NOVARTIS

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.

Commitment to Diversity and Inclusion

Novartis is committed to building an outstanding, inclusive work environment and diverse teams 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.

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
Apply to Job

Job ID
REQ-10086366

Global GMP Quality Auditor

Apply to Job

Source URL: *https://jobapi.novartis.com/req-10086366-global-gmp-quality-auditor*

List of links present in page

1. https://jobapi.novartis.com/req-10086366-global-gmp-quality-auditor
2. mailto:tas.mexico@novartis.com
3. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/INSURGENTES/Global-GMP-Quality-Auditor_REQ-10086366-1
4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/INSURGENTES/Global-GMP-Quality-Auditor_REQ-10086366-1