing demand).
- Ensures compliant shipment to customer for artwork changes approved only for packaging.

Essential Requirements:

- Minimum university degree in Natural sciences, Technical sciences or Economics.
- At least 2 years experience in various pharmaceutical functions e.g. supply chain / production and / or technical development.
- Active knowledge of English.
- Knowledge of Microsoft Office.
- Highly motivated, independent and self-initiative.

We offer temporary 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), 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.

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For Slovenia:

Kot Ekspert upravljanja življenjske krivulje izdelka (m/ ž /ostre

Odgovorni za implementacijo projektov življenjskega cikla izdelka (LC), usklajevanje in koordinacija vseh deležnikov in funkcij, ki so vključeni v projekt življenjskega cikla. Zagotavljanje pravilnosti in z zakonodajo ter standardi skladne implementacije projektov življenjskega cikla (lansiranje / spremembe / transferji / odprodaje oz. zmanjševanje dejavnosti) v skladu s projektnim načrtom in strategijo izvajanja ter ob upoštevanju zakonodaje, GMP standardov, SOP, ZVO, kodeksom ravnanja ter ostalimi internimi predpisi, dobrimi praksami in poslovnimi cilji.

Pridružite se nam in postanite naš naslednji talent!

Vaše ključne odgovornosti:

- Vodenje implementacije projektov življenjskega cikla izdelka na materialnem nivoju (SKU), z namenom pravilnosti in ustrezne prvedobave.
- Kreira, vzdržuje in komunicira podrobni projektni plan sprememb in datumov implementacije (Change Over Plan - COP) v skladu s strategijo/planom projekta in tako omogoči pravilno kreiranje lokalnih matičnih podatkov, planiranje aktivnosti na proizvodnih lokacijah in aktivnosti v drugih zvezah.
- Sodeluje s tehniki udeleženimi deli življenjskega cikla in usklaja aktivnosti s projektnim vodjem in strokovnjaki iz različnih funkcij (npr. članji projektne skupine, CPO DRA, planerji povpraševanja, taktični / operativni planerji, skrbniki podatkov in zdrugimi funkcijami v oskrbi).
- Je odgovoren za posodobljen assortiman na materialni ravni (SKUs) dodeljenih projektov. Inicira ustvarjanje materiala v odsotnosti skrbnika matičnih podatkov in posreduje predloge in ocene za ukinitev zastarelih materialov (SKUs).
- Spremlja regulatorne odobrivate sprememb in spremlja izvajanje sprememb v sistemu za nadzor sprememb (Change Control System), v skladu z regulatornim statusom in strategijo izvajanja.
- Zagotavlja skladnost z GMP, regulatornimi zahtevami, ZVO (vključno z upravljanjem zapisov) in nenehno izboljševanje kakovosti procesov za svoje področje je odgovornosti.
- Načrtuje in spremlja projektni plan sprememb (Change Over Plan) in izvedbo zastavljenih mejnikov projekta z vsemi vključenimi funkcijami in proaktivno obravnava / rešuje ali eskalira morebitne težave prek vodja projekta ali članov projektne skupine (npr. nepopolni matični podatki, ki se kažejo v manjkajočem povpraševanju (demands)).
- Zagotavlja ustreznost odprem vezanih na pakiranje izdelka s spremembami v ovojnini / artworkih).
- Zastopanje vizije podjetja, vrednot in skrb za dobre medsebojne odnose s poslovnimi partnerji.
- Odgovornost za osebni in strokovni razvoj.
- Izvajanje in upoštevanje vseh navodil in zahtev za zagotavljanje varnega dela, varovanja okolja in premoženja.
- Ostale naloge določene z letnim pogovorom o ciljih in s kazalniki uspešnosti.

- Druge naloge po navodilu nadrejenega in naloge na podlagi posebnega imenovanja.

Vaš doprinos k delovnemu mestu:

- Vsaj visoko šolska stopnja izobrazbe naravoslovnotehnične akademije ali ekonomskodružboslovne smere.
- Minimalno 2 leta izkušenj na različnih funkcijah v farmaciji, pr. oskrba/proizvodnja in/ali tehnični razvoj.
- Aktivno znanje angleškega jezika.
- Poznavanje orodja Microsoft Office.
- Visoka motiviranost za delo, samostojnost in samoinicativnost.

Z izbranim kandidatom bomo sklenili delovno razmerje za določen čas poskusno doba 6 mesecev. Prijavo oddajte z življenjepisom slovenskem in angleškim jezikom.

Kaj nudimo:

Konkurenčna plača, npr. paket, letni bonus, fleksibilen način dela, možnost dostopanja do urnika in delom od doma, zaposlitev v podjetju s certifikatom TOP Employer, pokojninsko shemo, shemo nagrajevanja in priznanja dosežkov, širjeni program promocije zdravja na področju telesnega, duševnega in socialnega zdravja, družbenega in ekološkega razvoja. Polni življenjekater dogodki, neomejene priložnosti za učenje in razvoj.

Predani smo raznolikosti in vključenosti

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si prizadeva ustvariti izjemno, vključno in delovno okolje in oblikovanje raznolikih timov, saj ti predstavljajo naše bolnike in skupnosti, ki jih oskrbujemo.

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>

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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
Ljubljana

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

Functional Area
Technical Operations

Job Type
Full time

Employment Type
Temporary (Fixed Term)

Shift Work
No

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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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