Principal Clinical Programmer

Job ID
REQ-10047309

4月 11, 2025

India

摘要

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. Drive the implementation of data analytics reports and dashboards for optimal data review by working with the users to establish robust user specifications and with programmers to implement the optimal output.
2. Translate business requirements into logical models and provide direction to the development team to translate business logic.
3. Lead authoring of the user requirements document, functional specifications and functional testing

scripts
4. Proactively identify or address needs for optimal data review working with users and programmers as appropriate.
5. Implement and execute robust project plans for delivery, ensuring customer needs are addressed in a timely manner,
6. Provide coordination between the project resources so that deadlines are met on deliverables.
7. Drive development of appropriate user training. Drive all necessary change management activities related to implementation of new data review tools / reports as related to data cleaning, review and visualization.
8. Provide understandable and actionable reports on clinical data and monitoring of clinical data for key stakeholders.
9. Provide quantitative analytical support to the global program teams, including providing support on analyzing reports and defining KPI Metrics.
10. Systematically sample / monitor utilization of reports and tools, perform and coordinate periodic review of outputs. Report findings and take necessary steps to ensure reports are being used for optimal data review via Metrics.
11. Create, file and maintain appropriate documentation
12. Work with the internal SMEs and key stakeholders in providing analysis and interpretation of clinical program/trial operational data.
13. Lead initiatives within Analytics, including training, to support overall compliance and adherence to data review through robust reports and tools.
14. Proactively identify and address areas of concerns to avoid issues and ensure consistency, accuracy and completeness of reported clinical data.
15. 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 and on GPTs for small projects.
16. May assist manager in coordinating and directing Clinical Reporting activities of External Partners (Contingent Workers, Functional Service Providers, Interns, etc.)
17. Program reports of advanced complexity from documented requirements, within the clinical reporting systems using SQL, PL/SQL, C#, VB script, SAS, Python, R.
18. Demonstrated experience in the concepts and use of Clinical Data Standards (e.g. SDTM, Novartis standard) for creation of report specifications or reports output.
19. Lead 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? https://www.novartis.com/about/strategy/people-and-culture

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: https://talentnetwork.novartis.com/network

Benefits and Rewards: Read our handbook to learn about all the ways we ’ ll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

部门
Development

Business Unit
Innovative Medicines

地点
India

站点
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

Apply to Job

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.

Job ID
REQ-10047309

Principal Clinical Programmer

Apply to Job

Source URL:
https://www.novartis.com.cn/careers/career-search/job/details/req-10047309-principal-clinical-programmer

List of links present in page

1. https://www.novartis.com/about/strategy/people-and-culture
2. https://talentnetwork.novartis.com/network
3. https://www.novartis.com/careers/benefits-rewards
4. https://novartis.wd3.myworkdayjobs.com/en-US/NovartisCareers/job/Hyderabad-Office/Principal-Clinical-ProgrammerREQ-10047309-1
5. mailto:diversityandincl.india@novartis.com
6. https://novartis.wd3.myworkdayjobs.com/en-US/NovartisCareers/job/Hyderabad-Office/Principal-Clinical-ProgrammerREQ-10047309-1