

IT Solution Design Expert (Regulatory Affairs)

Job ID
REQ-10018836
Nov 19, 2024
Spain

About the Role

Major accountabilities:

- Design and manage the development of modular, reusable, elegantly designed and maintainable software or solutions that supports the RA organization as well as other Cross Functional strategic initiatives.
- Participate fit-gap workshops with business providing effort estimates and solutions proposals.
- Hands on, solution driven, and customer focused. Develop and maintain the code repositories for which you are responsible and respond rapidly to bug reports or security vulnerability issues.
- Embody and integrate software development best practices into your everyday work and inspire others within the engineering community to emulate these practices.
- Collaborate extensively with your colleagues across Security, Compliance, Engineering, Project Management, Product Management, Product Service Management and Business Management.
- Continue support for moving Publishing other RA platforms to the cloud.
- Ensure delivered solutions adhere to architectural and development standards, best practices, and meet requirements as recommended in the architecture handbook.
- Ensure designed solutions are aligned with Data and Analytics strategy standards and roadmap. Bring innovations to the solutions and add measurable values to RA Business.

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.

Minimum Requirements:

- University degree in Information Technology, Computer Sciences, Life Sciences or similar.
- 10+ years of experience in IT technology development experience, preferably with 5+ years of experience working with Pharma Regulatory affairs (RA) business.
- Experience working in with RA Publishing business is a plus.
- Strong technical background with large scale projects on technology like JAVA/.net, Oracle, MS SQL, IIS webserver, PowerShell or similar end to end technical solution delivery.
- Experience on Publishing solutions like TRS, EFT and DMZ Gateway is preferred but not mandatory.
- Familiarity with concepts of data and system security and compliance in highly regulated environments

Languages :

- English.

Why Novartis: Our purpose is to reimagine medicine to improve and extend people's lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <https://www.novartis.com/about/strategy/people-and-culture>

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>

Join our Novartis Network: If this role is not suitable to your experience or career goals but you wish to stay connected to hear more about Novartis and our career opportunities, join the Novartis Network here: <https://talentnetwork.novartis.com/network>

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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