

## QA Operations Head (Associate Director)

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REQ-10083721  
июл 21, 2026  
США  
Available in: English

### Сводка

Join Novartis as our QA Operations Head and play a pivotal role in ensuring patients receive high-quality medicines that meet the highest standards of safety, compliance, and reliability. This is an opportunity to lead a critical Quality Operations organization at our new small molecule manufacturing site in Morrisville, partnering closely with Manufacturing, Quality Control, and technical teams to drive operational excellence, product quality, and regulatory compliance. As a visible site leader, you will empower and develop high-performing teams, champion continuous improvement, and help shape the future of quality within a dynamic and growing manufacturing environment.

### About the Role

#### Key Responsibilities

- Lead Quality Operations oversight to ensure compliance across manufacturing activities and QA Operational Excellence.
- Provide quality leadership for product release, batch record review, and disposition activities.
- Drive investigation of deviations, out-of-expectation results, and complaints, ensuring effective corrective actions.
- Lead continuous improvement initiatives to strengthen quality performance, compliance, and operational excellence.
- Support validation, technology transfer, commissioning, and implementation of new manufacturing processes and equipment.
- Ensure ongoing inspection readiness and successfully support internal, customer, and regulatory audits and inspections.
- Develop, coach, and lead high-performing teams while fostering a culture of accountability and learning.
- Manage resources, capacity planning, and quality metrics to achieve business and compliance objectives.
- Explore the potential of data and technology to support the manufacturing site's transformation journey in line with Novartis Technical Operations strategy.

#### Essential Requirements

- Bachelor of Science or Master of Science in Life Sciences, Pharmacy, Chemistry, Biotechnology, or related scientific discipline; advanced degree preferred.
- Minimum 8-10 years of relevant pharmaceutical experience including direct experience in oral solid dosage manufacturing which can include expert roles in Production, Manufacturing Science and Technology and/or Engineering, in addition to Quality.
- Demonstrated Quality Operations leadership experience with responsibility for product release, compliance, inspection readiness, people leadership, talent development, performance management, and cross-functional collaboration.
- Experience with Health Authority inspections (preferably FDA) and regulatory procedures
- Proven experience leading deviation investigations, corrective and preventive actions, validation activities, and continuous improvement initiatives.

#### Desirable Requirements

- Previous involvement with facility start-ups, qualifications, validations, transfers
- Qualifications or certification in Lean Management, Operational Excellence, or continuous improvement methodologies.

The salary for this position is expected to range between \$138,600 and \$257,400 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](#).

### #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:

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### 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](mailto: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.

Operations  
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Место  
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Company / Legal Entity  
U473 (FCRS = US473) Novartis Gene Therapies  
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Quality  
Job Type  
Full time  
Employment Type  
Regular  
Shift Work  
No

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