

Senior Clinical Programmer

Job ID REQ-10047311 Jun 26, 2025 India

Summary

Participate in the full lifecycle of producing key data and/or reports in support of data review reporting development including evaluation of requirements, design specifications, interface to programmers, report programming, coordinate validation and rollout activities along with providing quantitative analytical support. These tasks are to be performed independently or team based with minimal guidance and supervision.

About the Role

- 1. Produce and track reports for various line functions within Global Drug Development, used for ongoing monitoring of clinical data
- 2. Provide understandable and actionable reports on clinical data and monitoring of clinical data for key stakeholders
- 3. Author /co-author the user requirements document, functional specifications and functional testing scripts
- 4. Facilitate interaction with end-user on creating specifications and working with programmers or performing the programming activities for successful delivery.
- 5. To provide quantitative analytical support to the global program teams, including providing support on analyzing reports
- 6. Support the planning, execution and close-out of Clinical Programs/Trials.
- 7. Support the management in collation and delivery of analytics reports for critical decision making
- 8. Create, file and maintain appropriate documentation
- 9. Work with the internal SMEs and key stakeholders in providing analysis and interpretation of clinical program/trial operational data
- 10. Provide necessary training to end-user on best / appropriate and consistent use of various data review tools
- 11. Program reports of various complexity from documented requirements, within the clinical reporting systems using SQL, PL/SQL, C#, VB script, SAS, Python, R.
- 12. Good Knowledge of Novartis Clinical Data Standards and its implementation for creation of reports specifications or reports output
- 13. Support special projects of limited scope (sub team lead, local project, etc.) both clinical and non-clinical in nature.
- 14. Provide study level expertise and involvement in CTTs.
- 15. 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.

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?

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Division

Development

Business Unit

Innovative Medicines

Location

India

Site

Hyderabad (Office)

Company / Legal Entity

IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

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Accessibility and accommodation

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

Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.

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