

Global GMP Quality Auditor

Job ID

REQ-10086366

Aug 24, 2026

LOC_MX

About the Role

Key Responsibilities

- Plan, lead, and execute GMP audits across global operations and external partners.
- Assess compliance with regulations, quality standards, procedures, and guidance documents.
- Document audit observations, prepare reports, and communicate findings to stakeholders.
- Review and approve corrective and preventive action plans addressing audit findings.
- Provide expert guidance and training on audit activities.
- Escalate critical compliance issues and support timely mitigation activities when required.
- Maintain expertise in evolving regulations, industry standards, and quality expectations.

Essential Requirements

- Degree in Chemistry, Pharmacy, Biology, Engineering, or another related scientific discipline; other degrees with relevant experience may be considered.
- Minimum 10 years of broad experience in the pharmaceutical or medical device industry, including QA/QC management and manufacturing, development, or other relevant experience such as working at a regulatory health authority.
- Expertise in at least one of the following areas: drug product (DP) manufacturing, biologics, sterile manufacturing, laboratories, microbiology, packaging, active pharmaceutical ingredients (API), excipients, medical devices, computer system validation, or quality systems.
- Excellent knowledge of regulatory requirements, with experience and/or interaction with local Health Authorities and sporadically with other Health Authorities; sound and practical judgement in the interpretation and application of regulations and standards.
- Strong interpersonal skills, including diplomacy, persuasion, and stakeholder management.
- Willingness to travel approximately 60% globally.
- Advanced oral and written English communication skills.

Desirable Requirements

- 3 years auditing experience
- Proficiency in an additional language such as German, French, Italian, Chinese, or Spanish.

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Accessibility and accommodation

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Role Requirements

Division

DIV_TO

Business Unit

Quality

Location

LOC_MX

Site

INSURGENTES

Company / Legal Entity

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

Functional Area

FCT_QA

Job Type

Full time

Employment Type

Regular

Shift Work

No

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