

## QC Sample Management Technician (Wed - Sat AM)

Job ID  
REQ-10040065

2月 19, 2025

USA

### 摘要

The QC Sample Management Technician reporting to the QC Supervisor - Sample Management will support lab operations by performing routine sample management to support Quality Control (QC) Analytical laboratories

### About the Role

Major accountabilities:

### Position Summary

- Support lab operations by performing routine QC in-process and release test sample and QC retain sample management to support Quality Control (QC) Analytical laboratories.

## Core Responsibilities

- QC sample / retains sample handling including receipt, registration, storage, distribution and destruction
- Delivering samples to QC laboratories for testing as required
- Preparation of samples for shipment
- Performing sample inventory and disposal
- Ensure samples and materials are appropriately labelled and maintain sample tracking, labelling and chain of custody records in accordance with regulatory requirements and written procedures.
- Maintain the Sample Retain inventory room, refrigerators, and freezers.
- Ensure that Sample Retains are stored in appropriate locations.
- Ensures adequate storage areas are available in each sample storage area.
- Ensure Sample Retains are readily available for Shipping with proper documentation. Prepare for shipping testing to contract labs for chemistry and microbiology samples.
- Dispose of Sample Retains in accordance with cGMP regulations after retention time has passed.
- Organize Sample Retains by lot/batch number, documenting the storage location in logbooks/excel for ease of retrieval.
- Performs other related duties as assigned.

## Continuous Improvement

- Drive a culture of continuous improvement and proactively identify opportunities for efficiency enhancements and process improvements within the Quality Control procedures.

## Collaboration

- Interacting with outside customers and functional peer groups
- Communicate effectively with management regarding task completion, roadblocks, and needs.
- Support investigation of sample management-related issues: Work together with QA and Investigation/Compliance personnel and with stakeholders on addressing any non-compliance or sample management-related deviations, implementing corrective actions, and documenting findings.

## Compliance

- Ensure that sample management activities are completed in compliance with all applicable procedures, standards and GMP regulations.

## Key Performance Indicators

- Timely delivery on commitments with and departmental KPIs
- Timely responses and solutions to training-related issues
- Efficient and flexible usage of the available resources

- Compliance to all relevant company policies and guidelines
- Exhibiting core Novartis values and behaviors and fostering these within the team

## Ideal Background

### Education:

- HS diploma, Associate 's degree, Biotechnology certificate, or Bachelor 's degree in biology, chemistry, or other related degree concentration

### Languages:

- English

### Experience:

- 0-2 years of experience in a cGMP environment / commercial manufacturing.
- Experience in a regulated GMP environment, preferably within QC operations in a cell therapy company.

### Skills:

- Experience with handling Liquid Nitrogen (LN2) samples
- Experience with use and routine maintenance of sample handling equipment including BSCs, refrigerators, freezers and LN2 storage tanks.
- Experience with pipetting and aliquoting
- Experience with electronic laboratory management systems (such as LIMS)

## Competency Profile

- Strong communication skills for interacting with a variety of collaborators and partner groups
- Ability to work independently and as part of a team, self-motivation, adaptability, and a positive attitude
- Must demonstrate flexibility in adjusting to changing priorities and schedules.
- Ability to learn new techniques, perform multiple tasks simultaneously, keep accurate records, follow instructions, and comply with company policies
- Ability to function in a rapidly changing environment & handle multiple priorities.
- Ability to collaborate cross functionally to drive operational and quality excellence.
- Advanced organizational and time management skills.
- Communicate effectively with management regarding task completion, roadblocks, and needs.
- Customer service mindset
- Continuous improvement mind frame
- Highly organized with an attention to detail

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U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area  
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Job Type  
Full time

Employment Type  
Regular

Shift Work  
No

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