NOVARTIS

Associate Director / Director, Biostatistics

Job ID
REQ-10088039
Окт. 02, 2026
Великобритания

Сводка

Associate Director / Director Biostatistics
#LI-Hybrid
Location: London, UK

This role is based in London, UK. Novartis is unable to offer relocation support for this role. Please only apply if this location is accessible for you.

Biostatisticians at Novartis transform data into evidence that informs decisions across the product lifecycle and supports access to innovative medicines.

We are seeking an Associate Director/Director, HTA Biostatistics, to provide strategic statistical leadership for Health Technology Assessment, pricing and reimbursement and market access activities. The role will partner with International Value & Access, Health Economics and Outcomes Research, Medical Affairs, Clinical Development, Real-World Evidence, and external experts to deliver scientifically rigorous and payer-relevant evidence.
The Associate Director/Director will help advance Novartis HTA capabilities, support reimbursement dossiers in Novartis key markets, and contribute to the Joint Clinical Assessment dossiers as outlined in Regulation (EU) 2021/2282.

About the Role

Major accountabilities:

- Provide strategic statistical leadership for HTA, pricing and reimbursement, market access, and integrated evidence-generation activities across assigned programs.
- Partner with cross-functional and cross-regional teams to ensure HTA evidence is scientifically rigorous, decision-relevant, and aligned with intended use.
- Lead the implementation of efficient HTA methods. Demonstrated hands-on experience with population-adjustment methodology and evidence synthesis. Ability to select, implement, critique, and defend methods under payer scrutiny.
- Advanced practical experience in several additional methods relevant to HTA, such as survival extrapolation, treatment-switching adjustment, health-state utility analysis for mapping and application of preference weights, causal inference involving real-world evidence, missing-data methods, external controls and transportability.
- Strong understanding of drug development and the relationship between regulatory and payer requirements across the product lifecycle.
- Drive early and continuous evidence synthesis strategy to inform HTA planning and decision-making.
- Contribute to standards, reusable templates, automation, and best practices that improve the quality, consistency, and scalability of HTA analyses.
- Support responses to payers and HTA questions and contribute to scientific interactions with decision-making agencies, healthcare professionals, and external experts.
- Communicate complex statistical findings, assumptions, limitations, and recommendations clearly to statistical and non-statistical stakeholders.
- Contribute to publication work across assets.
- Plan, prioritize, and oversee project-level activities across parallel programs, ensuring effective resource management and delivery of high-quality statistical outputs.
- Support vendor selection and provide scientific and quality oversight for outsourced HTA activities
- Established scientific credibility demonstrated through peer-reviewed publications, conference presentations, methodological contributions, guidance development, or active leadership in relevant professional or HTA special-interest groups. Director candidates should demonstrate a sustained and visible external contribution.
- Coach and mentor biostatisticians in HTA statistical methodologies, processes, and scientific communication.

Essential requirements

- PhD in Statistics, Biostatistics, Mathematics or a related quantitative field with at least 8 years of relevant experience, or a master’s degree with at least 12 years of relevant experience.
- Demonstrated expertise in HTA, reimbursement evidence generation, and indirect treatment comparison methods
- Advanced understanding of drug development and the evidence needs, decision drivers, and review processes of major HTA and reimbursement bodies across multiple regions.
- Demonstrated ability to lead and deliver multiple complex, parallel projects across therapeutic areas and to influence senior decision-making in a global matrix environment.
- Excellent interpersonal and communication skills

Desirable requirements

- Experience supporting Joint Clinical Assessment requirements in Europe.
- Experience in Immunology and/or another major therapeutic area.
- Experience with health economics and outcomes research, real-world evidence, data science, automation, or AI-enabled evidence generation.

Benefits & Rewards

At Novartis, we’re committed to reimagining medicine together - and rewarding the people who make it happen.

Expected Annual Base Salary Range for role: 83,510.00 - 155,090.00 GBP Annual

The base salary offered is determined based on gender-neutral objectives, such as relevant skills, competencies and experience in accordance with the Novartis pay setting policy and upon joining Novartis will be reviewed periodically.

In addition to your base salary, you may be eligible for a performance-based bonus depending on certain performance parameters.

The rewards of being part of our team go far beyond base pay and incentives. We also offer a variety of competitive benefits in kind to help you thrive personally and professionally, such as insurance plans, retirement plans, wellbeing resources and global recognition programs. In addition, we provide flexible and hybrid working options, where possible, and minimum 14 weeks paid parental leave.

In addition to your base salary, you may be eligible for a performance-based bonus depending on certain performance parameters Long-term equity awards granted at group level may also be part of your package. Further details will be provided during the application process.

Read our brochure to learn more about our global total rewards offering:

https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf

Note: Benefits and compensation may vary by country and are subject to local legal requirements, including provisions of collective bargaining agreements where applicable. A full overview of your compensation package, including any relevant collective bargaining agreement details applicable to your role based on your employment location and Novartis employer entity, will be communicated separately to you during the application process.

Commitment to Diversity and Inclusion / EEO paragraph

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

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

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

Primary location salary range
£83,510.00 - £155,090.00
Дивизион
Development
Business Unit
Development
Место
Великобритания
Сайт
London (The Westworks)
Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No

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