NOVARTIS

Product Steward

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REQ-10081689
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США
Available in: English

Сводка

Location: Morrisville, North Carolina

Are you passionate about turning data into insights that improve product quality and manufacturing performance? Join Novartis as a Product Steward and become the technical expert responsible for maintaining deep product and process knowledge throughout the commercial lifecycle. In this highly visible role, you will use data analytics, process monitoring, and scientific expertise to ensure products remain in a state of control, drive continuous improvement initiatives, and support robust, reliable manufacturing operations. Working across multiple functions, you will play a key role in product stewardship, process validation, technical investigations, and operational excellence while helping deliver high-quality medicines to patients.

About the Role

Key Responsibilities

- Serve as the product subject matter expert throughout the commercial manufacturing lifecycle.
- Monitor process performance through data trending, statistical analysis, and ongoing product oversight.
- Lead technical investigations and drive effective corrective and preventive actions for process-related issues.
- Maintain product-specific quality risk assessments and support robust process control strategies.
- Ensure products remain in a continued state of validation through ongoing process verification activities.
- Partner with cross-functional teams to deliver process improvements, optimization initiatives, and manufacturing excellence projects.
- Support regulatory readiness, product transfers, and technical change implementation for assigned products.

Essential Requirements

- Bachelor’s degree in Pharmacy, Pharmaceutical Technology, Chemistry, or equivalent scientific degree.
- Minimum five years of experience in Oral Solid Dosage pharmaceutical manufacturing environments.
- Strong understanding of pharmaceutical manufacturing processes, regulatory requirements, and GMPs.
- Experience using data trending, statistical analysis, and process monitoring to drive decision-making.
- Proven ability to investigate process issues, identify root causes, and implement effective corrective actions.
- Demonstrated experience working collaboratively across technical, quality, and manufacturing functions.

Desirable Requirements

- Master of Science degree in Pharmacy, Pharmaceutical Technology, Chemistry, or a related scientific field.
- Experience supporting product transfers, process optimization initiatives, and manufacturing excellence programs.

The salary for this position is expected to range between $126,000 and $234,000 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 clickhere.

#LI-Onsite

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

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Production / Manufacturing
Место
США
Состояние
North Carolina
Сайт
Durham
Company / Legal Entity
U473 (FCRS = US473) Novartis Gene Therapies
Functional Area
Technical Operations
Job Type
Full time
Employment Type
Regular
Shift Work
No

Job ID
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Product Steward

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