Title: Clinical Scientist
Location: Worcester, MA (Hybrid)
Reports to: SVP, Head of Clinical Development

Join our team! At Mustang Bio we are driven by people. The patients we serve and the team we are building are the driving forces behind our mission to deliver life-changing first-in-class cell and gene therapies to patients with genetic diseases and aggressive forms of cancer.

Overview:

The Clinical Scientist serves in a clinical/medical research function and applies a broad knowledge of human clinical research, therapeutic area knowledge, and scientific resources to facilitate the design and implementation of clinical studies and the generation of regulatory required documentation. The primary objective and purpose of this position is ensuring that clinical studies are conducted in a scientific manner considering stated goals, objectives, and operational and regulatory requirements. Works closely with Medical Affairs, Biometrics, Translational Science, and other Clinical Development staff and develops approach to data review from a medical and scientific perspective. This position is based in an office or home office environment. Minimal travel to Business Units and/or Corporate Offices may be required (hybrid 2-3 days onsite).

Essential Job Functions:

- Review ongoing clinical trial data to identify safety concerns or potential safety concerns across therapeutic areas and all programs
- Independently develop data review and data integrity approaches for individual clinical trials as well as integrating data from our alliance partners and scientific research collaborations
- Assist in the preparation and review of key study regulatory documents, submissions, and annual reports
- Develop scientific/medical training materials for clinical operations and cross-functional teams
- Assist Clinical Operations teams with preparation and presentation of materials at site initiation visits, as necessary
- Support the Clinical Operations department with study start-up and study maintenance activities, as needed
- Assist Clinical Development team with preparing presentations for scientific conferences, clinical study investigator meetings, board meetings and expert advisory meetings, as well as developing manuscripts for publication in peer-reviewed journals
- Perform or assist with other clinical scientific activities, as assigned

Experience & Qualifications:

- Minimum of MPH, MS, PharmD, PhD or MD in field relevant to clinical research preferred
- Minimum 5 years of experience with PharmD, PhD or MD or minimum 8 years with MS
- Must be able to use scientific and clinical knowledge to develop relevant medical/scientific materials for clinical trials
- Excellent writing and organizational management abilities
- Knowledge of clinical operations and FDA and HIPAA rules, ISO regulations, and ICH guidelines
- Knowledge of good document management practices
- Experience in a matrixed work environment
- Ability to independently anticipate and resolve problems
- Ability to independently lead cross-functional efforts and teams
- Proficiency in clinical data interpretation
- Excellent skills in oral and written communications
- Ability to work independently with minimal guidance as well as collaboratively within a team setting
- Travel <10%