

Snr Specialist, IT Business Analyst, Technical Design (Regulatory Affairs)

Job ID

REQ-10018835

Aug 13, 2024

Spain

About the Role

Role Responsibilities:

- Engage with global business associates and leverage the appropriate teams and functions to determine requirements and deliver data-driven recommendations to improve efficiency and add value.
- Analyses the business domain and author business requirements. Coordinate and facilitate ongoing reviews of business processes.
- Ensure consistency and traceability between user requirements, functional specifications, and testing & validation. Support the validation and testing as appropriate.
- Flag issues/ changes/ risks to the Project Manager and workstream leads. Work together with a product squad in delivering the Product's roadmap. Actively participate in sprint planning discussions and ensure sprint functional deliverables (prioritized backlog, user stories completed and demonstrated etc.) are on track.
- Liaise with vendor, Novartis internal IT teams and business to ensure documentation is at the appropriate level of details and that the requirements are accurately interpreted and implemented.
- Act as interface between business and Implementation partners. Review the sprint demos and ensure that gaps are documented.

Role Requirements:

- Bachelor's degree in engineering or pharmaceutical discipline. An advanced degree (MBA, MS etc.) and related accreditations (IIBA, Veeva, Agile certifications etc.) is a plus.
- 7+ years of IT Business Analysis experience with excellent communication skills.
- Must have proven strong knowledge of SDLC, Validation & Compliance
- Proficiency with tools such as Jira, Confluence, HPQC, Business process modelling tools
- Experience in Data migration and System integration related projects.
- Multi-national global experience in interacting with senior management, collaborating across boundaries and relationship management, and influencing without authority.
- Experience in Regulatory Affairs business processes is a plus (e.g. Registration Management, Submission Management, Submission Content management, Submission Publishing & Clinical Publishing, Product Labelling)

Desirable:

- Implementation experience of Veeva Submission and Submission Archive module is a plus.
- Experience in Managing GxP Projects and Related Fields is a plus

Language: English

Benefits & Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: [Novartis Life Handbook](#)

Benefits in Spain include Company Pension plan; Life and Accidental Insurance; Meals; Allowance or Canteen in the office; Flexible working hours.

You'll receive: You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook. <https://www.novartis.com/careers/benefits-rewards>

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Commitment to Diversity & Inclusion:

We are committed to building an outstanding, inclusive work environment and diverse teams representative of the patients and communities we serve.

This role is based in Barcelona, Spain. Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you

Role Requirements

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>

Division

Operations

Business Unit

CTS

Location

Spain

Site

Barcelona Gran Vía

Company / Legal Entity

ES06 (FCRS = ES006) Novartis Farmacéutica, S.A.

Functional Area

Technology Transformation

Job Type

Full time

Employment Type

Regular

Shift Work

No

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