Senior Specialist IT MES

Job ID
REQ-10067292

11月 25, 2025

Spain

摘要

Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

Location: Barcelona, Hyderabad or Ljubljana
#LI-Hybrid (12 days/month in office)

Are you looking for an exciting opportunity to work with the latest technologies, collaborate with a top-performing team, and be surrounded by highly skilled professionals? We are seeking a talented and motivated associate to join our Novartis Manufacturing IT team.

About the Role

Key Responsibilities:

- **System Installation and Configuration:** Perform the installation and configuration of the MES software, working closely with the vendor and IT teams. Ensure proper integration with existing systems and data sources and troubleshoot any technical issues that may arise during the installation process.
- **Validation:** Develop and execute validation protocols (e.g. Installation Qualification, Operational and Performance Qualification) to verify the functionality and compliance of our MES with regulatory standards. Your meticulous approach will ensure that our system meets all necessary requirements. In collaboration with business stakeholders, you will ensure that overall validation lifecycle is in place and maintained.
- **Vendor Collaboration:** Collaborate with our trusted vendor to gather user requirements and ensure the system's capabilities align with the organization's manufacturing processes. Provide input on system design, assess technical feasibility, and manage the procurement process in coordination with the vendor.
- **Integration and Digitalization**: Integrate our solutions into Novartis ecosystem. You will be responsible for end-to-end integration between the system ensuring that relevant data is exchanged and available per business requirements. MES plays an important role on our digital journey hence you will have a chance to impact how we transform our ways of working.
- **Continuous Improvement**: Continuously monitor and evaluate the performance of our MES to identify areas for optimization. Work closely with business stakeholders on translation of business requirements into a technical solution. You will be part of the business community where you will have a chance to work with different sites worldwide.
- **Successful Implementation**: Ensure the successful implementation of the Manufacturing Execution System within project timelines, budget, and quality standards. This includes completing all necessary tasks, such as requirements gathering, server provisioning, installation, and configuration, while adhering to project goals and delivering the system on time and within budget.
- **Compliance with Validation Protocols and Regulatory Requirements**: Validate the functionality and compliance of the MES system by executing validation protocols and ensuring adherence to regulatory standards. The absence of critical or major observations during internal and external inspections indicates a successful compliance record.
- **Stable Operations and System Uptime**: Maintain stable operations of the MES system with minimal downtime and interruptions. Measure and track system uptime to ensure optimal availability and performance, meeting the defined Key Performance Indicators (KPIs) as per the Service Level Agreement.
- **User Satisfaction**: Gather feedback from end-users to evaluate their satisfaction with the MES system. Conduct regular surveys or interviews to assess the usability, effectiveness, and user experience of the system. A high level of user satisfaction indicates a successful implementation that meets the needs and expectations of the users.

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.

Essential Requirements:

- Bachelor's degree in a computer science, engineering or information technology discipline. An advanced degree and related accreditations a plus.

- Experience in GxP environment and understanding of pharmaceutical manufacturing processes and technologies are an advantage but not essential.
- Fluent English (written and verbal)
- Experience in MES systems.
- Digital and Tech savvy

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

Accessibility and accommodation:
Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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: 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
Information Technology

地点
Spain

站点
Barcelona Gran V í a

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

Alternative Location 1
Hyderabad (Office), India

Alternative Location 2
Ljubljana, Slovenia

Functional Area
Technology Transformation

Job Type
Full time

Employment Type
Regular

Shift Work
No

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