Director/Senior Medical Director

Remote

Summary

Reporting to the Chief Medical Officer, the Director/Senior Medical Director, conducts and oversees Clinical Science activities in support of Sierra business objectives.

Responsibilities

- Oversees the medical aspects of the cross-functional early stage development programs for Sierra.
- Contributes to the development of and executes the clinical development strategy for early stage assets based on the science and the available clinical and preclinical data, while balancing regulatory, operational, scientific and business drivers in the development of such a plan.
- Ensures alignment and approval of senior management prior to implementation and execution of the strategy.
- Oversees, leads and contributes to the team(s) responsible (including internal and CRO staff) for medical aspects of the execution of clinical studies and overarching clinical development plan including medical monitoring and pharmacovigilance activities.
- Oversees and participates in all aspects of clinical studies for a program including protocol development, study execution, analysis and reporting of data, and regulatory submissions.
- Designs, vets or validates (for example, with KOLs), and contributes to the clinical development strategy for a molecule/company and participates in the design of appropriate clinical development plans to achieve that strategy.
- Assists with in-licensing, evaluating and assessing clinical development strategies for new technologies and potential new pipeline products.
- Liaises with KOL's, developing relationships and strategy.
- Collation, interpretation, presentation and dissemination of clinical data and outcomes to stakeholders internal and external to firm Including preparation of materials to senior management or external advisors, ensuring effective presentation of data.
- Works closely with clinical operations, PM, Safety, Regulatory, Biometrics and other internal colleagues to ensure effective and efficient execution of studies and mitigation of risks.
- May supervise other members of Clinical Science team.
- Regular international travel to attend meetings is required.
- Other related duties as assigned.

Qualifications and Requirements

- MD degree or equivalent, Oncology or other relevant drug development experience or the equivalent combination of education and experience.
- Minimum 3 years clinical research or clinical development experience for Director and +5 years' experience for Senior Director, with previous experience in a similar environment.
- Phase I to III drug development experience preferred.
- Knowledge of EMA, FDA and related regulations.

- Team player with professional presence coupled with excellent verbal and written skills.
- Ability to maintain effective working relationships with staff members, investigators and medical community.
- Ability to work effectively in a rapidly changing and scaling environment.
- Must be able to travel without any legal restrictions across international borders.

How to Apply

To apply for this role, please submit your CV and cover letter in PDF format to hr@sierraoncology.com. Please indicate the position title in the subject line of your email.

We thank you in advance for your interest in Sierra. We will contact you directly should we wish to arrange a meeting to discuss this position further.