

# Senior Principal Statistical Programmer

Job ID  
REQ-10075745  
апр 14, 2026  
Индия

## Сводка

Responsible for all statistical programming/data review reporting and analytics development aspects of several studies, a medium to large sized project or project-level activities. Acts as a key collaborator and strategic partner in ensuring that drug-development plans are executed efficiently with timely and high-quality deliverables. Complies with project / study standards and specifications following internal and regulatory guidelines. Oversees programming style, quality of statistical reporting & compliance with timelines.

## About the Role

### Key Responsibilities

- Lead statistical programming activities for several studies or drive the implementation of data analytics reports. Make decisions and propose strategies at study or project level.
- May act as functional manager for local associates including providing supervision and advice on functional expertise and processes.
- Build and maintain effective working relationship with cross-functional teams, able to summarize and discuss status of deliverables and critical aspects (timelines, scope, resource plan), e.g. as representative in study or project-level team.
- Ensure project-level standardization. Provide and implement programming solutions; ensure knowledge sharing.
- Act as expert in problem-solving aspects.
- Ensure timely and quality development and validation of datasets and outputs for regulatory submissions/interactions, safety reports, publications, post-marketing activities etc. Leads/co-leads novel projects within the team -Generates innovative ideas within own team and /or project team /functional community

### Essential Requirement

- Demonstrates strong proficiency in **SAS** for the analysis and summarization of clinical trial data.
- Has served as a **Trial Programmer** or in a comparable programming role with end-to-end study responsibility.
- Possesses experience in the **development and/or review of critical study documents**, including Protocols, eCRFs, Data Transfer Specifications, SAPs, and mock shells, ensuring consistency with study objectives and regulatory expectations.
- Shows openness to adopting **R and other programming languages**, with a willingness to embrace emerging technologies such as **AI/ML**.
- Exhibits a comprehensive understanding of **CDISC data standards** and their application across clinical studies.
- Minimum 6 years + with Graduation. (MSc preferred)

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Дивизион

Development

Business Unit

Development

Место

Индия

Сайт

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**

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