

Manufacturing Specialist

Job ID
REQ-10076161
май 14, 2026
Мексика

Сводка

#LI-Hybrid

Location: Mexico City, Mexico

Relocation Support: This role is based in Mexico City, Mexico. Novartis is unable to offer relocation support: please only apply if accessible.

When something unexpected happens in manufacturing, you'll be the person who brings clarity, structure, and momentum - so production can continue compliantly and patients aren't kept waiting. In this role, you'll lead deviation and investigation work end-to-end: assessing criticality and product impact, authoring robust investigations, and driving root-cause analysis using practical investigation tools. You'll translate findings into effective Corrective and Preventive Actions, ensure implementation through Good Manufacturing Practice systems, and verify that actions truly prevent recurrence. You'll also support the operation by generating manufacturing orders in the Manufacturing Execution System when needed and delivering targeted training that reinforces strong quality behaviors.

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 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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>

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.

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

Дивизион
Operations
Business Unit
Production / Manufacturing

Место
Мексика

Сайт
INSURGENTES

Company / Legal Entity

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

Functional Area

Technical Operations
Job Type
Full time
Employment Type
Regular
Shift Work
No

Ajustes de accesibilidad

Novartis tiene el compromiso de trabajar y proporcionar adaptaciones razonables para personas con discapacidad. Si, debido a una condición médica o discapacidad, necesita una adaptación razonable para cualquier parte del proceso de contratación, o para desempeñar las funciones esenciales de un puesto, envíe un correo electrónico a tas.mexico@novartis.com y permítanos conocer la naturaleza de su solicitud y su información de contacto. Incluya el número de posición en su mensaje.

Job ID
REQ-10076161

Manufacturing Specialist

[Apply to Job](#)

Job ID
REQ-10076161

Manufacturing Specialist

[Apply to Job](#)

Source URL: <https://www.novartis.ru/kr-ko/careers/career-search/job/details/req-10076161-manufacturing-specialist-es-es>

List of links present in page

1. <https://www.novartis.com/about/strategy/people-and-culture>
2. https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf
3. <mailto:tas.mexico@novartis.com>
4. https://novartis.wd3.myworkdayjobs.com/es/Novartis_Careers/job/INSURGENTES/Manufacturing-Specialist_REQ-10076161-1
5. https://novartis.wd3.myworkdayjobs.com/es/Novartis_Careers/job/INSURGENTES/Manufacturing-Specialist_REQ-10076161-1