Company: Pfizer Inc.
Salary range:
Job Title: Associate Director, Clinician Medical Monitor (MD Required)
Brief description of the position: Pfizer Vaccines has a very active vaccine pipeline and is currently developing several vaccines. This position is to join the COVID vaccine program and/or the RSV vaccine program. Both are conducting large vaccine efficacy studies. This position would provide an opportunity to join a very dynamic team as physician/medical monitor and work on clinical trials and other related vaccine development work.

Pfizer’s Pearl River site has been involved in vaccine development for over 100 years. Currently, hundreds of staff at the site support Pfizer’s large vaccine pipeline as well as its approved vaccines creating a large vibrant professional community committed to vaccine development. The site is in commuting distance of Westchester County NY, upper Manhattan, and Bergen County NJ.

ROLE SUMMARY

The Clinician Medical Monitor (CMM) is accountable for providing medical and scientific expertise and oversight for Global Clinical Trials and serves as a single point of accountability for design, execution, monitoring, delivery and reporting of one or more clinical studies to ensure patient safety. Additional study level activities include presentation of study results to internal and external committees or advisory boards, presentation of data at international scientific meetings and publication of study results in peer reviewed journals.

The CMM will also act as a medical monitor for select clinical trials, including phase 2B and phase 3 registration studies. The CMM is accountable for patient safety for subjects participating in Pfizer clinical trials and provides medical guidance during the design, execution, and reporting for clinical studies.

In addition to study level activities, the CMM will participate in program level activities including authoring/reviewing safety and efficacy summaries, clinical overviews, investigator brochures, risk management plans, periodic safety update and clinical sections of product labels.

In addition to work on specific assets the CMM will participate in standing committees, review compounds for potential in-licensing, including performance of due diligence reviews, and provide assistance to new business development on market opportunities and the target product profile.

For a complete job listing please visit: https://pfizer.wd1.myworkdayjobs.com/PfizerCareers/job/United-States---New-York---Pearl-River/Associate-Director--Clinician-Medical-Monitor--MD-Required--4814842-3

Qualifications you are seeking: QUALIFICATIONS

Candidate demonstrates a breadth of diverse leadership experiences and capabilities including: the ability to influence and collaborate with peers, develop and coach others, oversee and guide the work of other colleagues to achieve meaningful outcomes and create business impact.
MD degree

Licensed by a health authority to prescribe medicines (independent of supervision) for at least one year (post "intern/houseman" year), and has utilized the license to prescribe medicines in a patient care setting for an aggregate duration of at least one year.

M.D. required, internal medicine or surgical certification is highly desirable. Training and experience in infectious diseases or infection control experience in the hospital setting is preferred

Knowledge of Good Clinical Practice

Experience in small molecule or vaccine clinical development and conduct of clinical trials for treatment or prevention of diseases in hospitalized patients is preferred

Experience in assessment of adverse events and safety among hospitalized patients participating in therapeutic clinical trials is preferred

Skilled in protocol design, interpretation, and medical monitoring

General therapeutic area knowledge

Excellent written and oral communication

Capacity to adapt to a fast-pace and changing environment
Phone:
Fax:
Address:

Website: