

## **Director or Senior Director or Executive Director of Pharmacovigilance Physician/Clinical Development/Oncology**

Hengrui Therapeutics, Inc. is recruiting for a Director or Senior Director or Executive Director of Pharmacovigilance Physician/Clinical Development/Oncology, located in Princeton, NJ.

Hengrui Therapeutics is a biotechnology startup, focusing on early stage clinical development of innovative therapeutic solutions in areas of major unmet medical needs, such as oncology, immunology/inflammation, metabolic disorders, and other therapeutic areas. We offer abundant opportunities for motivated pharmaceutical professionals to pursue their scientific/clinical interests, make an impact and maximize their growth potential.

The individual for the position will be responsible for providing ongoing medical evaluation/review and assessment of the risk-benefit profile for our drugs in development, while managing all safety issues. He or she will be responsible for the development of overall risk management plans, pharmacovigilance strategies and risk minimizations activities.

The individual will be responsible to oversee, prepare and review scheduled periodic/annual reports (FDA) and Periodic Safety Update Reports (FDA and Global FDA). He or she will be one of the main contributors of INDs, IMPDs and other regulatory submissions, and play an important role in the required regulatory interactions with the FDA, EMEA and other regulatory authorities.

The role requires a motivated, energetic and highly entrepreneurial individual who demonstrates outstanding scientific and pharmacovigilance knowledge and the highest personal and ethical standards. The successful candidate must be equally comfortable among the team as well as working independently.

Other functional groups represented on the team include clinical oncology, operations/project management, regulatory affairs, biostatistics, clinical pharmacology, commercial assessment functions, and functional contractors.

### **Required Education & Qualifications**

The position requires a minimum of an advanced degree in medicine. A minimum 3 years Drug Safety/Pharmacovigilance experience in a pre-approval setting. Direct experience in the Oncology area preferred. Experienced in safety reviews of AEs/SAEs, IBs, protocols, clinical study reports, informed consents, and other safety-related documents.

Strong familiarity of GCP, ICH and Global regulations. Excellent communication, writing and analytic skills.

Experience in Clinical Safety, Pharmacovigilance and risk management within the pharmaceutical industry is required, at least 5 years for Director, 7 years for Senior Director, 10 years for Executive Director. Effective oral and written communication and presentation skills are essential. Able to successfully lead as well as be a team player on cross-functional, cross-cultural teams in a matrix environment, to be solutions oriented, adaptable, deliver desired results and consistently meet goals are required. Travel required, may be up to 20% annually.

**Primary Location** North America-United States-Princeton, NJ

**Organization** Hengrui Therapeutics, Inc.

**Job Function** Full-Time, PV/Clinical Development, MD

**\*Please include salary requirements when submitting CV/Resume to:  
[hr@hengruitx.com](mailto:hr@hengruitx.com)**