

# Sterility Assurance Expert

Job ID

REQ-10075576

Apr 27, 2026

LOC\_US

## About the Role

### Key Responsibilities

- Serve as site sterility assurance expert for aseptic processing and microbial control programs
- Design, maintain, and improve sterility assurance strategies aligned with global regulatory expectations
- Lead sterility-related deviations, contamination investigations, and risk-based decision making
- Own environmental and personnel monitoring strategies, including trending, escalation, and continuous improvement
- Author, review, and approve sterility assurance documentation, validations, and technical rationales
- Represent sterility assurance during regulatory inspections, audits, and quality governance forums
- Partner cross-functionally to resolve sterility risks and strengthen inspection readiness and compliance

### Essential Requirements

- Bachelor's degree in a scientific or technical field, preferably Microbiology or a related discipline
- Extensive experience supporting sterility assurance programs in regulated biopharmaceutical manufacturing environments
- Deep understanding of aseptic processing, microbial control strategies, and contamination risk management
- Proven experience leading sterility-related investigations, environmental monitoring programs, and corrective action planning
- Strong knowledge of global regulatory expectations, including United States and European health authority requirements
- Ability to apply scientific judgment, analyze complex data, and communicate clear, compliant recommendations

### Desirable Requirements

- Experience supporting sterility assurance activities for aseptic cell or gene therapy manufacturing
- Direct participation in regulatory inspections with ownership of sterility assurance topics and responses

The pay range for this position at commencement of employment is expected to be between \$108,500 and \$201,500 per year; however, while salary ranges are effective for a defined period, fluctuations in the job market may necessitate adjustments. Final pay determinations will depend on a variety of factors, including but not limited to geographic location, experience level, knowledge, skills, and abilities. The total compensation package may also include other elements, such as a performance-based bonus and a full range of medical, financial, and other benefits, including retirement programs and paid time off. Details of participation in these benefit plans will be provided if an offer of employment is made. Employment with Novartis is at-will, and the company reserves the right to modify compensation at any time based on individual, business, or market factors.

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## Role Requirements

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Division

DIV\_TO

Business Unit

Quality

Location

LOC\_US

Site

Morris Plains

Company / Legal Entity

U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area

FCT\_QA

Job Type

Full time

Employment Type

Regular

Shift Work

No

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