

Associate Director External Service Provider Quality Assurance

Job ID REQ-10042563 Mar 03, 2025 Irlanda

Resumen

As Associate Director External Service Provider QA, you'll have the thrilling opportunity to oversee the implementation of top-notch quality standards, cutting-edge processes, and innovative tools and systems.

You will play a pivotal role in managing external service providers supporting Novartis Global Clinical Trials in R&D Quality, ensuring that our partners meet the highest standards of excellence.

About the Role

Deadline for applications: 10th of March 2025

Major Accountabilities:

- Responsible for review and approval of External Service Providers (ESP) qualification for global clinical trials.
- Responsible for negotiation and execution Quality Assurance Agreements (QAA) / quality terms with ESPs.
- Review and approval of external service provider Quality Risk Assessments (QRAs) to enable
 identification and evaluation of various metrics, risks, trends, and potential quality and performance
 issues with the ESP in a proactive manner. Ensure communication and support mitigation of actions for
 potential risks.
- Responsible for review and approval of quality issues related to ESPs and ensure appropriate escalation of major and critical issues. Support assessment of serious breach and reporting to health authorities.
- Collaboration with business partners such as, Vendor Partnership and Governance, Global Medical
 Affairs, other applicable Vendor Business Offices and Procurement to ensure their involvement in the risk
 evaluation and timely communication to the business and follow-up on required actions.
- Review quality metrics, monitoring and reporting including follow-up with line functions and escalation.
- Ensure inspection readiness of ESP related activities and support for internal and external audits and health authority (HA) inspections pertaining ESP management.
- Ensure compliance with regulatory requirements (GCP, GLP, GVP, GMP) and continuous improvement of quality relevant processes within area of responsibility.

Obligatory requirements:

 Education: Minimum a bachelor's degree in Life Sciences, Pharmacy or Medicine, or other related discipline required

- 12+ years' experience in pharmaceutical development and excellent knowledge of the quality management system, clinical operations processes and vendor management related activities is preferred.
- Good knowledge of GCP/GLP, GMP, GVP.
- Demonstrated leadership in implementing robust processes and quality systems, and setting global quality standards in a regulated area, including controlled documentation for the pharmaceutical development area.
- Experience in managing External Service Providers and knowledge of quality standards and regulatory requirements. Experience in interactive response technology (IRT), eCOA and Imaging services is added advantage.
- Thorough technical understanding of quality system, clinical trial process collecting, analyzing, and monitoring of Third-party Key Quality Indicators.
- Experience in data analysis and trending using available tools.
- Demonstrated effective management and establishment of successful international and cross-divisional collaborations.
- Fluent English, written and spoken

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Accessibility and accommodation: Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to diversity.inclusion_ch@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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División Development **Business Unit** Innovative Medicines Ubicación Irlanda Sitio Dublin (NOCC) Company / Legal Entity IE02 (FCRS = IE002) Novartis Ireland Ltd Alternative Location 1 Barcelona Gran Vía, España Alternative Location 2 London (The Westworks), Reino Unido **Functional Area**

Quality

Job Type

Full time

Employment Type

Regular

Shift Work

No

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