NOVARTIS

Expert Clinical Programmer

Job ID
REQ-10084428
Jul 28, 2026
LOC_IN

About the Role

Major accountabilities:

- Produce and track reports, analytics or visualizations for various line functions within Global Drug Development, used for ongoing monitoring of clinical data
- Provide understandable and actionable reports on clinical data and monitoring of clinical data for key stakeholders
- Author the user requirements document, functional specifications and functional testing scripts
- Facilitate interaction with end-user on creating specifications and working with programmers or performing the programming activities for successful delivery.
- To provide quantitative analytical support to the global program teams, including providing support on analyzing reports
- Support the planning, execution and close-out of Clinical Programs/Trials.
- Support the management in collation and delivery of analytics reports for critical decision making
- Create, file and maintain appropriate documentation
- Work with the internal SMEs and key stakeholders in providing analysis and interpretation of clinical program/trial operational data
- Provide necessary training to end-user on best / appropriate and consistent use of various data review tools
- Support or lead special projects of limited scope (sub team lead, local project, etc.) both clinical and non-clinical in nature.
- Provide study level expertise and involvement in CTTs.
- Lead or provide support to special projects both clinical and non-clinical in nature or in general areas spanning various responsibilities but not limited to systems issues, processes, user support, training, etc.

Minimum Requirements:

- At least 6 years’ experience in clinical review and reporting programming, business analytics and/or clinical trial setup, gained
- in the pharmaceutical industry, CRO or Life Science related industry as well as the following:
- Excellent knowledge of programming languages (SQL, PL/SQL,
- C#, VB script, SAS, Python, R)
- • Excellent knowledge of Data Review and/or Business Intelligence tools (such as Spotfire, Review)
- Knowledge of clinical data management systems and/or relational databases (e.g. OC/RDC, INFORM, RAVE) as applied to
- clinical trials
- Attention to detail, quality, time management and customer focus development. Understanding of Drug Development Process, ICH-GCP, CDISC standards and Health Authority guidelines and regulations.

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_IN
Site
Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Alternative Location 1
LOC_IN
Functional Area
FCT_RD
Job Type
Full time
Employment Type
Regular
Shift Work
No
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