NOVARTIS

Regulatory Associate Manager, Maintenance and CTA

Job ID
REQ-10091936

十一月 12, 2026

Spain

摘要

#LI-Hybrid

Location: Barcelona, Spain

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

Make an impact where science, strategy and execution come together. As Regulatory Associate Manager, Maintenance and CTA, you ’ ll help deliver high-quality regulatory activities across an assigned portfolio, supporting lifecycle management activities through to timely submission and approval. Working closely with regulatory leads and cross-functional teams, you ’ ll apply your regulatory knowledge to identify challenges, support health authority interactions and keep critical activities moving forward. You ’ ll also have the opportunity to contribute to continuous improvement and automation initiatives, helping us work smarter while delivering for patients.

About the Role

Key Responsibilities

- Deliver high-quality lifecycle management submissions in line with regulatory requirements.
- Build strong product and local regulatory knowledge to support assigned portfolio activities.
- Identify regulatory risks, gaps and trade-offs, helping minimise delays and enable timely approvals.
- Coordinate timely responses to health authority queries, supporting clear and robust regulatory justifications.
- Maintain accurate product information within regulatory compliance databases and information management systems.
- Support preparation and finalisation of documentation for health authority interactions.
- Contribute to continuous improvement, simplification and automation initiatives across Regulatory Affairs.

Essential Requirements

- Fluent English and Spanish, with strong written and verbal communication skills.
- Degree in a scientific or healthcare discipline, or equivalent relevant pharmaceutical industry expertise.
- Strong knowledge of regulatory requirements, guidelines and processes supporting lifecycle management and clinical trial applications.
- Experience supporting regulatory submissions, approvals, compliance and interactions with health authorities.
- Ability to interpret safety and efficacy data and understand product labelling requirements.
- Strong planning, organisational and communication skills, with the ability to work effectively across global, cross-functional teams.

Desirable Requirements

- Fluency in Portuguese.

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)

Primary location salary range
€ 36,100.00 - € 67,100.00

部门
Development

Business Unit
Development

地点
Spain

站点
Barcelona Gran V í a

Company / Legal Entity
ES06 (FCRS = ES006) Novartis Farmac é utica, S.A.

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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(kPlayer.plugins["share"]) { config.plugins.share = { disable: true }; } config.ui.uiComponents = []; if (kPlayer.plugins["googleAnalytics"]) { config.plugins.googleTagManager = {}; config.plugins.googleTagManager.customEventsTracking = {}; config.plugins.googleTagManager.containerId = 'GTM-57RJQ5'; config.plugins.googleTagManager.customEventsTracking.custom = []; config.plugins.googleTagManager.customEventsTracking = { preset: { coreEvents: true, UIEvents: false, playlistEvents: false, castEvents: false } }; }

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