

Product Steward

Job ID

REQ-10081689

Jul 05, 2026

LOC_US

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 click [here](#).

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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>

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.

[Read our handbook \(PDF 30 MB\)](#)

Division

DIV_TO

Business Unit

Production / Manufacturing

Location

LOC_US

Site

Durham

Company / Legal Entity

U473 (FCRS = US473) Novartis Gene Therapies

Functional Area

FCT_TO

Job Type

Full time

Employment Type

Regular

Shift Work

No

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