NOVARTIS

Associate Expert Drug Supply

Job ID
REQ-10090772
Oct 29, 2026
LOC_IT

About the Role

Major accountabilities:

- Work in shifts, is responsible to perform the necessary activities for the preparation, production and packaging for shipping of sterile radioactive drugs
- Execute all assigned activities in compliance with internal procedures (HSE, EU GMP Annex 1 / Annex 3, others) and in compliance with applicable laws and regulations
- Perform routine setup and cleaning, microbiological verification of pharmaceutical areas and equipment and all areas of the Plant with a need for periodic cleaning activities based on GMP
- Perform the preparation of all raw materials needed for the production (batch related kit preparation)
- According with the Production Supervisor, perform the management of material stocks and waste materials
- Follow the GDP in printing, filling and signing all documents (ALCOA+)
- Promptly report to the Production Supervisor and the Qualified Person and/or the HSE Manager any deviation and/ or out of specification detected during the production process
- Perform logistics activities (transportation and documentation related) about the incoming of materials, refilling of the stock areas and packaging materials for shipping.
- Perform activities related to waste management (transportation and documentation related) for radioactive waste
- Support external specialized personnel in carrying out ordinary and extraordinary maintenance/qualification activities
- Collaborate with the Production Supervisor to the training of new personnel
- Support the site quality system by providing input from the production area and sustaining the continuous improvement of the facility
- Observe all safety requirements (D.Lgs. 81/08; D.lgs 101/95, Global and local HSE SOPs), GMP regulations (AIFA, FDA), and housekeeping practices while performing duties.
- Provides continuous feedback and is the point of contact for generating improvements and changes aimed at optimizing the quality and efficiency of production processes.
- Responsible for performing technical tests and supporting the qualification and validation phases of processes and products.
- undefined

Education & Qualifications

Technical/Scientific background/education (High School) is mandatory

University degree in Science is desirable, Pharmacy or Chemical Engineering, Pharmaceutical

Technology or equivalent job experience.

Languages

- English: desirable
- Italian (Local Language): proficient

Experiences

- Minimum 2 years experience in GMP manufacturing support / technical role
- Previous experience in sterile injectable pharmaceutical companies (radiopharmaceutical experience is desirable)

Competencies

- Good scientific and technical understanding
- Team player with strong team spirit
- Change management, adaptability, ability to work under pressure
- Good understanding or capacity to quickly understand production processes
- Quality and compliance skills
- Good understanding of regulatory requirements across multiple health authorities

Technical / Functional Skills & Knowledge

- Good working knowledge/understanding of typical manufacturing IT system (SCADA, HMI etc)
- Good office software applications

Benefits & Rewards

At Novartis, we’re committed to reimagining medicine together - and rewarding the people who make it happen.

Expected Annual Base Salary Range for role: €30,200 to €56,000

The base salary offered is determined based on gender-neutral objectives, such as relevant skills, competencies and experience in accordance with the Novartis pay setting policy and upon joining Novartis will be reviewed periodically.

In addition to your base salary, you may be eligible for a performance-based bonus depending on certain performance parameters.

The rewards of being part of our team go far beyond base pay and incentives. We also offer a variety of competitive benefits in kind to help you thrive personally and professionally, such as insurance plans, retirement plans, wellbeing resources and global recognition programs. In addition, we provide flexible and hybrid working options, where possible, and minimum 14 weeks paid parental leave.

Pay equity is a fundamental principle of our employment policy and reflects our commitment to create a diverse, equitable and inclusive environment that treats all employees with dignity and respect, as outlined in our Code of Ethics.

Read our brochure to learn more about our global total rewards offering:

https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf

Note: Benefits and compensation may vary by country and are subject to local legal requirements, including provisions of collective bargaining agreements where applicable. A full overview of your compensation package, including any relevant collective bargaining agreement details applicable to your role based on your employment location and Novartis employer entity, will be communicated separately to you during the application process.

Commitment to Diversity and Inclusion / EEO paragraph

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

Role Requirements

Primary location salary range
€30,200.00 - €56,000.00
Division
DIV_GD
Business Unit
Development
Location
LOC_IT
Site
Ivrea
Company / Legal Entity
IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl
Functional Area
FCT_RD
Job Type
Full time

Employment Type
Regular
Shift Work
No
Apply to Job

Job ID
REQ-10090772

Associate Expert Drug Supply

Apply to Job

Source URL: *https://jobapi.novartis.com/req-10090772-associate-expert-drug-supply*

List of links present in page

1. https://jobapi.novartis.com/req-10090772-associate-expert-drug-supply
2. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
3. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Ivrea/Associate-Expert-Drug-Supply_REQ-10090772
4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Ivrea/Associate-Expert-Drug-Supply_REQ-10090772