

## Senior Principal I Principal Biostatistician

Job ID  
REQ-10061483

9月 04, 2025

USA

### 摘要

Fournir un soutien d'expert et des connaissances fonctionnelles et techniques pour assurer l'intégrité / validité scientifique pour le développement clinique, le développement précoce et / ou les projets de recherche. Participer au cycle de vie complet de la production de données clés et / ou de rapports à l'appui de la élaboration de rapports d'examen des données, y compris l'évaluation des exigences, les spécifications de conception, l'interface avec les programmeurs, la programmation des rapports, la coordination des activités de validation et de déploiement ainsi que la fourniture d'un soutien analytique quantitatif. Fournir un soutien statistique pour les soumissions réglementaires, y compris la planification, l'analyse et la production de rapports sur l'innocuité clinique et l'efficacité. Peut également fournir un soutien statistique à la recherche ou à d'autres domaines de R&D.

Responsable de conseiller / diriger la planification, le développement et la mise en œuvre de technologies conformes à l'industrie (CDISC et réglementaires) conformes aux normes de données cliniques, d'infrastructure ou d'automatisation conformes à l'industrie (CDISC et réglementaires). Fournir un support expert et une orientation client exceptionnelle aux utilisateurs professionnels et aux équipes sur leur utilisation, notamment:  
Outils de collecte standard de données dans EDC (CRF, vérifications des modifications, dérivations, configurations de base)

-Sp é cifications de transfert de donn é es  
Donn é es d ' analyse/Normes TFL/D é finir  
Solutions / technologies d ' automatisation  
-nfrastructure d ' entreprise, r è gles d ' affaires et directives.

## About the Role

### Your Key Responsibilities:

- Responsible for all statistical tasks on the assigned trials.
- Protocol development in alignment with the development plan, developing statistical analysis plan, reporting activities.
- Contribute to planning and execution of exploratory analyses and statistical consultation within your cross-functional teams.
- Initiate, drive and implement novel methods and innovative trial designs in alignment with the Lead Statistician.
- Collaborate with clinical, regulatory and other strategic functions to drive quantitative decision making in assigned indications/program with oversight.
- Collaborate cross-functionally (e.g. data management, programming, medical writing) to ensure timeliness and quality of statistical deliverables.
- Contribute to external engagement with consultants, advisory boards, health authorities, congresses, and scientific meetings

Video Link [Meet the Data Analytics team](#)

This position will be located at the East Hanover, NJ site and will not have the ability to be located remotely.

### Role Requirements:

#### Essential Requirements:

- PhD with 3+ years ' experience preferred OR MS with 7+ years ' experience
- Fluent English (oral and written)
- Strong communication and presentation skills

### Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between \$119,700 and \$222,300 per year. The final salary offered is determined based on factors like, but not limited to, relevant skills and experience, and upon joining

Novartis will be reviewed periodically. Novartis may change the published salary range based on company and market factors.

Your compensation will include a performance-based cash incentive and, depending on the level of the role, eligibility to be considered for annual equity awards. US-based eligible employees will

receive a comprehensive benefits package that includes health, life and disability benefits, a 401(k) with company contribution and match, and a variety of other benefits. In addition, employees are eligible for a generous time off package including vacation, personal days, holidays and other leaves. To learn more about the culture, rewards and benefits we offer our people click [here](#).

#### Why Novartis:

Our purpose is to reimagine medicine to improve and extend people's lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <https://www.novartis.com/about/strategy/people-and-culture>

#### You'll receive:

You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook.

<https://www.novartis.com/careers/benefits-rewards>

**Accessibility and Reasonable Accommodations:** The Novartis Group of Companies are 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 application process, or in order to perform the essential functions of a position, please send an e-mail to [tas.nacomms@novartis.com](mailto:tas.nacomms@novartis.com) call +1(877)395-2339 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

**Join our Novartis Network:** If this role is not suitable to your experience or career goals but you wish to stay connected to learn more about Novartis and our career opportunities, join the Novartis Network here: <https://talentnetwork.novartis.com/network>

**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>

## EEO Statement:

The Novartis Group of Companies are Equal Opportunity Employers. We do not discriminate in recruitment, hiring, training, promotion or other employment practices for reasons of race, color, religion, sex, national origin, age, sexual orientation, gender identity or expression, marital or veteran status, disability, or any other legally protected status.

## Accessibility & Reasonable Accommodations

The Novartis Group of Companies are 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 application process, or to perform the essential functions of a position, please send an e-mail to [us.reasonableaccommodations@novartis.com](mailto:us.reasonableaccommodations@novartis.com) or call +1(877)395-2339 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

部门  
Development

Business Unit  
Universal Hierarchy Node

地点  
USA

状态  
New Jersey

站点  
East Hanover

Company / Legal Entity  
U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area

Research & Development

Job Type

Full time

Employment Type

CDI

Shift Work

No

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2. <https://www.novartis.com/sites/novartiscom/files/novartis-life-handbook.pdf>
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