

# Clinical Data Strategy Expert, Associate Director

Job ID  
REQ-10080343  
Июн. 24, 2026  
США  
Available in: English

## Сводка

Cambridge MA  
Internal: Associate Director  
LI#-Hybrid

The Associate Director, Clinical Data Strategy is a strategic leadership role within Translational Medicine (TM), responsible for shaping and driving end-to-end clinical data strategies at the program and trial level.

This role ensures that clinical data is planned, structured, and leveraged effectively from early development through execution, enabling high-quality, decision-ready data to support portfolio progression and data-driven decision-making.

Working in close partnership with cross-functional stakeholders, the Associate Director provides strategic oversight, governance, and expertise to align data strategy with scientific, operational, and regulatory objectives across the TM portfolio.

## About the Role

### Key Responsibilities

#### Lead Clinical Data Strategy

- Define and drive study- and program-level data strategies in the Neuroscience (primary) and other (secondary) disease areas ensuring alignment of assessment selection, performance, collection, and data flow with scientific, operational, and regulatory objectives

#### Drive Cross-Functional Alignment

- Partner with Clinical, Medical, Data Management, Biostatistics, and Therapeutic Area experts to ensure assessment selection is feasible, well-defined, and executable

#### Provide Portfolio-Level Oversight

- Guide data and assessment strategy consistency across studies/programs/therapy areas, ensuring prioritization, risk visibility, and alignment with TM portfolio objectives

#### Advance Innovation & Quality

- Lead improvements in data and endpoint planning, acquisition, and usability, including optimization of complex or novel endpoint data capture and quality, and adoption of innovative technologies and streamlined processes

### Essential Requirements:

- Advanced scientific degree with relevant experience in clinical trial management, data management, or PRO/COA usage in a Pharmaceutical/CRO or clinical site/functional endpoint environment
- Strong understanding of clinical development, study planning, and trial operations with direct experience in Neuroscience disease populations
- Demonstrated ability to lead cross-functional initiatives and influence stakeholders
- Track record of driving strategic planning, innovation, and process improvement

The salary for this position is expected to range between: \$152,800-\$283,400/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:** 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 \(PDF 30 MB\)](#)

### EEO Statement:

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protected status.

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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.

Дивизион

Biomedical Research

Business Unit

Research

Место

США

Состояние

Massachusetts

Сайт

Cambridge (USA)

Company / Legal Entity

U175 (FCRS = US175) Novartis Institutes for BioMedical Research, Inc.

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

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