NOVARTIS

Sr Manufacturing Technical MES Specialist

Job ID
REQ-10087150
Сен. 16, 2026
США

Сводка

Location: Morris Plains, New Jersey

Imagine helping bring life-changing cell and gene therapies to patients by transforming how manufacturing operations are executed and documented. As a Sr Manufacturing Technical MES Specialist at Novartis, you will play a critical role in advancing digital manufacturing capabilities through the design, implementation, and optimization of Manufacturing Execution Systems. Working closely with manufacturing, quality, supply chain, and technical teams, you will drive paperless manufacturing solutions, enhance operational efficiency, and ensure the compliant execution of complex production processes. This is an opportunity to shape the future of advanced therapy manufacturing while making a meaningful impact on patients worldwide.

About the Role

Key Responsibilities

- Own end-to-end Manufacturing Execution System processes supporting commercial, clinical, and future manufacturing programs.
- Design, configure, test, implement, and continuously improve electronic Master Batch Records and manufacturing workflows.
- Provide Manufacturing Execution System production support and resolve system issues to ensure uninterrupted operations.
- Collaborate with manufacturing, supply chain, quality, technical development, and validation teams to optimize processes.
- Manage Manufacturing Execution System updates, change controls, corrective actions, and documentation revisions to maintain compliance.
- Support system integrations and master data management across enterprise platforms, including laboratory and enterprise resource planning systems.
- Coach and train key users and subject matter experts on batch record design, execution, and best practices.

Essential Requirements

- Bachelor’s degree or higher, or equivalent relevant work experience.
- Minimum 2 to 5 years of Manufacturing Execution System experience, including Master Batch Record design and creation.
- 5 years of experience in a regulated current Good Manufacturing Practice environment, preferred.
- Demonstrated expertise in technical Manufacturing Execution System Master Batch Record design, testing, implementation, and maintenance.
- Knowledge of Manufacturing Execution System interfaces and integrations, including enterprise resource planning and laboratory information management systems.
- Strong business requirements gathering, project management, communication, and documentation skills with exceptional attention to detail.

The salary for this position is expected to range between $98,700 and $183,300 per year. The final salary offered is determined based on factors like, but not limited to, relevant skills and experience, and upon joining Novartis will be reviewed periodically. Novartis may change the published salary range based on company and market factors. Your compensation will include a performance-based cash incentive and, depending on the level of the role, eligibility to be considered for annual equity awards. US-based eligible employees will receive a comprehensive benefits package that includes health, life and disability benefits, a 401(k) with company contribution and match, and a variety of other benefits. In addition, employees are eligible for a generous time off package including vacation, personal days, holidays and other leaves.

To learn more about the culture, rewards and benefits we offer our people click here.

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)

EEO Statement:

The Novartis Group of Companies are Equal Opportunity Employers. We do not discriminate in recruitment, hiring, training, promotion or other employment practices for reasons of race, color, religion, sex, national origin, age, sexual orientation, gender identity or expression, marital or veteran status, disability, or any other legally protected status.

Accessibility & Reasonable Accommodations

The Novartis Group of Companies are 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 application process, or to perform the essential functions of a position, please send an e-mail to us.reasonableaccommodations@novartis.com or call +1(877)395-2339 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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Operations
Business Unit
Production / Manufacturing
Место
США
Состояние
New Jersey

Сайт
Morris Plains
Company / Legal Entity
U014 (FCRS = US014) Novartis Pharmaceuticals Corporation
Functional Area
Technical Operations
Job Type
Full time
Employment Type
Regular
Shift Work
No

Job ID
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Sr Manufacturing Technical MES Specialist

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List of links present in page

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