Position:

- Principal Research Scientist-Companion Animal Product Development

Company Overview:

Elanco is a global research-based company that develops and delivers product and services to enhance animal health and production. We value innovation, both in scientific research and daily operations, and strive to cultivate a collaborative work environment for more than 7,000 employees in more than 70 countries. Founded in 1954, Elanco is a division of Eli Lilly and Company.

Basic Qualifications:

- Doctor of Veterinary Medicine (DVM or equivalent) or PhD. degree
- 8 years of research experience with documentation of and conducting research in a regulated environment related to effectiveness of animal health therapeutics and/or parasiticides.

Additional Information:

Lilly is an EEO/Affirmative Action Employer and does not discriminate on the basis of age, race, color, religion, gender, sexual orientation, gender identity, gender expression, national origin, protected veteran status, disability or any other legally protected status.

Travel Percentage:

- 10-25%

Responsibilities:

Elanco Animal Health Research and Development is pursuing the development of pharmaceutical drugs and parasiticides for dogs and cats to fulfill a vision of companionship enriching life. This role contributes to the development of New Animal Health Products by designing, planning, coordinating, implementing, and reporting pilot and pivotal studies in compliance with global regulatory standards and guidelines. Authoring of study protocols, selecting Contract Research Organizations (CROs) to conduct studies as well as other study functions during planning, implementation and execution of studies (directly or via CROs) are also included. The individual in this role is expected to perform key leadership and technical responsibilities which provide innovative solutions, a positive and motivating work environment, ultimately delivering the Elanco pipeline.

Technical Expertise:

- Design efficacy program strategy within a Companion Animal product development team contributing to the development of New Animal Health Products.
- Manage and direct operational aspects of *in vivo* effectiveness programs including laboratory

studies and clinical field trials.

- Plan and implement all activities associated with studies including protocol development and preparation of final study reports according to relevant internal & global regulatory standards including GCP, VICH, and GLPs.
- Manage project timeline and budget deliverables
- Co-ordination of clinical trial material supplies and all study materials as necessary.
- Co-ordination of and collaboration within a multi-disciplinary project effectiveness team.
- Oversee Design/Develop study data output, and reports.
- Identify and select CROs and establish contract agreements, and maintain positive business collaboration.
- Liaise with Elanco/Lilly personnel e.g. regulatory, new product planning/marketing teams, procurement, legal, QA. Liaise externally with key opinion leaders, clinical experts and other contacts.
- Prepare regulatory effectiveness technical section in partnership with Elanco Regulatory Affairs and assist with Regulatory Submission.

Leadership:

- Proactively search for solutions.
- Work effectively and flexibly within and across all Elanco R&D teams and external collaborators to achieve overall Elanco R&D deliverables.
- Create a positive work environment that is aligned with company objectives.
- Provide and accept challenges to deliver innovative technical solutions and create an innovative culture.

Compliance:

Follow appropriate standard operating procedures to ensure compliance with applicable regulatory requirements and corporate quality standards.

Additional Skills/Preferences:

- Applicants who have Diplomate status (parasitology speciality) in the American College of Veterinary Microbiologists (or equivalent board certification) or are seeking parasitology board

certification are preferred

- Experience in companion animal parasitology preferred
- Excellent knowledge of GCPs, VICH and other Guidelines and GLPs.
- Previous experience working with external contract research organizations.
- Proficiency in oral and written English. Any additional language is beneficial.

If interested in being considered for the role, please apply via the link below:

https://xjobs.brassring.com/TGWebHost/jobdetails.aspx?partnerid=25428&siteid=5645&AReq=24509BR&Codes=HM