NOVARTIS

Senior Principal Clinical Data Scientist

Job ID
REQ-10051889
Jul 31, 2025
LOC_IE

About the Role

Major accountabilities:

- Lead functional activities for a medium to large sized project in phase I to IV clinical studies in Novartis Global Development Organization.
- Co-ordinate activities of Data Managers either internally or externally.
- Make data Mgmt decisions and propose strategies at study or project level.
- Ensure application of consistent data Mgmt processes, influence increased standardization and documentation across assigned project/programs -Comply with company, department and industry standards and processes.
- Provide and implement data Mgmt solutions; ensure knowledge sharing.
- Leads process and training deliverables within multiple platforms, franchises or therapeutic areasDevelops strategies to ensure effective training and knowledge retention.
- Progresses DO towards complete, compliant, agile and simple end to end processes and effective training (Protocol/Measure through Analysis and Reporting).
- Drives towards agreed deliverables, proactively addressing potential issues before they become problematic -Build and maintain effective working relationship with cross-functional teams, able to summarize and discuss status of deliverables and critical data Mgmt aspects.
- Represents DO in all audits and inspections, centralizing and aligning the team in audit preparation, readiness and response.
- Act as subject matter expert (SME) or, as assigned, lead process improvement/non clinical project initiatives.
- Develops risk Mgmt strategies to prevent data quality/coding issues from derailing projects -Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt - Distribution of marketing samples (where applicable)

Minimum Requirements:

- Degree / Masters qualified in a relevant area
- Ideally 9+ years’ experience in Drug Development with at least 8 years' in Clinical Data Management
- Experience working across several end to end studies
- Strong leadership, collaboration and organizational skills with proven ability to successfully manage simultaneous trials and meet deadlines.
- A background of coaching and mentoring team members as required, ensuring that data management associates on the individual programs are aware of the risks, priorities, goals and impact of the work contribution
- Excellent understanding of clinical trials methodology, GCP and medical terminology
- Proven ability to interrogate and view data through various programming/GUI techniques.

Commitment to Diversity & Inclusion:

We are committed to building an outstanding, inclusive work environment and diverseteams 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)

Division
DIV_GD
Business Unit
Development
Location
LOC_IE
Site
Dublin (NOCC)
Company / Legal Entity
IE02 (FCRS = IE002) Novartis Ireland Ltd
Functional Area
FCT_RD
Job Type
Full time
Employment Type
Regular
Shift Work
No
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