

## Site Quality Head, Florida (AD level)

Job ID  
REQ-10072294  
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CША

### Сводка

Job Title: Site Quality Head (Associate Director level)

#LI-Onsite  
Location: Winter Park, Florida

At Novartis, we are redefining the future of cancer care through Radioligand Therapy (RLT) - a powerful convergence of nuclear medicine and precision oncology. As we expand our global RLT manufacturing footprint, we are building a team of passionate, purpose driven quality leaders who are inspired to make a meaningful impact. This is a unique opportunity to play a critical role in the start-up and ongoing quality operations of a new RLT manufacturing site, helping ensure the delivery of life changing therapies to patients around the world.

As the Site Quality Head in Winter Park, Florida, you'll shape the quality foundation for a new Radioligand Therapy manufacturing site—combining rigorous compliance with strong, visible leadership to help deliver time-sensitive therapies reliably and safely. You'll lead and develop a high-performing Quality organization, partner closely with site and global stakeholders, and set the standard for inspection readiness, robust management of product quality issues, and continuous improvement—so the site starts up strong and scales with confidence.

### About the Role

#### Key Responsibilities

- Lead site quality strategy and governance to meet Novartis standards and current Good Manufacturing Practice.
- Build, coach, and develop the Quality team, strengthening capability, engagement, and safe working practices.
- Drive plant startup, expansions, and technology transfer, ensuring compliant planning, commissioning, qualification, and validation activities.
- Lead inspection readiness and represent the site during health authority, corporate, and internal audits.
- Oversee deviations, investigations, out-of-specification events, and corrective and preventive actions through effective closure.
- Partner with Manufacturing and cross-functional leaders to enable compliant, efficient operations and risk-based decision making.
- Define and monitor site quality performance indicators, driving continuous improvement and timely escalation of risks.

#### Essential Requirements

- Bachelor's degree in life sciences or a related scientific discipline.
- Ten years of experience in a GMP pharmaceutical manufacturing environment, including laboratory operations and Aseptic experience, and at least three years of combined relevant experience in Quality Assurance and/or Quality Control roles.
- In-depth knowledge of cGMP and United States Food and Drug Administration regulations and International Council for Harmonization regulations. Understanding of US Pharmacopeia, European Pharmacopeia, and American Chemical Society standards.
- Proven success leading health authority inspections and delivering robust remediation and sustained compliance improvements.
- Demonstrated leadership in matrix organizations with excellent communication, organizational, and stakeholder management skills.
- Experience applying continuous improvement methods such as Lean Six Sigma, Total Quality Management, and 5S workplace organization.

#### Desirable Requirements

- Prior experience with site start-up or rapid site expansion
- Experience or training in Radioligand Therapies, radiopharmaceuticals and/or radiation safety.

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](#).

**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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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.

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Production / Manufacturing  
Место  
США  
Состояние  
Florida  
Сайт  
Winter Park (Florida)  
Company / Legal Entity  
U473 (FCRS = US473) Novartis Gene Therapies  
Functional Area  
Quality  
Job Type  
Full time  
Employment Type  
Regular  
Shift Work  
No

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