

# Manufacturing Specialist

Job ID

REQ-10076161

May 14, 2026

LOC\_MX

## About the Role

### Key Responsibilities

- Open and assess deviations, determining criticality within defined timelines
- Evaluate product impact of deviations in alignment with batch release activities
- Author and own investigations, ensuring clear scope, root cause, and timely closure
- Apply structured root cause analysis tools to identify product and process deviations
- Develop, document, and implement effective Corrective and Preventive Actions
- Verify robustness and effectiveness of critical and major investigations
- Execute experiments or manufacturing runs to support investigation outcomes
- Collaborate cross-functionally to ensure compliant production during deviations
- Generate manufacturing orders within the Manufacturing Execution System as required
- Deliver targeted training to reinforce quality practices and compliance standards

### Essential Requirements

- Bachelor's degree in a scientific field, with two to five years of pharmaceutical industry experience
- Proven experience working in a Good Manufacturing Practice production environment, preferably aseptic or sterile
- Demonstrated experience leading deviation investigations and corrective and preventive action activities
- Strong knowledge of current Good Manufacturing Practice regulations and regulatory expectations for pharmaceutical manufacturing
- Experience applying structured root cause analysis methods to product and process issues
- Fluent English and Spanish communication skills, both written and spoken

This role includes participation in a rotating shift model, which involves work on weekends for part of the schedule, with overall working time remaining five days per week and details agreed in advance.

### Commitment to Diversity and Inclusion

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

### Accessibility and accommodation

Novartis is committed to work with and provide reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to perform the essential functions of a position, please send an e-mail to [tas.mexico@novartis.com](mailto:tas.mexico@novartis.com) and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

## Role Requirements

**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>

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[Read our handbook \(PDF 30 MB\)](#)

Division

DIV\_TO

Business Unit

Production / Manufacturing

Location

LOC\_MX

Site

INSURGENTES

Company / Legal Entity

MX06 (FCRS = MX006) Novartis Farmacéutica S.A. de C.V.

Functional Area

FCT\_TO

Job Type

Full time

Employment Type

Regular

Shift Work

No

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