

# **Associate Director, Manufacturing Operations**

Job ID REQ-10030322 Nov 18, 2024 USA

#### **Sommario**

Location: Morris Plains, NJ #LI-Onsite

#### About this role:

The Associate Director of Manufacturing Operations for Morris Plains Cell and Gene Therapy is responsible for the overall operations within the manufacturing area operating over multiple shifts 24/7/365. The Associate Director provides direction, leadership, and guidance to support roles that directly affect daily operations which are expected to produce and deliver product with high quality in a safe, compliant, efficient, and cost effective manner.

#### **About the Role**

#### **Key Responsibilities:**

- Accountable for all aspects of staffing, including recruiting talent, and strategic/succession planning.
- Ensures adequate resources and cross training to meet the demands of a multi product processing facility.
- Supports the operations site strategy and ensure tactics are aligned with strategy
- Contribute to site financial and business goals
- Performs prioritization of core operations, continuous improvements of projects
- Designs and optimizes Manufacturing process to meet demands of multiple products
- Serves as escalation point for Manufacturing issues that occur on weekdays and weekends
- Collaborates with all support functions to ensure that the production plan is met
- Supports process improvement initiatives and ensures that Operational Excellence is embedded in the team.
- Represents manufacturing during Health Authority inspections

#### **Essential Requirements:**

- Bachelor's Degree in Biotechnology, Biopharmaceutical, Pharmaceutical Technology, Chemistry, Microbiology, or equivalent required. Advanced Degree preferred.
- 8+ years' experience in cGMP required, with aseptic and cell therapy manufacturing highly desirable
- 8+ years' direct management experience
- Demonstrated experience leading large multi levels teams (shop floor leaders)
- Project management, Lean, Operational Excellence, Product/Process Development or Regulatory experience a plus

#### **Commitment to Diversity and Inclusion:**

Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.

**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? <a href="https://www.novartis.com/about/strategy/people-and-culture">https://www.novartis.com/about/strategy/people-and-culture</a>

**Join our Novartis Network:** Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: <a href="https://talentnetwork.novartis.com/network">https://talentnetwork.novartis.com/network</a>

**Benefits and Rewards:** Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <a href="https://www.novartis.com/careers/benefits-rewards">https://www.novartis.com/careers/benefits-rewards</a>

#### **EEO Statement:**

The Novartis Group of Companies are Equal Opportunity Employers who are focused on building and advancing a culture of inclusion that values and celebrates individual differences, uniqueness, backgrounds and perspectives. 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. We are committed to fostering a diverse and inclusive workplace that reflects the world around us and connects us to the patients, customers and communities we serve.

### **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 <u>us.reasonableaccommodations@novartis.com</u> 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.

Divisione

Operations

**Business Unit** 

Innovative Medicines

Posizione

USA

Sito

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 Apply to Job

Job ID REQ-10030322

# **Associate Director, Manufacturing Operations**

# Apply to Job

**Source URL:** https://www.adacap.com/careers/career-search/job/details/req-10030322-associate-director-manufacturing-operations

# List of links present in page

- 1. https://www.novartis.com/about/strategy/people-and-culture
- 2. https://talentnetwork.novartis.com/network
- 3. https://www.novartis.com/careers/benefits-rewards
- 4. mailto:us.reasonableaccommodations@novartis.com
- 5. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\_Careers/job/Morris-Plains/Associate-Director--Manufacturing-Operations\_REQ-10030322