Stealth BIOTHERAPEUTICS | Leading Mitochondrial Medicine

Position Title: Director, Clinical Development

Position Summary:

The Director, Clinical Development will report to our Chief Clinical Development Officer with primary responsibility to advance the development of Stealth's clinical assets. In addition to planning, implementing, and managing clinical research studies in collaboration with internal and external stakeholders to ensure trial integrity and success, responsibilities will include: providing strategic input and development support for clinical plans, evaluating and interpreting clinical data, reviewing and authoring study-related documents, monitoring patient data and data collection status, and delivering protocol-related training to CROs and clinical trial sites. Further responsibilities include developing and maintaining strong relationships with key opinion leaders and investigators to gain insights for clinical trial planning purposes.

Responsibilities:

- Contributes to the creation of the clinical development plan in partnership with cross-functional team members as well as KOLs and CROs
- Authors, manages, and reviews study materials for competent authorities' submissions; gains insights from clinical and scientific experts
- Analyzes and interprets clinical trial data, collaborating with program lead to conduct data review
- Troubleshoots internal and external conflicts to ensure trial integrity and success
- Develops and presents protocol training for CROs and trial sites
- Engages with trial sites and CROs regularly to monitor subject data and data collection status, ensure adherence to protocol, and evaluate consistency of data
- Develops and maintains knowledge of the therapeutic area, current medical practice, and pharmaceutical regulations to help ensure best practices
- Establishes and develops professional relationships with external experts to gain clinical insights for clinical trial planning
- Provides educational and scientific support to external experts as needed
- Provides scientific support for publications, presentations, and conferences
- Participates in scientific advisory boards
- Provides effective clinical presentations to internal and external audiences

Competencies:

- Ability to proactively predict issues, resolve problems, and make sound decisions
- Excellent oral and written skills; ability to effectively communicate complex information
- Ability to analyze & interpret clinical data
- Strong organizational abilities and attention to detail
- Ability to work independently and on a team
- Ability to effectively collaborate with internal and external stakeholders
- Strong understanding of drug drug interactions
- Ability to gain insights from clinical and scientific experts
- Self-starter with the ability to "flex" between strategy and operational execution

Requirements:

- Bachelor's degree in science with 7 years related experience or Master's degree and 4 years related experience; PharmD preferred
- Understands scientific & clinical research process
- Experience participating in ad boards and working with KOLs highly preferred

- Approximately 30% travel