

Clinical Pathologist, Preclinical Safety

Job ID
REQ-10024580
Oct 14, 2024
Estados Unidos

Resumen

#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

Benefits and Rewards:

Read our handbook to learn about all the ways we'll help you thrive personally and professionally: [Novartis Life Handbook](#)

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.

Novartis Compensation and Benefit Summary: The pay range for this position at commencement of employment is expected to be between \$183,200 and \$274,800/year; however, while salary ranges are effective from 1/1/24 through 12/31/24, 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.

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:
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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>

EEO Statement:

The Novartis Group of Companies are Equal Opportunity Employers who are focused on building and advancing a culture of inclusion that values and celebrates individual differences, uniqueness, backgrounds and perspectives. 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. We are committed to fostering a diverse and inclusive workplace that reflects the world around us and connects us to the patients, customers and communities we serve.

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 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.

División

Biomedical Research

Business Unit

Pharma Research

Ubicación

Estados Unidos

Sitio

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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