

Clinical Quality Assurance - Program Lead

Job ID REQ-10061165

9月 10, 2025

Spain

摘要

-Bieten Sie Qualit ä tssicherung und Compliance-Aufsicht für Entwicklungs- und Forschungsfunktionen. Fördern Sie die Überwachung von Qualit ä tsmanagementsystemen und -initiativen innerhalb der globalen, regionalen und Länderorganisation und stellen Sie die Einhaltung der geltenden regulatorischen Anforderungen der Gesundheitsbehörden (z. B. GCP, GLP, GMP, PV, IP) und der Novartis-Verfahren und Qualit ä tsstandards sicher. Vorbild für gutes Qualit ä tsverhalten bei gleichzeitiger Förderung einer Qualit ä tskultur (z. B. richtiges erstes Mal usw.), um die nichtqualitativ hochwertigen Stakeholder (z. B. NIBR, GDD) positiv zu beeinflussen. Entwickeln, vorantreiben und/oder unterst ützen Sie Qualit ä tsplaninitiativen, um Die Organisationsstrategie, Mission und Vision zu erreichen.

About the Role

Key Responsibilities:

- Proactively provide QA leadership to the business strategy for assigned programs/trials by ensuring considerable organization awareness (e.g. Interrelationship of departments and business priorities),
- Drive implementation of quality strategy within Global Clinical Team (GCT)/ Clinical Trial Team (CTT) under responsibility
- Regularly monitor the implementation of the annual Quality Plan pertaining to the assigned programs/studies
- Ensure adequate oversight of proactive quality risk management process in the overseen areas including quality risk assessments and submission/inspection readiness activities and ensure that Clinical Trial Process (CTP) are in control
- Provide robust and clear quality oversight in the following areas of clinical development:
 - Support/collaborate with key stakeholders (e.g., Country Development Quality (CDQ), Development Units (DUs), GCT and/or CTT members) to ensure that risks are detected and remediated.
 - Support core governance for quality incident management for critical and major deviations pertinent to the programs being assigned and ensure timely escalation when required.
 - Provide Good Clinical Practice (GCP) guidance to day-to-day questions arising from Clinical trials deliverables.
 - Collaborate with Country Development QA and External Service Providers (ESP) QA to drive initiatives relevant to internal monitoring and outsourced activities Quality oversight.
 - Support inspections preparation and facilitation in collaboration with other QA groups within Research & Development Quality (RDQ).
 - Support audits and inspections follow-up activities including Corrective & preventative Actions (CAPA) preparation.
- Actively leverage audit/inspection outcomes/trends to sustain improvement in clinical trials conduct.
- Active participation in continuous improvement initiatives (including Work streams) and ensure that areas identified as weaknesses are properly being addressed and executed for sustainability
- Be QA point of contact for the defined trials and attend the meetings and ensure quality is embedded in the decision taking processes.

Essential Requirements:

- Bachelor's degree in life science or healthcare field required. Advanced degree or equivalent education/degree in life sciences/healthcare preferred (PhD/MD/ PharmD/ Masters).
- 7 years of involvement in regulated activities (GCP/ Pharmacovigilance (PV)), clinical development and/or QA positions.
- Broad understanding of global expectations of Health Authorities in the area of Clinical Development and profound understanding of the science of product
- · development.
- Ability to work independently and in a global/matrix environment.
- 3 or more years 'experience in managing projects.
- Strong skills in GCP, quality and/or clinical development

Why Novartis:

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部门 Development

Business Unit Innovative Medicines

地点 Spain

站点

Barcelona Gran V í a

Company / Legal Entity ES06 (FCRS = ES006) Novartis Farmac é utica, S.A.

Alternative Location 1 Basel (City), Switzerland

Alternative Location 2 Dublin (NOCC), Ireland

Alternative Location 3 Home Worker - England/Wales, United Kingdom

Alternative Location 4 London (The Westworks), United Kingdom

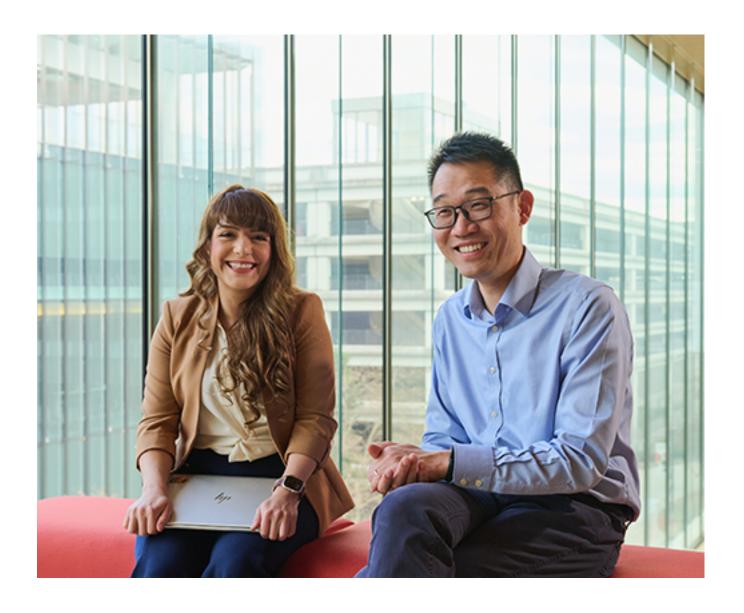
Functional Area Quality

Job Type Full time

Employment Type Regul ä r

Shift Work No

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