

# Principal Statistical Programmer

Job ID  
REQ-10075744  
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 5 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

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

### **Accessibility and accommodation**

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