

Senior Manager, Internal Audit – Tech Ops

Job ID

REQ-10077262

Jul 16, 2026

LOC_ES

About the Role

Key Responsibilities

- Lead audit and advisory engagements end-to-end across manufacturing, supply chain, and technical operations.
- Deliver risk-based insights by assessing root causes and recommending practical, value-adding improvements.
- Ensure consistent, high-quality execution aligned with Internal Audit standards and quality assurance programs.
- Prepare clear, impactful audit and advisory reports for senior operations stakeholders.
- Act as a trusted advisor to senior Operations stakeholders, providing independent challenge and practical insight.
- Contribute to the continuous improvement of audit methodology, supporting consistent, high-quality execution and AI/digitally enabled audit practices.
- Contribute to operations audit planning by identifying emerging risks across manufacturing and supply processes.
- Coach and develop auditors through onboarding, peer coaching, and knowledge sharing.
- Champion data-driven and digitally enabled audit practices in collaboration with Audit Intelligence Enablement teams.

Please be advised that this role will have a global travel requirement of 30%-40%.

Essential Requirements

- University degree in technical or scientific discipline related to manufacturing or supply chain management.
- Extensive experience in Internal Audit or assurance roles within manufacturing, supply, or regulated operations.
- Strong understanding of pharmaceutical manufacturing, supply chain, and quality environments.
- Proven ability to lead complex audit or advisory engagements and influence senior operations stakeholders.
- Hands-on experience using data analytics, digital tools, and artificial intelligence (AI) to enhance audit outcomes in complex or regulated environments.
- Solid knowledge of pharmaceutical regulatory frameworks, including GMP, GDP, FDA, EMA, and related standards.
- Fluent English, with the ability to communicate credibly at senior leadership level; additional European languages are an advantage.

Desirable Requirements

- Preferred background in Pharmaceuticals, Life Sciences, Biotechnology, or Big Four audit/consulting with a healthcare or regulated operations focus.
- Professional certifications are a plus, including CIA (Certified Internal Auditor) and/or CRMA (Certification in Risk Management Assurance).

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

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>

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

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

Primary location salary range

€59,100.00 - €109,700.00

Division

DIV_NI

Business Unit

Audit & Compliance

Location

LOC_ES

Site

Barcelona Gran Vía

Company / Legal Entity

ES06 (FCRS = ES006) Novartis Farmacéutica, S.A.

Functional Area

FCT_FN

Job Type

Full time

Employment Type

Regular

Shift Work

No

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