

# Senior Clinical Sciences Trial Leader

Job ID  
REQ-10044734  
Mar 25, 2025  
Japan

## Summary

1. Responsible for financial and resource decisions within scope of assigned authority  
2. Responsible for the availability of high quality, Biomedical Research data according to agreed timelines and budget to enable no delays in strategic decision making and drug registration. External impact: Novartis perceived as a credible, ethical, and preferred business partner.

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## About the Role

- Study Leader and/or Clinical Scientist for predominantly medium to high complexity, global studies and may provide additional Clinical Sciences support to high priority, high complexity, global studies.
- Lead the clinical protocol development process in collaboration with the Medical Lead and other line functions; responsible author for clinical protocols, amendments, etc.; contribute to the medical/scientific input given for the development of study-related documents and processes which resides in other line functions; contribute to the development of clinical sections of study-level regulatory documents.
- Lead development of strategic and scientific input into study concept, feasibility, and ability to execute; develops and implements study-level operational execution plan in partnership with key cross functional partners, if applicable.
- Collaborate with key cross functional partners to identify and select strategic and high performing sites to ensure recruitment commitments are met.

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## Work Experience:

- Bachelors in life science/healthcare required; Advanced degree or equivalent education/degree in life sciences/ healthcare preferred (PhD/MD/ PharmD/ Masters).
- Minimum 6+ years' experience in clinical trials development.
- Fluent English (oral and written)
- For TCO: Strong understanding of oncology/hematology and demonstrates high learning agility.
- Demonstrated ability to confidently drive complex collaborations through unpredictable circumstances

and higher paced changes.

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**Languages:**

- English.

**Benefits and Rewards:**

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

[novartis-life-handbook.pdf](#)

**Commitment to Diversity and Inclusion**

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

**Accessibility and accommodation**

Novartis is 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 recruitment process, or in order to perform the essential functions of a position, please send an e-mail to [midcareer-r.japan@novartis.com](mailto:midcareer-r.japan@novartis.com) and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

[novartis-life-handbook.pdf](#)

[midcareer-r.japan@novartis.com](mailto:midcareer-r.japan@novartis.com)

**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>

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**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>

Division

Biomedical Research

Business Unit

Pharma Research

Location

Japan

Site

Toranomon (NPKK Head Office)

Company / Legal Entity

JP05 (FCRS = JP005) Novartis Pharma K.K.

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

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