

## Clinical Pathologist, Preclinical Safety

Job ID  
REQ-10024580

1月 10, 2025

USA

### 摘要

#LI-Hybrid

Internal title: Senior Principal Scientist

This position will be located at our Cambridge MA or East Hanover, NJ site (Cambridge, MA preferred) and will not have the ability to be located remotely.

About the role:

As a Clinical Pathologist within the Preclinical Safety organization within BR's Translational Medicine organization, you will apply your expertise to research and preclinical safety studies through sound scientific assessment and integration of laboratory data, including non-traditional safety biomarker results, and providing, and reviewing for well-written interpretive reports. You will contribute to an integrated understanding of the patho mechanisms and human safety significance of non-clinical study findings as well as actively participate in the drug development decisions through input as Project Pathologist on a diverse portfolio of drug platforms and targets, and have the opportunity to

lead investigations for preclinical safety issue resolution.

## About the Role

Key responsibilities:

- Effectively assess for toxicological and pharmacological effects of small molecules and biologics in preclinical studies, with a focus on clinical pathology and relevant soluble biomarkers.
- Collaborate with global teams of pathologists and other scientists in design of preclinical safety studies and interpretation and communication of study findings, including the impact of the data on the safety profile and clinical development
- Provide expert review of protocols, results, and reports generated at contract laboratories and maintain effective relationships with CRO clinical pathologists, scientists, and technical staff to ensure high quality, on-time reporting.
- Contribute to scientific and technical support for identification, development, qualification, and validation of translational biomarkers targeted to support study safety assessments.
- Support continuous development of laboratory analytical capabilities and career development of the technical team in collaboration with the laboratory manager.
- Serve as Project Pathologist on assigned compounds, targets, and therapeutic areas
- Keep up to date on scientific literature relevant to your work on project and target teams, and to novel soluble biomarkers and applicable technologies

Essential Requirements:

- A Doctor of Veterinary Medicine (DVM) or equivalent related experience
- Board Certification in Clinical Pathology by American/European College of Veterinary Pathologists (ACVP or ECVCP)
- Preferred PhD in a biological science
- Some experience in toxicological pathology desired
- Experience preferred in development of novel biomarkers and immunoassays and/or other technical immunological methods (flow cytometry, multiplex technology, etc.), and application in safety assessment, preferably within pharmaceutical/biotech or contract research organization
- Experience preferred in interpreting clinical pathology data for laboratory animals, including

nonhuman primates and rodents

- Some understanding of the drug development process, preferably including experience working directly on drug development project teams and Regulatory interactions and submissions preferred
- Excellent verbal and written communication skills- Ability to work cross-functionally within a team and matrix environment

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Novartis Compensation and Benefit Summary: The pay range for this position at commencement of employment is expected to be between \$168,000 and \$312,000/year; however, while salary ranges are effective from 1/1/25 through 12/31/25, fluctuations in the job market may necessitate adjustments to pay ranges during this period. Further, final pay determinations will depend on various factors, including, but not limited to geographical location, experience level, knowledge, skills, and abilities. The total compensation package for this position may also include other elements, including a sign-on bonus, restricted stock units, and discretionary awards in addition to a full range of medical, financial, and/or other benefits (including 401(k) eligibility and various paid time off benefits, such as vacation, sick time, and parental leave), dependent on the position offered. Details of participation in these benefit plans will be provided if an employee receives an offer of employment. If hired, employee will be in an “at-will position” and the Company reserves the right to modify base salary (as well as any other discretionary payment or compensation program) at any time, including for reasons related to individual performance, Company or individual department/team performance, and market factors.

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部门

Biomedical Research

Business Unit

Pharma Research

地点

USA

状态

Massachusetts

站点

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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